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Bwrdd Iechyd Prifysgol
Hywel Dda
University Health Board

Pwyllgor Cyllid a Pherfformiad/ Finance and Performance Committee

DYDDIAD Y CYFARFOD: DATE OF MEETING:	27 August 2026
TEITL YR ADRODDIAD: TITLE OF REPORT:	Macmillan Cancer Support Neighbourhood Investment Partnership Business Case
CYFARWYDDWR ARWEINIOL: LEAD DIRECTOR:	Dr Ardiana Gjini, Executive Director of Public Health
SWYDDOG ADRODD: REPORTING OFFICER:	Dr Bruce Bolam, Deputy Director/Consultant in Public Health Rhian Evans, Principal Project Manager

Pwrpas yr Adroddiad (dewiswch fel yn addas)

Purpose of the Report (select as appropriate)

Ar Gyfer Penderfyniad/ For Decision

ADRODDIAD SCAA SBAR REPORT

Sefyllfa / Situation

The Finance and Performance Committee is asked to scrutinise the affordability, value for money, financial risk, delivery confidence and governance route for the Macmillan Cancer Support Neighbourhood Investment Partnership business case (**Attachment 1**). The partnership aims to improve population health, reduce health inequalities, strengthen community resilience and support cancer prevention through a neighbourhood-based model of prevention and early intervention.

The strategic intent of the business case was supported at the 12 August 2026 meeting of Formal Executive Team. However, further assurance is required before the Health Board enters into any binding funding agreement. A staged governance route is therefore proposed.

It is recommended that the 24 September 2026 meeting of Board be asked to approve the strategic intent of the partnership and authorise continued negotiation of the final agreement. This would not authorise execution of the agreement or create a binding financial commitment. The grant agreement would then return to the 27 October 2026 meeting of this Committee for further deliberation. The agreement would then be submitted for final approval at the 26 November 2026 Board meeting.

Cefndir / Background

The partnership has been developed in response to increasing healthcare demand, rising prevalence of cancer and long-term conditions, persistent health inequalities and the need to strengthen prevention and community-based support across Mid and West Wales. The proposal aligns with:

- A Healthier Mid and West Wales Strategy.
- Community by Design.
- The Social Model for Health and Wellbeing.
- The 20four7 Prevention Model.
- Welsh Government prevention priorities.
- Value-Based Healthcare principles.
- Macmillan Cancer Support's Neighbourhood Health Programme.

The business case has been developed by a project team led by Public Health, Primary Care, Value-Based Health Care and Research and Innovation staff, and through workshops and consultation with Macmillan Cancer Support and key internal and external stakeholders, including third sector organisations and local authorities across the three counties in the Hywel Dda University Health Board (HDdUHB) region.

The business case relates to two separate funding streams:

1. **Social Investment Fund (Repayable)**

The business case proposes approximately **£9.35 million of repayable social investment funding** to support programme implementation, workforce development, evaluation and service delivery over a five-year Neighbourhood Prevention Programme. Repayment would be governed through an outcomes-based funding agreement, with payments linked to agreed delivery, outcome and healthcare-utilisation measures and capped at the value of social investment funding received.

2. **Anchor Grant (Non-Repayable)**

The proposal includes approximately **£2 million of non-repayable Anchor Grant funding**, intended to be invested directly by Macmillan Cancer Support into community and third-sector infrastructure, neighbourhood coordination, community development, volunteering programmes, micro-grants and local prevention initiatives, in alignment with the overall partnership objectives. This element does not create a repayment liability for the Health Board.

The total programme value is therefore currently approximately **£11.35 million over five years**, comprising £9.35 million repayable social investment and £2.0 million non-repayable Anchor Grant funding, within a proposed eight-year partnership and repayment framework.

The proposed agreement may create a material future liability over the life of the contract and should therefore return to Board for final approval before execution, following Director of Finance oversight, procurement advice, legal review and completion of due diligence.

Asesiad / Assessment

The business case sets out the core financial, delivery, evaluation and governance architecture for the proposed partnership. The proposed repayable exposure, annual repayment profile, benefits realisation framework, gateway model, programme risks and key mitigations have been developed and scrutinised through Executive review. The principal issues are summarised below.

Financial and Benefits Assessment

The repayable exposure is the £9.35 million social investment element of the partnership, with total outcomes payments capped at the value of funding received. The current business case sets out an indicative annual repayment profile, with payments beginning at approximately £353,000 in Year 2, rising to approximately £834,000 in Year 3, £1.25 million in Year 4 and £1.67 million in Year 5, followed by approximately £1.75 million annually in Years 6 to 8. This extended profile is intended to avoid concentrating repayment during early implementation, when programme costs are being incurred but longer-term utilisation benefits have not yet fully developed.

Financial benefits have been modelled as cost avoidance and moderated future demand growth rather than guaranteed cash-releasing savings. Benefits modelling has been undertaken using two complementary approaches: TriTech's conservative estimate of approximately £5.2 million over five years, based mainly on smoking and alcohol-related interventions, and National Association of Primary Care modelling indicating potential benefits of up to £12.4 million over five years, based on broader assumptions about participant outcomes and reduced healthcare utilisation. The range reflects the inherent uncertainty in prevention modelling and provides an appropriate basis for testing the programme through implementation and evaluation, with payment contingent on benefits realisation.

The proposal should be assessed through a Value-Based Healthcare lens. The question is not whether prevention produces immediate cash-releasing savings, but whether it delivers sufficient value through improved outcomes, reduced inequalities, better experience, stronger community resilience and moderated future demand, relative to the resources and financial exposure involved. This avoids applying a narrower and higher test to prevention than that applied to other investments when they are nationally directed, improve access or quality, or respond to immediate service pressures, even where they may increase recurrent cost.

Evaluation and Benefits Realisation

The business case includes an evaluation framework combining reach, engagement, retention, behavioural outcomes, patient activation, clinical indicators, healthcare utilisation, longer-term disease outcomes and wider outcomes such as wellbeing, social connection and community participation. It also proposes independent evaluation and validation, including matched comparison methods where feasible. These arrangements are intended to reduce uncertainty progressively as the participant cohort and follow-up period increase. The remaining task is to translate this framework into final contractual schedules, including agreed measures, thresholds, validation rules and reporting arrangements.

Governance and Risk

The business case includes a staged implementation model across three neighbourhoods, formal gateway reviews, a programme risk register, separation of the non-repayable Anchor Grant from the repayable social investment element, alignment of recruitment and delivery commitments with confirmed funding, and a proposed programme governance structure. These arrangements provide important controls over implementation, financial exposure and future expansion.

The main outstanding matters are confirmation that the final outcomes methodology and independent validation arrangements are robust; the maximum Health Board liability and annual repayment profile are explicit and affordable; the contractual fallback mechanism is acceptable; gateway, variation, withholding and termination provisions preserve appropriate

Health Board control; workforce and delivery commitments do not run ahead of confirmed funding; procurement and accounting treatment are confirmed; and the final contractual terms remain consistent with the business case and financial parameters approved by Board.

The recommended approval route is for Board on 24 September 2026 to consider the strategic intent of the partnership. This would authorise continued negotiation and development of the agreement but would not authorise execution or create a binding financial commitment. The grant agreement would then return to the 27 October 2026 meeting of this Committee for further deliberation. The agreement would then be submitted for final approval at the 26 November 2026 Board meeting.

Argymhelliad / Recommendation

It is recommended that the Finance and Performance Committee:

- **SUPPORT** the strategic direction of the Macmillan Cancer Support Neighbourhood Investment Partnership.
- **NOTE** that this does not constitute approval of the final business case, funding agreement or any binding financial commitment at this stage.
- **NOTE** that further work is required on the legal, contractual and financial detail before formal approval can be considered.
- **AGREE** that a final business case will return to the next meeting of the Finance and Performance Committee for scrutiny before any onward recommendation to Board.

Amcanion: (rhaid cwblhau)

Objectives: (must be completed)

Cyfeirnod Cylch Gorchwyl y Pwyllgor:

Committee ToR Reference:

3.1.9 Conduct detailed scrutiny of all aspects of financial performance, the financial implications of significant revenue expenditure (all those over £1million requiring Board approval), business cases (except those that are capital and digital in nature), projects, and proposed investment decisions on behalf of the Board.

Cyfeirnod Cofrestr Risg Datix, Teitl a Sgôr Risg Cyfredol:
Datix Risk Register Reference and Score:

The proposal contributes directly to the mitigation of the following principal risks:

Risk 1198, Ability to shift care into the community, current score 16

Risk 1194, Increasing uptake and access to public health interventions, current score 12

Risk 1193, Failure to address health inequalities, current score 9

Risk 1188, Effective leveraging within partnerships, current score 9.

It also supports the wider strategic response to Risk 1192, Wrong value set for best health and wellbeing,

	current score 12, by accelerating the shift towards prevention, neighbourhood working and community-based support.
Parthau Ansawdd: Domains of Quality Quality and Engagement Act (sharepoint.com)	7. All apply
Galluogwyr Ansawdd: Enablers of Quality: Quality and Engagement Act (sharepoint.com)	6. All Apply
Amcanion Strategol y BIP: UHB Strategic Objectives:	All Strategic Objectives are applicable
Nodau Cynllunio 2026/27 Planning Goals 2026/27	3. 20Four7 Population Health 4. Community by Design 5. People First, Digital Always 6. Safe, Timely, High Quality Care 7. Future Orientated
Hypergyswllt i Amcanion Llesiant HDdUHB 2026 Hyperlink to HDdUHB Well-being Objectives 2026	5. Foster a learning culture by sharing knowledge and applying new ways of working that enhance local and global well-being 6. Work with partners and communities to nurture and support everyone to achieve their best well-being 7. Improve health and well-being by focusing on prevention and early intervention, while working in partnership to ensure everyone has equal access to care and opportunities for good health
Model Cymdeithasol ar gyfer Iechyd a Llesiant : (wedi'i gwblhau ar gyfer newid strategol a/neu newidiadau sylweddol i wasanaethau) A Social Model for Health and Wellbeing : (complete for strategic change and/or significant service changes)	7. All Apply
Gwybodaeth Ychwanegol: Further Information:	
Ar sail tystiolaeth: Evidence Base:	Attachment 1: Macmillan Cancer Support Neighbourhood Investment Partnership: Business Case; Attachment 2: Modelling Benefits; Appendix 1: Health Coaching current position; Appendix 2: Brazil model; Appendix 3: CHWW Business Case; Appendix 4: Demonstrator project report; Appendix 5: Communications and Engagement Plan; Appendix 6: Evidence Reviews and Literature Synthesis; Appendix 7: Macmillan GANTT chart;

	Appendix 8: Macmillan Funding Agreement Template; Appendix 9: Macmillan Quality Impact Assessment; Appendix 10: Macmillan Health Equity Impact Assessment; Appendix 11: Macmillan Equality Impact Assessment.
Rhestr Termau: Glossary of Terms:	Please see full business case (Attachment 1)
Partïon / Pwyllgorau â ymgynhorwyd ymlaen llaw y cyfarfod hwn: Parties / Committees consulted prior to this meeting:	Development has involved Public Health, Primary Care, Value-Based Health Care, Research & Innovation, Macmillan Cancer Support, local authority partners, peak third sector and patient representation organisations and internal service stakeholders.

Effaith: (rhaid cwblhau) Impact: (must be completed)	
Ariannol / Gwerth am Arian: Financial / Service:	The programme will provide approximately £11.35 million of investment, comprising £9.35 million of repayable outcomes-based social investment funding and a £2.0 million non-repayable Anchor Grant. The main financial issue is that repayment may need to be funded from future Health Board resources because benefits are expected mainly through broader value, cost avoidance and moderated demand rather than guaranteed cash-releasing savings. Final approval should therefore be subject to confirmation of affordability, source of repayment, accounting treatment, procurement route, capped liability, fallback provisions and Board approval before execution.
Ansawdd / Gofal Claf: Quality / Patient Care:	The programme is expected to have a positive impact on quality and patient care by improving access to personalised, preventive and community-based support. The Quality Impact Assessment is included as Appendix 9 and is being finalised through internal Health Board processes, including completion of the metrics and risk-scoring sections, before final approval.

Gweithlu: Workforce:	The proposal includes 17 WTE Health Coaches and programme support, alongside neighbourhood-based Community Health and Wellbeing Worker capacity to be commissioned through the third sector. The principal workforce issue is the management of fixed-term recruitment, supervision, redeployment and exit arrangements, ensuring workforce commitments do not run ahead of confirmed funding and gateway decisions.
Tegwch lechyd: Health Equity:	The programme is explicitly intended to reduce health inequalities by targeting neighbourhoods with higher need, improving access to prevention and early intervention, and strengthening local community assets. An equity-focused impact assessment has been completed and is included as Appendix 10.
Risg: Risk:	Key risks relate to future financial liability, affordability, benefits realisation, contractual fallback, gateway control, recurrent sustainability, workforce liabilities, delivery capacity and partner alignment. The staged approval route mitigates these risks by separating strategic-intent approval from final contractual approval, allowing further deliberation by the Committee if required and preserving gateway decision-points during programme delivery.
Cyfreithiol: Legal:	Further legal and risk advice is being sought because of the novel outcomes-based repayment model. Execution of any binding agreement must remain subject to satisfactory assurance on funding and repayment provisions, liabilities, procurement, data sharing, variation, withholding, gateway, fallback and termination arrangements. The final agreement should return to Board for approval before execution because the proposed contract value and potential liability are material to the Health Board's financial and governance position.

Enw Da: Reputational:	The partnership offers a significant reputational opportunity through partnership with Macmillan Cancer Support and visible investment in prevention and communities. Reputational risks include failure to deliver anticipated benefits, unclear public messaging, or entering financial commitments without sufficient assurance.
Gyfrinachedd: Privacy:	The programme will involve referral, assessment, outcomes monitoring and information sharing between partners. Data protection, confidentiality, information governance and data sharing arrangements will reflect established Health Board policies and procedures.
Cydraddoldeb: Equality:	The proposal is expected to have positive equality impacts by improving access to prevention and support in communities experiencing higher levels of need. An Equality Impact Assessment screening has been undertaken and is provided as Appendix 11. This positive evaluation means it has been unnecessary to complete a full Equality Impact Assessment.

Macmillan Cancer Support Neighbourhood Investment Partnership: Business Case



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Executive Summary

This business case seeks approval for a five-year Neighbourhood Prevention Programme. The programme will help Hywel Dda University Health Board (HDdUHB) improve early intervention, prevention and neighbourhood-based support for people affected by cancer and other long-term health conditions.

Funding from Macmillan Cancer Support will be used to develop, test and evaluate a new Community by Design approach in three high need neighbourhoods across Carmarthenshire, Ceredigion and Pembrokeshire. The programme aims to strengthen local communities, improve access to preventive services, and help people stay healthier for longer, reducing the need for avoidable healthcare interventions.

The programme is aligned with HDdUHB's strategic objectives, Welsh Government priorities, Value-Based Health Care principles and the Macmillan Neighbourhood Health Programme. It responds to the increasing demands on our health services, the growing prevalence of cancer and other long-term conditions and persistent health inequalities by investing in the wider determinants of health and shifting resources upstream towards prevention and community-based solutions.

The programme will be funded through a combination of repayable Macmillan Social Investment funding and a non-repayable Anchor Grant. This provides a unique opportunity to develop, test and evaluate a new model of neighbourhood health that addresses the wider determinants of health whilst strengthening community capacity and resilience.

Proposed Model

The programme consists of three integrated components:

Neighbourhood Prevention Support

- Anchor Grant funding for local neighbourhood asset mapping, coordination and community development initiatives, vital to ensuring the other two components can function effectively.
- Strengthens local community infrastructure to enable sustainable prevention and wellbeing activities at neighbourhood level.
- Macmillan will directly commission and fund the third-sector to deliver this element of the programme, with participant inclusion and exclusion criteria established during service design and implementation.

Community Health and Wellbeing Workers (CHWWs)

- Proactive neighbourhood outreach through relationship-based engagement with households experiencing the greatest disadvantage.
- Earlier identification of unmet need, improved access to services and strengthened community resilience.
- Expected to engage 3,000 residents annually.



Health Coaching

- Personalised behaviour change support focused on smoking cessation, alcohol reduction, diet and physical activity.
- Referrals from primary care, community services and priority access pathways including cancer, cardiovascular, respiratory and diabetes services.
- Expected to support 3,000 individuals annually.
- Identification, referral, triage and participant eligibility processes for this component are outlined on pages 10–11.

The programme will also fund neighbourhood-based training and support for ‘Making Every Contact Count’, together with digital tools to support behaviour change, improve access to services, and enable effective monitoring and evaluation of programme impact.

Expected Benefits

The programme aims to deliver:

- Improved health behaviours and self-management.
- Increased patient activation and wellbeing.
- Earlier identification of need and reduced health inequalities.
- Greater utilisation of community assets and support networks.
- Improved integration between health, care and community services.
- Reduction in avoidable demand on primary, community and acute healthcare services.
- Long-term reductions in preventable disease, including cancer, cardiovascular disease and diabetes.

Evaluation and Benefits Realisation

A comprehensive evaluation will be delivered by the TriTech Institute and the HDdUHB Research and Innovation Team with external advisors from the National Association of Primary Care. This will assess how well the programme is working, including its impact on health outcomes, patient experience, use of services and value for money.

The programme will be introduced in one neighbourhood in each county during the first three years. Each neighbourhood intervention will be established for at least three years, with ongoing evaluation. This phased approach will allow learning from early implementation to shape future investment and delivery decisions for the programme and the region more broadly.

Progress will be measured through a set of jointly agreed key performance indicators and annual evaluation reports. Alongside the formal evaluation, the Programme Board will use a Quality Improvement approach, including Plan-Do-Study-Act (PDSA) cycles, to test changes, learn from experience and refine delivery as the programme develops. This will support continuous improvement, ensure services respond to local needs, and help spread effective practice across all elements of the programme.



Delivery and Governance

The programme will be delivered over five years (October 2026 – September 2031). Whilst comparable programmes are typically funded over three years, a five-year period has been agreed to enable the programme to establish neighbourhood infrastructure, test and refine the model, generate robust evidence of impact and fully assess the medium-term benefits of neighbourhood-based prevention. This extended timeframe will support learning, adaptation, benefits realisation and the development of a sustainable commissioning model.

The programme will be governed through a dedicated Programme Board chaired by the Executive Director of Public Health as Senior Responsible Owner, supported by established partnership, assurance and corporate governance arrangements within HDdUHB documented in the Project Governance section of this business case, below.

Financial Summary

The programme seeks £11.35 million of investment over five years, combining £9.35 million of repayable Social Investment Fund funding with £2 million of non-repayable Anchor Grant funding to scale a population health and prevention model that integrates health coaching, community-based support, digital interventions, workforce development and evaluation, delivering long-term health improvement and sustainable demand reduction across the system. Further detail on the investment profile and funding requirements is provided on page 29 of the business case.

The KPIs that will trigger repayment of the Social Investment Fund are outlined on pages 29–30 of the business case. These measures form the basis of the outcomes framework against which programme performance will be assessed. Discussions with Macmillan regarding the repayment mechanism, including the timing, profile and period over which any repayments would be made by the Health Board should the agreed KPIs be achieved, are ongoing and have not yet been finalised.

The proposed funding model includes a repayable loan component, with repayments commencing in Year 2. Although the programme is forecast to generate substantial financial and non-financial benefits, recurrent funding will be required to support ongoing delivery costs beyond the period of initial investment. In the longer term, consideration should be given to the development of pooled funding arrangements with partner organisations, recognising the broader system-wide benefits generated by the programme and supporting a sustainable funding model.



Background

Strategic alignment

This proposal aligns with HDdUHB's strategic direction, Welsh Government policy, Value-Based Healthcare principles and Macmillan's neighbourhood health ambitions. It supports a shift from reactive care to prevention, early support and community-based action, with a clear focus on reducing inequalities, improving outcomes that matter to people, and easing avoidable demand on NHS services.

In particular, the proposed programme brings together three complementary strategic drivers of the HDdUHB A healthier Mid and West Wales- Healthier Lives, Well Lived strategy:

- **Social Model for Health and Wellbeing** – recognising that health is shaped by social, economic, environmental and community conditions, not healthcare alone.
- **Community by Design** – organising support around neighbourhoods, local assets and resident participation.
- **20four7 Prevention Model** – focusing prevention on the 20% most deprived communities, four major behavioural risk factors and seven priority health areas, including cancer, that can prevent the most avoidable harm in our communities.
- **Delivering Value Based Health Care Across the Life-Course: Our Strategic Approach (2026-2031)** - Priority 3: Build the capacity and capability of staff to apply VBHC across planning and operational priorities

Within this programme, the 20four7 model provides the NHS delivery approach for better neighbourhood working. It provides a platform to bring together integrated health and care teams, community organisations and voluntary sector partners to support prevention, wellbeing and early intervention at a local level. By linking clinical services with community-based support, the model and this proposed programme aims to improve outcomes for local people while reducing avoidable demand on acute services.

Funding context

Social Investment Fund (Repayable Grant)

Macmillan and Social Finance have developed social investment mechanisms that provide up-front investment for new community-based models of care. The funding is designed to help systems shift care from hospitals into neighbourhoods and communities. Repayment is linked to the achievement of agreed outcomes and reductions in health care utilisation.

Key features:

- Provides initial capital to establish new neighbourhood care models;
- Focuses on prevention, anticipatory care and community-based support;
- Repayment is linked to demonstrable improvements and savings, and



- Creates a sustainable mechanism for reinvesting resources back into communities.

Anchor Grant (Non-Repayable)

Alongside its social investment, Macmillan Cancer Support will provide a non-recurrent anchor grant to support the community capacity and local infrastructure needed for effective neighbourhood prevention. The grant will be awarded directly by Macmillan to voluntary community and social enterprise organisations and community partners to fund:

- Community capacity building and organisational development;
- Coordination of local partnerships and neighbourhood networks;
- Small grants for community-led prevention initiatives;
- Workforce, volunteer and peer training and development;
- Strengthening community assets, social connections and local participation, and
- Other enabling activities that improve the sustainability of neighbourhood prevention.

A neighbourhood-based approach

Cancer outcomes are influenced by much more than clinical treatment alone. Supporting people and communities to prevent illness, access help earlier, maintain wellbeing during and after treatment, and remain connected to local sources of support are recognised as critical components of effective, sustainable cancer and other chronic disease care.

Community by Design assumes that services, including those for cancer and other chronic diseases, should be organised around neighbourhoods wherever possible, bringing care, support and decision-making closer to people's homes and communities. Key characteristics of a Community by Design neighbourhood include:

- Community-focused – designed around the needs, strengths and aspirations of local people.
- Locally accessible – residents can reach key services, facilities and community assets easily, preferably through walking, cycling or public transport.
- Integrated services – health, social care, voluntary sector and community services work together around the population's needs.
- Strong sense of place and identity – reflecting local culture, heritage, language and community networks.
- Supports wellbeing and prevention – enabling people to remain healthy, independent and connected within their community.
- Flexible to geography – Neighbourhood boundaries reflect how people live their lives, including travel patterns, natural communities, existing service catchments and local relationships, rather than fixed population thresholds or administrative boundaries.



For the purposes of this business case, a neighbourhood is defined as a place-based community, typically centred around a population that can access most of its daily health, social, community and local services within an accessible and well-connected area. It is characterised by strong social relationships, shared local identity, integrated services, and active participation by and accountability to residents in shaping the places where they live, work and connect.

Implementation will adopt a phased, learning-led approach, commencing in Llanelli as the initial implementation site due to established neighbourhood need, local readiness, and the presence of existing community engagement mechanisms and partner infrastructure. This will enable the programme to mobilise at pace in 2027 while testing and refining the model and generating early learning on workforce development, governance, referral pathways, community engagement, evaluation and integration with wider neighbourhood prevention activity. Subject to agreed decision gates and evaluation findings, the programme will then expand to Pembrokeshire and Ceredigion during Years 2 and 3.

The introduction of Community by Design in Wales has presented an opportunity for Health Boards to consider how Primary Care Clusters can be reconfigured to inform and develop systematic wide change through creating Connected Communities (neighbourhoods). Bringing together professional collaboratives to work around natural population groupings (which will differ from the current Cluster footprints), these will feed up into Integrated System groups which could either be developed on a County wide basis (Pembrokeshire and Ceredigion) or a sub County basis in Carmarthenshire (South and North). Developing the model in this way will support our communities in identifying what is most important to them, as well as driving systematic service change that addresses population health needs and prevention.

Further work needs to be undertaken to agree a methodology for defining the natural population groupings as in a primarily rural Health Board using a population basis with a minimum and maximum number might be prohibitive in allowing for more smaller, rural groupings to occur.

Future implementation will be shaped through community co-design with residents, primary care, local authorities, voluntary sector partners and wider stakeholders. Whilst early analysis suggests that neighbourhoods in Pembrokeshire and Ceredigion should be selected based on population health need, geographical deprivation, delivery readiness and existing community capacity, the precise sequencing, delivery footprints and service model adaptations will be determined locally. This will ensure that each neighbourhood model reflects local assets, community priorities, existing provision and the differing challenges associated with urban, rural and coastal communities.

The phased implementation and investment profile have also been deliberately designed to provide assurance regarding the achievement of the programme's Key Performance Indicators (KPIs) and the repayment of the Social Investment Fund. The gradual expansion from one neighbourhood in Year 1, to two neighbourhoods in Year 2 and three neighbourhoods in Year 3 creates defined delivery milestones against which progress can be assessed, benefits realised and repayments triggered. Each phase is linked to a clear set of operational, clinical and system-



value KPIs, including participation rates, patient activation, improvements in cardiovascular and metabolic risk factors, reductions in healthcare utilisation and independently validated demand avoidance.

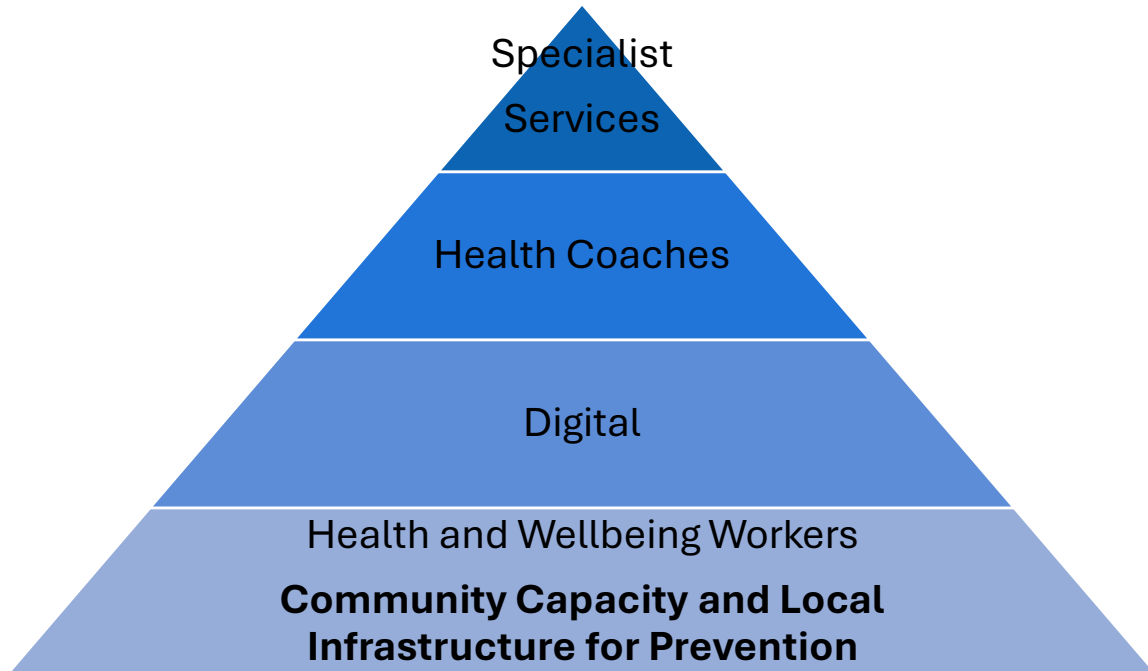
An explicit exit strategy has been incorporated into the programme design, with formal gateway reviews at the end of each implementation phase. These gateways will assess delivery performance, KPI achievement, financial performance, realised benefits and organisational readiness for further scale-up. Should the programme fail to demonstrate sufficient progress against agreed milestones, a decision can be taken to pause, redesign or discontinue further rollout, thereby limiting financial exposure and protecting the Social Investment Fund investment. This staged approach provides confidence that investment is released and repaid in line with demonstrable outcomes and ensures robust governance throughout the life of the programme.

This adaptive approach recognises that neighbourhoods across a large and predominantly rural Health Board area are not uniform. Some areas may benefit from town-centred delivery models serving defined neighbourhood clusters, whilst others may require hub-and-spoke, outreach or networked approaches to support more dispersed populations. By combining a common set of principles with local flexibility, the programme will be able to respond to place-specific needs while maintaining a consistent focus on prevention, community wellbeing and reducing health inequalities, including those that may otherwise remain hidden within rural communities.



The Proposed Programme Model

Figure 1 illustrates the relationship between the key components of the programme and how they work together to support neighbourhood-based prevention and wellbeing.



Programme Component 1: Health Coaching

Overview

The Health Coaching Programme will provide structured, personalised support to help people improve their health, reduce modifiable risk factors and manage their wellbeing with greater confidence.

This component builds on two successful HDdUHB initiatives. The Health Coaching Service, developed as an extension of local smoking cessation services, received more than 500 referrals over a 12-month period, with around 50% of participants achieving their primary wellbeing goals. Evaluation identified improvements in confidence, motivation, self-management and sustained behaviour change. Relatedly, a local Lifestyle Medicine programme was selected for development through the nationally recognised Bevan Exemplar Programme and demonstrated improvements in participants' mental wellbeing, confidence, physical activity and social connectedness, with many continuing healthy lifestyle activities and peer-support networks after the programme ended.

The proposed service will support people at risk of poorer outcomes from cancer and other long-term conditions, including those affected by smoking, alcohol use, poor diet, excess weight, low physical activity, low confidence in self-management or wider barriers to health and wellbeing.

Health Coaching is the individual-level behaviour change component of the Neighbourhood Prevention Programme. It will work alongside Community Health and Wellbeing Workers, and existing social prescribing, community assets and specialist services. Its purpose is to turn lifestyle advice into practical, achievable action, based on what matters to the person.

The service will have broad access, but will actively prioritise people with the greatest capacity to benefit. This means people whose outcomes, treatment readiness, quality of life or avoidable healthcare use could improve most through timely, personalised behaviour-change support.

Capacity Modelling

The proposed model is based on a workforce of **17.0 Whole Time Equivalent (WTE) Health Coaches, 3.0 WTE Health Coach Assistants** and **1.0 WTE band 6 Clinical Lead** supported by appropriate clinical and operational supervision. Capacity assumptions have been informed by pilot learning, existing health coaching services, demand modelling and realistic productivity expectations.

Based on current modelling, the service is expected to receive approximately **4,998 referrals annually**, with an anticipated engagement rate of **80-85%**. Allowing for DNA (Did Not Attend) rates and attrition of **20-25%**, the service is projected to support around **3,060 individuals each year** through a meaningful health coaching intervention over 12 weeks.

Each Health Coach is expected to manage an active caseload of approximately **45-60 individuals** at any one time and deliver around **180 completed interventions**



per WTE per annum. Workforce planning recognises the need to balance direct patient contact with administration, multidisciplinary working, supervision, quality improvement and professional development activities to maintain a safe and effective service.

Activity modelling presented in this business case is based on the assumptions above. However, the number of participants engaged over the length of the five-year program is anticipated to expand significantly with group-based and increasing use of digital approaches, including telephone and video consultations, secure messaging and digital self-management tools. These approaches will increase accessibility, increase productivity and support efficient use of workforce capacity without compromising personalised care.

Sensitivity testing indicates that the proposed workforce can accommodate moderate increases in demand through optimisation of digital delivery and group-based interventions. Ongoing monitoring of referral volumes, caseload complexity, waiting times, DNA rates and outcomes will ensure workforce capacity remains aligned to demand and supports continuous service improvement.

Why it is needed

Many cancer and long-term condition outcomes are influenced by modifiable behavioural risk factors, including smoking, alcohol use, unhealthy weight, poor diet and low physical activity. Around four in ten new cancer cases in Wales were linked to potentially modifiable risk factors, including smoking, overweight and obesity. And this burden is not equally shared: after accounting for age, people living in the most deprived fifth of areas are 1.2 times more likely to get cancer than people living in the least deprived fifth.

Health Coaching provides a practical route to earlier prevention and better self-management. It supports people to set goals, build confidence, address barriers and connect to the right community or specialist support. It is a supported self-management intervention that helps people build knowledge, skills and confidence to become active participants in their care.

Identification, Referral and Triage

Health Coaches will be embedded as part of multidisciplinary teams in primary care, with clear links to cancer and other specialist services and pathways, alongside existing social prescribing and new Community Health and Wellbeing Workers as part of the neighbourhood model. The service will support three broad groups:

1. People at increased risk of avoidable poor outcomes because of modifiable behavioural risk factors.
2. People living with or beyond cancer, where health coaching can support recovery, confidence, wellbeing, treatment readiness and connection to local support.
3. People referred from other priority pathways, including cancer, cardiovascular and planned-care pathways.



Health Coaching will complement, not replace, existing specialist services for smoking cessation, alcohol and other drug treatment, weight management, mental health support, and cancer and other chronic disease management. Participants must:

- Be aged 18 years or over.
- Provide informed consent to participate in the programme.
- Meet one or more lifestyle risk criteria including alcohol risk (AUDIT-C), smoking/vaping, overweight or obesity (BMI), other lifestyle risk factors, or referral from priority pathways including cancer, cardiovascular, respiratory and diabetes.
- Be willing to engage in personalised health coaching.

As the service expands and number of a referrals increase, a RAG approach will be refined and developed to aid prioritisation of those with greatest capacity to benefit.

Red — highest priority / coordinated support

People in this group have high need and high potential benefit, but will require active case management with primary and secondary care services. This may include people with multiple lifestyle risks, cancer-related needs, poorly controlled long-term condition risks, treatment-readiness concerns, repeated avoidable healthcare use, or significant barriers linked to deprivation or wider determinants of health.

Support will include rapid contact, a named Health Coach, a structured assessment, more intensive follow-up and links to CHWWs, social prescribing, community support and / or specialist services.

Amber — priority targeted support

People in this group have clear modifiable risk and a realistic opportunity to improve outcomes through structured coaching. This may include people with one or more significant behavioural risks, early long-term condition risk, cancer prevention or recovery needs, or moderate support needs.

Support will include a structured programme of one-to-one or group coaching, goal setting, behaviour-change planning, community connection and planned follow-up.

Green — open-access / supported self-management

People in this group have lower-complexity needs or are ready to benefit from support following brief advice, digital support, group sessions, MECC conversations, community activities or self-management resources.

Support will include light-touch coaching, digital tools, signposting, group offers and an easy route back into the service if needs increase.

Specialist or urgent needs

Where safeguarding, acute clinical instability, severe mental health crisis or complex substance misuse is identified, Health Coaching will not be the primary intervention at that point. The person will be supported into the appropriate specialist pathway,



with health coaching offered later or alongside specialist care where safe and appropriate.

The triage process ensures individuals receive the most appropriate level of support while enabling effective management of referrals from high-volume prevention and priority pathways.

Personalised Health Coaching

Participants receive an initial assessment followed by a structured programme of support focused on behaviour change and self-management. Each participant works with a Health Coach to identify personal goals, develop action plans and build confidence to make sustainable lifestyle changes.

For people affected by cancer, the offer will focus on practical lifestyle support that complements specialist cancer care, for example reducing cancer-related risk factors while waiting for diagnosis or treatment. The offer will complement specialist cancer services by focusing on prevention, recovery, wellbeing and the wider factors that help people live well before, during and after cancer.

Outputs and outcomes

Consistent Programme Outputs

- Health Coaches embedded within primary care and community settings.
- Standardised triage and referral processes.
- Structured health and lifestyle assessments completed.
- Personalised behaviour change plans developed.
- Referrals and signposting to community and specialist services.
- Ongoing coaching and follow-up support.
- Consistent outcome measurement and evaluation.

Illustrative Service Outputs

Based on pilot activity and projected scale-up, excluding service expansion to group-based and digital interventions:

- Approximately 4,998 referrals annually.
- Around 3,060 individuals supported each year.
- Increased engagement with healthy lifestyle interventions.
- Increased referrals from cancer services and other priority pathways, with approximately 30% of Health Coaching activity supporting cancer prevention, recovery and wellbeing.
- Improved access to preventative and community-based support.
- Increased uptake of behaviour change interventions.

Contribution to Programme Outcomes

This component contributes to the wider prevention programme by delivering:

- Earlier identification of behavioural and lifestyle risk factors.



- Improved health behaviours and self-management.
- Increased patient activation, confidence and resilience.
- Reduced risk of chronic disease development and progression.
- Better support for people living with long-term conditions.
- Improved wellbeing and quality of life.
- Better integration between primary care, community services and specialist pathways.
- Reduced long-term demand on health and care services.
- Stronger prevention and support for priority pathways, with cancer support highlighted through risk reduction, personalised wellbeing support, recovery after treatment, and links to community and specialist services.

Clinical Governance

The Band 6 Clinical Lead will provide professional and operational leadership for the Health Coach workforce, ensuring robust clinical governance arrangements are embedded across the service. The post holder will be responsible for maintaining safe, effective and consistently high-quality delivery of health coaching interventions through regular supervision, competency assessment, audit, and quality assurance processes. They will ensure adherence to organisational policies, information governance requirements, safeguarding procedures, and agreed escalation pathways, while supporting continuous professional development and workforce capability. This dedicated leadership capacity will strengthen oversight, mitigate clinical and operational risks, support service evaluation and improvement, and provide assurance that health coaching services are delivered consistently, safely and in line with the Health Board's policies and procedures.

Digital Enablement

Digital enablement will provide support to the Health Coaching Service and wider Neighbourhood Prevention Programme, building upon existing HDdUHB digital infrastructure and capabilities. This includes the use of REDCap for health coaching data capture, outcome monitoring and service evaluation; integration with existing primary care and GP-facing systems where appropriate; and the ORCHA platform to provide access to clinically reviewed and quality-assured health and wellbeing applications. The programme will also align with and support referral or signposting pathways into established prevention and wellbeing services, including Waiting Well, Keeping Well, smoking cessation, weight management and other population health initiatives.

The digital component will facilitate targeted service improvements through streamlined referral and triage processes, structured and standardised data capture, automated reminders, behaviour-change nudges, follow-up prompts, participant self-management resources and access to approved digital health applications. These capabilities will support Health Coaches in maintaining engagement between sessions, reinforcing agreed goals and outcomes, and managing caseloads efficiently whilst providing a consistent user experience.



Digital tools will complement, rather than replace, face-to-face, telephone, group-based and community-delivered interventions. This blended approach will increase service reach and flexibility while ensuring that individuals who are digitally excluded, have accessibility requirements, or prefer traditional forms of support are not disadvantaged. The programme will adhere to the principles of digital inclusion and equitable access, ensuring that all participants can engage with services in a manner appropriate to their needs. As the programme matures, selected digital capabilities may also be extended to Community Health and Wellbeing Workers, supporting referral tracking, follow-up activity, signposting and outcome monitoring. Routine data collected through REDCap and related systems will support ongoing monitoring, evaluation, quality improvement and benefits realisation. This will include measurement of activity levels, participant engagement, programme completion rates, Patient Activation Measure (PAM) scores, health and wellbeing outcomes, and healthcare utilisation metrics where appropriate and available. Data insights will inform service development, population health planning and evidence-based decision making across the prevention agenda.

All digital developments will be subject to appropriate governance, assurance and technical review processes. This will include compliance with Information Governance requirements, UK GDPR and NHS Wales data protection standards; completion of Data Protection Impact Assessments where required; and adherence to Health Board records management policies. Appropriate cyber security assessments will be undertaken to ensure alignment with NHS Wales Cyber Resilience standards and organisational security controls. In addition, any new or enhanced digital functionality that may influence clinical decision-making, patient care pathways or the communication of health-related information will be assessed through the Health Board's Clinical Safety Framework, including clinical safety review and hazard assessment where applicable.

Future digital investment will be delivered in a phased and affordable manner, aligned to the Health Board's Digital, Data and Technology Strategy, enterprise architecture principles and wider population health objectives. Development will focus on interoperability, accessibility, data quality, cyber resilience and sustainable service design, whilst maintaining alignment with the Health Board's evolving data platform and strategic direction for integrated digital care.

A key principle of the digital approach will be to place greater control and ownership in the hands of citizens, supporting the Health Board's ambition of **"Patient First, Digital Always"**. Through the use of a Patient Engagement Portal and associated digital tools, individuals will be able to access personalised information, resources, appointments, self-management support, educational content and approved health and wellbeing applications at a time and place convenient to them. This will enable people to take a more active role in managing their own health and wellbeing, supporting behaviour change, prevention and early intervention. Digital engagement capabilities will facilitate two-way communication, timely reminders, progress tracking and access to personalised goals, helping individuals become active participants in their care rather than passive recipients of services. The approach supports the wider strategic objective of delivering more proactive, preventative and person-centred care whilst reducing unnecessary barriers to accessing support.



Future digital investment will be delivered in a phased and affordable manner, aligned to the Health Board's Digital, Data and Technology Strategy and the principles of **Patient First, Digital Always**. Development will focus on empowering citizens through digital engagement, self-service and self-management capabilities, whilst maintaining the highest standards of clinical safety, information governance, cyber resilience, accessibility and interoperability across the wider health and care ecosystem.

Making Every Contact Count Workforce Development

Making Every Contact Count (MECC) will be an additional cross-cutting workforce development enabler of the Neighbourhood Prevention Programme. It will provide a common prevention conversation framework across Health Coaching, CHWWs, community partners and relevant frontline teams, helping staff and volunteers make brief, practical behaviour-change conversations part of everyday contact with residents.

Training, refresher sessions and practical conversation tools will help staff and partners raise risk factors sensitively, promote screening awareness and early help-seeking, and connect people to Health Coaching, CHWWs, smoking cessation, weight management, alcohol support, community assets and specialist pathways.

MECC will support the programme's cancer prevention and support ambition, including the expectation that around 30% of programme activity is aligned to cancer prevention and people affected by cancer. Activity will be monitored through training uptake, staff confidence, use of brief intervention approaches where recorded, and referrals or signposting into wider prevention support.

Summary

Health Coaching will provide a broad but prioritised prevention and wellbeing offer for people at risk of poorer cancer and long-term condition outcomes. The service will remain open and inclusive, while using a redesigned RAG model to prioritise people with the greatest capacity to benefit.

The component will strengthen prevention, support cancer recovery and wellbeing, improve self-management and connect people to community and specialist support. It will also provide a measurable pathway from individual behaviour change to improved outcomes and reduced avoidable healthcare use over time.

Digital tools and MECC workforce development will support this component by improving referral, follow-up, behaviour-change prompts, self-management support and consistent prevention conversations across Health Coaching, CHWWs and wider neighbourhood partners. Further detail regarding the current Health Coaching service is provided in Appendix 1.



Programme Component 2: Community Health and Wellbeing Workers (CHWWs)

Overview

Community Health and Wellbeing Workers (CHWWs) provide a proactive, relationship-based approach to prevention, working directly with households in communities rather than waiting for people to access services. Operating at neighbourhood level, CHWWs act as trusted connectors between residents, community resources and health and care services, helping to identify unmet need early and support people before issues escalate into crisis. The model complements existing services, including Health Coaching and social prescribing, by providing ongoing engagement, practical support and continuity for residents.

Why it is Needed

Many people living in deprived or underserved communities experience barriers to accessing support and may not engage with traditional health services until problems become more serious. Health inequalities are often driven by wider social, economic and environmental factors that cannot be addressed through clinical care alone.

The CHWW model responds to this challenge by taking a preventative, community-based approach. Through direct neighbourhood outreach and trusted relationships, CHWWs identify health, wellbeing, social and practical needs earlier, improve access to support, strengthen community connections and help communities build resilience and self-management capacity.

The Model

CHWWs work within defined neighbourhoods linked to GP practices and Primary Care clusters, alongside Health Coaches and wider community partners.

Using a structured household engagement approach, CHWWs:

- Proactively engage with households
- Build trusted relationships with residents
- Identify unmet health, wellbeing and social needs
- Support service navigation and access
- Connect people to community, voluntary and statutory services
- Encourage self-management and prevention
- Strengthen community participation and local resilience

The model adopts a whole-household approach, recognising that health and wellbeing are influenced by family, social and community circumstances as well as individual factors.



Delivery Approach

Each CHWW is responsible for a defined locality and works closely with GP practices, Health Coaches, community organisations and other local services. The approach focuses on:

- Early identification of need
- Chronic disease prevention and wellbeing promotion
- Supported referrals and follow-up
- Reducing barriers to access
- Strengthening links between communities and services
- Building community capacity and local solutions

The model is intended to generate long-term relationships within neighbourhoods, creating a visible and trusted presence within communities.

Expected Outputs

System Outputs

- Households proactively contacted and engaged
- Residents supported through personalised interventions
- Needs assessments completed
- Referrals and community connections facilitated
- Increased utilisation of community assets and services
- Improved intelligence regarding local population needs

Community Outputs

- Increased access to preventative support
- Improved uptake of health and wellbeing services
- Greater community participation and connectedness
- Improved confidence and self-management among residents
- Stronger relationships between communities and local services

Illustrative Reach

Based on similar UK services, including Westminster and Cornwall, a team of four CHWWs and a team lead is expected to support around 500 households a year once established. Across three neighbourhood teams, this equates to focused engagement with at least 3,000 people annually, alongside wider community contacts and referrals into tailored health, wellbeing and cancer prevention support.

Contribution to Programme Outcomes

The CHWW model contributes to the wider programme by delivering:

- Earlier identification and management of need
- Improved access to prevention and wellbeing support
- Increased self-management and health confidence
- Reduced social isolation and loneliness
- Improved uptake of screening, immunisation and preventative interventions



- Better connection between residents and community assets
- Reduced avoidable demand on primary, urgent and emergency care services
- Improved understanding of population need and health inequalities

Summary

The Community Health & Wellbeing Workers model offers Hywel Dda University Health Board a practical opportunity to shift from a reactive healthcare model towards a preventative, community-based approach that addresses the root causes of ill health. The programme is designed to reduce avoidable demand on primary and urgent care services by proactively supporting households with issues such as poverty, housing difficulties, social isolation, low health literacy, and access to benefits and community support.

A key benefit for HDdUHB is the potential to release valuable clinical capacity within primary care. The business case highlights that around 20% of GP consultations relate to non-medical issues, equating to an estimated 460,000 consultations annually across the Health Board population. By identifying and addressing these wider determinants of health earlier, CHWWs can reduce repeat GP appointments, improve care navigation, and enable clinicians to focus on patients with complex medical needs.

The programme also supports the Health Board's strategic priorities around prevention, reducing inequalities, and delivering care closer to home. Through proactive engagement with vulnerable and hard-to-reach households, CHWWs can improve access to preventative services including vaccinations, screening programmes, and long-term condition reviews. Evidence from existing UK programmes has demonstrated increased uptake of vaccinations and health checks, alongside reductions in GP consultations.

For HDdUHBs rural and deprived communities, the model offers a mechanism to tackle health inequalities by connecting residents to welfare advice, housing support, employment opportunities, and community resources. This whole-household approach aims to improve wellbeing, reduce loneliness, strengthen self-management of chronic conditions, and increase community resilience.

Beyond direct patient benefits, the CHWWs will generate valuable population insight, helping HDdUHB better understand unmet need within communities and target future resources more effectively. It also supports integrated working between health services, local authorities, voluntary sector organisations, and community partners, strengthening the place-based care agenda. Further information regarding the CHWW model is provided in Appendices 2, 3 and 4.



Programme Component 3: Community Capacity and Local Infrastructure for Prevention

Overview

Community Capacity and Local Infrastructure for Prevention provides the neighbourhood infrastructure needed to enable prevention, wellbeing and early intervention. It connects CHWWs, Health Coaches, residents, community organisations, local assets and statutory services so that prevention activity is coordinated and locally relevant.

The model is not a standalone service. It creates the conditions for prevention activity to flourish in communities, reflecting the Social Model for Health and Wellbeing and the wider social, economic, environmental and community factors that shape health.

Why it is Needed

CHWWs, Health Coaches and wider prevention services depend on strong local infrastructure. Across West Wales, community assets vary in capacity, accessibility and connection to statutory services, limiting opportunities for:

- Early intervention and prevention
- Social prescribing and community referrals
- Volunteering and community participation
- Peer support and community-led activities
- Coordinated support for people and families

This component strengthens the local systems that connect people to support and helps create a more consistent, equitable prevention offer across neighbourhoods.

The Model

The programme combines a consistent infrastructure framework with locally determined priorities, including a cancer prevention and support focus, so communities can respond to local needs within a common approach.

Core Neighbourhood Infrastructure

Every neighbourhood is supported by:

- A Neighbourhood Coordinator function to connect partners and coordinate activity
- Community engagement and co-design processes, such as a locally-appropriate multi-sector forum for engaging residents, patients, community organisations, CHWWs, Health Coaches and statutory partners in shared programme planning
- Community asset and systems mapping



- Community prevention micro-grants
- Communications and participation resources
- A shared learning, monitoring and evaluation framework

Together, these elements provide the foundation for coordinated prevention activity and stronger links between communities and services.

Neighbourhood Prevention Planning and Alignment

Enabled by the Neighbourhood Coordinator function, local partnerships will use community engagement, asset mapping and local intelligence to:

- Understand local strengths, assets and community capacity
- Identify unmet need and health inequalities
- Map relationships across community and statutory services
- Agree locally relevant prevention priorities
- Coordinate action and investment opportunities
- Review and adapt priorities based on learning and changing needs

This will bridge strategic priorities and practical delivery, improving coordination and reducing duplication across programmes.

Flexible Local Delivery

While the core infrastructure is consistent, activities will vary according to local priorities. Partnerships may focus on:

- Strengthening community organisations
- Developing peer support networks
- Recruiting and supporting volunteers
- Improving transport and access to services
- Expanding social prescribing opportunities
- Supporting cancer and other chronic disease carers and targeted population groups
- Developing community hubs and spaces
- Improving outreach, communication and participation

This supports locally owned solutions within a common programme framework.

Expected Outputs

- Consistent System Outputs
- Community Wellbeing Partnerships established
- Local delivery aligned with wider strategies and plans
- Community asset mapping completed
- Community prevention funds distributed
- Regular co-design and learning cycles implemented
- Shared evaluation and learning framework in place
- Illustrative Community Outputs



- New or strengthened community-led initiatives
- Increased volunteering opportunities
- Expanded peer support networks
- Stronger connections between community organisations and local assets
- Contribution to Programme Outcomes

This component strengthens the community conditions required for prevention by supporting:

- Increased community capacity and participation
- Better integration between statutory and community services
- Improved access to community-based support
- Stronger coordination of local assets and services
- Earlier intervention and reduced health inequalities

Summary

Community Capacity and Local Infrastructure for Prevention provides the foundation for sustainable neighbourhood prevention. It brings together residents, community organisations, CHWWs, Health Coaches and statutory partners to strengthen local capacity, improve access to support and support earlier intervention.

Co-production is central to the model. Through engagement, asset mapping and neighbourhood partnerships, residents and partners will shape support around local strengths, needs and priorities while working within a consistent prevention framework.

Anchor Grant funding, paid directly to third-sector organisations and community partners, will strengthen community capacity, local assets, social connections and neighbourhood infrastructure, helping communities build resilience and develop sustainable, locally owned responses to health and wellbeing challenges. Over time, this will support stronger, more connected communities that are better able to prevent ill health, reduce inequalities and improve population wellbeing. Further information regarding the Communication Plan is provided in Appendix 5.

Spread and Scale

The programme has been designed as a scalable neighbourhood prevention model that can be adapted and implemented across diverse communities within HDdUHB and, subject to evaluation findings, more widely across Wales. The phased implementation approach will enable learning from each neighbourhood to inform subsequent delivery, ensuring that effective interventions, governance arrangements and partnership models are refined before wider adoption. Through robust evaluation, quality improvement methodologies and continuous learning cycles, the programme will identify the components that deliver the greatest impact and value. Evidence generated through the programme will support the development of commissioning guidance, implementation resources and workforce models that



facilitate replication and sustainable scale-up. A core principle of future spread and scale will be the promotion of **equitable access to prevention services and outcomes**, ensuring that communities experiencing the greatest levels of deprivation, rurality, exclusion or health inequality are able to benefit from the model. While delivery approaches may vary to reflect local context, the programme will maintain a consistent focus on reducing unwarranted variation, improving equity of service provision and ensuring that all communities have fair access to prevention, wellbeing and community-based support. By combining a consistent core framework with flexibility to respond to local needs, the programme will create a transferable model for reducing health inequalities, strengthening community capacity, improving population health outcomes and supporting a long-term shift towards prevention and neighbourhood-based care.



Benefits Realisation/Resource Releasing Opportunities

The proposed Neighbourhood Prevention Model is expected to deliver measurable benefits across population health, behaviour change, cancer prevention and support, service utilisation and healthcare demand over the short, medium and long term. Approximately 30% of programme activity will be aligned to cancer prevention and support, including reductions in smoking, alcohol-related harm, unhealthy weight and physical inactivity; improved access to screening and early support; and stronger community support for people affected by cancer. The projected outcomes presented below are based on modelling assumptions developed by TriTech, drawing on published literature, evidence reviews, established prevention programmes and the best available data at the time of analysis. These assumptions have been used to estimate participant engagement, sustained behaviour change, reductions in disease risk and the resulting impact on healthcare utilisation. Together, the measures demonstrate the anticipated pathway from participation in preventative interventions to improved health outcomes, reduced incidence of long-term conditions including preventable cancers, and lower demand on primary, community and acute healthcare services. The framework provides a basis for ongoing evaluation and benefits realisation, enabling the programme to assess both immediate improvements in health behaviours and longer-term value through reductions in avoidable healthcare activity and preventable disease burden.

The sensitivity analysis demonstrates a lower and upper range of potential resource-releasing opportunities, reflecting different modelling assumptions. Using the more conservative methodology developed by **Tritech**, the programme is estimated to generate approximately **£5.2 million** in resource release over five years. In contrast, modelling undertaken by the **National Association of Primary Care (NAPC)** suggests that benefits could be as high as **£12.4 million** over the same period. Further information regarding the evidence reviews and literature search is provided in appendix 6.



Tritech Modelling Assumptions

Table 1: Assumed Behaviour Change Maintenance (Relapse) Rates

Relapse rate(who mainatins)										
Type	year 1	year 2	year 3	year 4	year 5	year 6	year 7	year 8	year 9	year 10
Smoking	0.4	0.8	0.9	0.9	0.9	0.9	0.9	0.9	0.9	0.9
Alcohol	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
Healthy living	0.7	0.7	0.7	0.8	0.9	0.9	0.9	0.9	0.9	0.9

Table 2: Projected Cumulative Behaviour Change Outcomes Over a 10-Year Period

Cumaltive numbers										
Type	Year 1	Year 2	Year 3	Year 4	Year 5	Year 6	Year 7	Year 8	Year 9	Year 10
smoking stopped	392	705	987	1241	1469	1675	1860	2026	2176	2311
Alcohol stopped	551	826	964	1033	1067	1084	1093	1097	1099	1101
Healthy living	1125	1912	2463	2904	3300	3657	3979	4268	4528	4763



Table 3: Projected Reductions in NHS Activity and Healthcare Utilisation Attributable to Behaviour Change Interventions

NHS demand reduction (lower)										
Smoking	Year 1	Year 2	Year 3	Year 4	Year 5	Year 6	Year 7	Year 8	Year 9	Year 10
GP appointments	783	1410	1974	2482	2939	3350	3720	4053	4353	4622
Outpatient appointments	118	212	296	372	441	502	558	608	653	693
Unscheduled care contacts	20	35	49	62	73	84	93	101	109	116
ED attendances	16	28	39	50	59	67	74	81	87	92
Hospital admissions	8	14	20	25	29	33	37	41	44	46
Alcohol	Year 1	Year 2	Year 3	Year 4	Year 5	Year 6	Year 7	Year 8	Year 9	Year 10
GP appointments	2203	3305	3856	4131	4269	4338	4372	4389	4398	4402
Outpatient appointments	110	165	193	207	213	217	219	219	220	220
Unscheduled care contacts	28	41	48	52	53	54	55	55	55	55
ED attendances	33	50	58	62	64	65	66	66	66	66
Hospital admissions	33	50	58	62	64	65	66	66	66	66
Total/year	Year 1	Year 2	Year 3	Year 4	Year 5	Year 6	Year 7	Year 8	Year 9	Year 10
GP appointments	2987	4715	5830	6613	7207	7687	8092	8442	8750	9024
Outpatient appointments	228	377	489	579	654	719	777	827	873	913
Unscheduled care contacts	47	77	98	114	127	138	148	156	164	171
ED attendances	49	78	97	112	123	132	140	147	153	158
Hospital admissions	41	64	78	87	93	99	103	106	109	112

3 year total	
GP appointments	13531
Outpatient appointments	1093
Unscheduled care contacts	221
ED attendances	224
Hospital admissions	182

5 year total	
GP appointments	27351
Outpatient appointments	2326
Unscheduled care contacts	462
ED attendances	458
Hospital admissions	362

10 year total	
GP appointments	69347
Outpatient appointments	6436
Unscheduled care contacts	1238
ED attendances	1189
Hospital admissions	892

Table 4: Projected Reduction in Major Long-Term Conditions Attributable to Programme Interventions

3 year prediction	
Cancer reduction	36
CVD reduction	37
Diabetes reduction	261

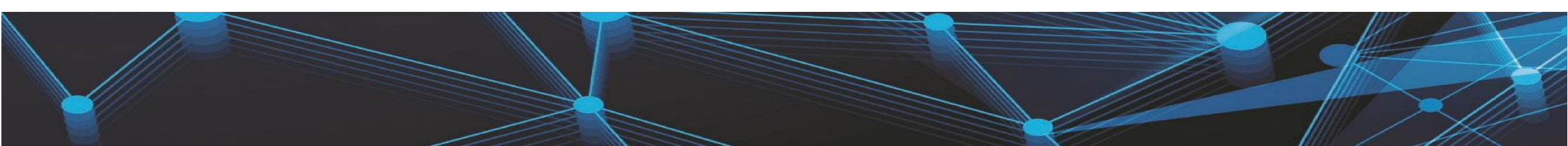
5 year prediction	
Cancer reduction	77
CVD reduction	80
Diabetes reduction	486

10 year prediction	
Cancer reduction	203
CVD reduction	212
Diabetes reduction	1169



Table 5: Potential Resource Releasing Opportunities from the Tritech Modelling

		3 year total		5 year total		10 year total	
		Activity	£	Activity	£	Activity	£
Smoking/Alcohol	GP appointments	13,532	405,960	27,352	820,560	69,347	2,080,410
	Outpatient appointments	1,094	234,116	2,327	497,978	6,436	1,377,304
	Unscheduled care contacts	222	27,676	463	57,721	1,240	154,587
	ED attendances	224	72,128	459	147,798	1,189	382,858
	Hospital admissions	183	717,818	363	1,423,868	892	3,498,870
	Total	15,255	1,457,698	30,964	2,947,924	79,104	7,494,029
Reduction in Prevalence	Cancer	36	383,652	77	820,589	203	2,163,371
	CVD	37	167,388	80	361,920	212	959,088
	Diabetes	261	596,094	486	1,108,546	1,169	2,665,091
	Total	334	1,147,134	643	2,291,055	1,584	5,787,550
Total Resource Released			2,604,832		5,238,979		13,281,579



National Association of Primary Care Modelling Assumptions

This KPI Framework sets out a structured approach for measuring programme performance, outcomes and value across the full pathway of care, from initial engagement through to population-level impact. Organised around six stages, Reach and Engagement, Activity, Behaviour Change, Health Improvement, Healthcare Demand, and Population Impact, the framework provides a clear line of sight between programme delivery and the outcomes that matter to individuals, the Health Board and funders. By combining activity, outcome, cost and comparative measures, the framework is designed to support continuous learning, demonstrate value for money, and provide a robust basis for outcomes-based funding and performance management.

Table 6: Cohort-Based Intervention Model provided by the National Association of Primary Care

Cohort	Cohort ideas	Intervention	Hours	Cost per engaged person	Savings per person	FTE/coach time allocation	People supported per year	Total cost	Total benefit	Net benefit	ROI
Red	Recent discharge, recurrent COPD exacerbations, persistent high users, poorly controlled LTCs	Named coach, structured assessment, approximately 6 one-to-one contacts, CHWW and clinical coordination	6.00	£205	£279	60%	3,276	£672,000	£913,409	£241,409	1.4
Amber	Significant behavioural risk, early LTC deterioration, pain, anxiety/depression, cancer recovery	Blended one-to-one and group coaching, planned follow-up, community connection	1.50	£51	£93	20%	4,368	£224,000	£404,159	£180,159	1.8
Green	Lower-complexity lifestyle risk, rising BMI, moderate anxiety, motivated self-referrers	Digital, group, MECC, community support and easy re-entry	0.12	£4	£22	20%	54,600	£224,000	£1,176,755	£952,755	5.3
Over 1 year							62,244	£1,120,000	£2,494,323	£1,374,323	2.2
Over 5 years							311,220	£5,600,000	£12,471,617	£6,871,617	2.2



Project Costs

Table 7 summarises the estimated costs associated with implementing and sustaining the programme over a five-year period. The investment includes funding for health coaching staff, workforce development, digital support, evaluation, programme management, community health and wellbeing workers, and local prevention infrastructure. Around 30% of funded activity will be planned and monitored as cancer prevention and support activity, including behaviour change support, MECC for cancer prevention, referral pathways from cancer services, community support for people affected by cancer, and local infrastructure that improves access to prevention, screening and wellbeing support. The total investment required is estimated at £8.68 million, comprising £8 million from the Social Investment Fund and £2 million from the Anchor Grant, with expenditure reflecting the phased expansion of services during the early years of implementation.

Table 7: Five-Year Programme Investment Costs by Funding Source

Social Investment Fund (Repayable)	Year 1	Year 2	Year 3	Year 4	Year 5	Yr 1-5 total
Health Coaching Staff 1 WTE Band 6 Clinical Lead 17 WTE Band 5 Health Coaches 3 Band 4 Health Coach Assistants	£607,858	£900,651	£1,041,869	£1,041,869	£1,041,869	£4,634,116
Community Health and Wellbeing Workers 15 WTE Band 3 CHWW	£176,670	£353,340	£530,010	£530,010	£530,010	£2,120,040
Training Costs	£18,500	£18,500	£18,500	£18,500	£18,500	£92,500
Making Every Contact Count Training	£50,000	£50,000	£50,000	£50,000	£50,000	£250,000
Laptops	£2,834	£2,835	£2,836	0.00	0.00	£8,505
National Advisory Function	£10,800	£10,800	£10,800	£10,800	£10,800	£54,000
Patient Activation Measures Licences	£47,165	£0	£0	£0	£0	£47,165
Digital Support 2 WTE Band 6 Digital Project Managers Digital enablement costs	£200,000	£200,000	£200,000	£200,000	£200,000	£1,000,000
Tritech Evaluation	£85,645	£85,645	£85,645	£85,645	£85,645	£428,225
Project staff cost 1 WTE Band 7 Project Manager 0.6 WTE Band 6 Project Support 0.2 WTE Band 9 Public Health Consultant	£144,870	£144,870	£144,870	£144,870	£144,870	£724,350
Total	£1,344,342	£1,766,641	£2,084,530	£2,081,694	£2,081,694	£9,358,901
Anchor Grant (Non-Repayable)	Year 1	Year 2	Year 3	Year 4	Year 5	Yr 1-5 total
Community Capacity and Local Infrastructure for prevention	£166,000	£332,000	£499,000	£499,000	£499,000	£1,995,000
Total	£166,000	£332,000	£499,000	£499,000	£499,000	£1,995,000
Total						£11,353,901



Proposed Repayment Plan

The proposed Macmillan Repayment Plan is structured to provide HDdUHB with significant upfront investment to support programme delivery, combining a non-repayable Anchor Grant of £2 million with a repayable Macmillan loan of £9.36 million over the life of the scheme. Repayments are performance-based and linked to the achievement of healthcare consumption reductions, with repayment triggers commencing from Year 2 once 80% of the healthcare consumption reduction modelling is realised. The repayment profile is phased, beginning at 20% in Year 2 (£353k), increasing to 40% in Year 3 (£834k), 60% in Year 4 (£1.25m) and 80% in Year 5 (£1.67m), before reaching full repayment levels from Years 6 to 8 (£1.75m annually). This approach enables HDdUHB to benefit from early investment and implementation support while aligning repayments to the delivery of measurable value and system-wide efficiencies, creating a sustainable model in which financial returns are generated through reduced healthcare utilisation and reinvested into ongoing service transformation.

Table 8: Proposed Repayment Plan

Macmillan Repayment Plan

	Anchor Grant (Non-Repayable)	Macmillan Loan	In Year Repayment %	Hywel Dda Repayment
Year 1	£166,000	£1,344,342		£0
Year 2	£332,000	£1,766,641	20%	£353,328
Year 3	£499,000	£2,084,530	40%	£833,812
Year 4	£499,000	£2,081,694	60%	£1,249,016
Year 5	£499,000	£2,081,694	80%	£1,665,355
Year 6				£1,752,463
Year 7				£1,752,463
Year 8				£1,752,463
Total	£1,995,000	£9,358,901		£9,358,901



Key Performance Indicators

The following measures have been identified to monitor the effectiveness, impact and value of the project, providing a balanced assessment of service activity, participant engagement, cancer prevention and support, clinical outcomes, healthcare utilisation and long-term population health benefits. Reporting will specifically track the programme's expectation that around 30% of activity supports cancer prevention, earlier support and people affected by cancer. Together, the measures will enable the programme to evaluate progress towards its prevention objectives, demonstrate return on investment and support ongoing service improvement and future scaling decisions.

Table 9: Key Performance Indicators, Targets and Reporting Arrangements

Key Performance Indicators (KPI)						
Indicators of Service performance which will be reported monthly/ quarterly/ annually.						
KPI No.	Description	Numerator	Denominator	Target Measure	Frequency of reporting	Responsible Owner
1	Number of people seen by the intervention	Participants enrolled	N/A	Annual target	Monthly	Service Lead
2	Retention rate (%)	Participants completing intervention	Participants enrolled	>=80%	Monthly	Service Lead
3	Improvement in PAM	Participants improving PAM	Participants with PAM scores	Increase in patients classified as PAM >2	Quarterly	Service Lead
4	Reduction in Blood Pressure	Change from baseline	Patients assessed	10 mmhg in systolic (approx.20% reduction in CVD risk)	Quarterly	Service Lead
5	Reduction in HbA1c	Change from baseline	Patients assessed	Below Prediabetic range	Quarterly	Service Lead
6	Reduction in Lipids	Change from baseline	Patients assessed	1 mmol reduction in Idlc (approx.20% reduction in CVD risk)	Quarterly	Service Lead
7	Reduction in BMI	Change from baseline	Patients assessed	Below a 25 kpms	Quarterly	Service Lead
8	GP appointments avoided	Estimated avoided	N/A	Cumulative total	Quarterly	Tritech

9	Outpatient appointments avoided	Estimated avoided	N/A	Cumulative total	Quarterly	Tritech
10	Unscheduled care contacts avoided	Estimated avoided	N/A	Cumulative total	Quarterly	Tritech
11	ED attendances avoided	Estimated avoided	N/A	Cumulative total	Quarterly	Tritech
12	Hospital admissions avoided	Estimated avoided	N/A	Cumulative total	Quarterly	Tritech
13	Cancer cases averted	Predicted cases prevented	N/A	3/5/10 year total	Annually	Tritech
14	Strokes/heart attacks averted	Predicted events prevented	N/A	3/5/10 year total	Annually	Tritech
15	Diabetes cases averted	Predicted cases prevented	N/A	3/5/10 year total	Annually	Tritech



Risks, Assumptions, Issues and Dependencies Log

The successful delivery of the project depends upon effective management of strategic, operational and organisational factors that may influence programme outcomes. This RAID Log provides a high-level framework for identifying and monitoring the key **Risks, Assumptions, Issues and Dependencies** associated with the development, implementation and evaluation of the programme. It supports robust governance by providing visibility of factors that could impact delivery, benefits realisation, timescales, stakeholder engagement and long-term sustainability. The RAID Log will be maintained throughout the programme lifecycle and reviewed through established governance arrangements to support informed decision-making, proactive mitigation and successful delivery of the programme's ambitions to improve population health, reduce health inequalities and shift investment towards prevention and community-based care.

Table 10: Programme RAID Log (Risks, Assumptions, Issues and Dependencies)

ID	Type	Description	Impact	Mitigation / Action	Owner	Status
R1	Risk	Long-term sustainability after funding ends	Loss of services/community trust	Develop sustainability plan and future funding routes	Programme Board	Open
R2	Risk	Failure to demonstrate ROI	Ability to repay investment	Implement robust evaluation and KPI framework	Public Health / Tritech	Open
R3	Risk	Rurality and hidden deprivation not captured	Increased inequalities	Use local intelligence and rural pilots	Public Health	Open
R4	Risk	Duplication with existing services	Confusion and inefficiency	Map services and align roles	Commissioning	Open



R5	Risk	Insufficient co-production and engagement	Reduced uptake and impact	Community reference groups and champions	Programme Manager	Open
A1	Assumption	Macmillan funding and Board approval secured	Programme cannot proceed otherwise	Maintain executive engagement	SRO	Open
I1	Issue	Short timescales for business case	Potential design gaps	Continue consultation during mobilisation	Project Manager	Open
D1	Dependency	Board and funding approvals	Programme start delays	Maintain governance timetable	SRO	Active

Macmillan Commissioning Template

A **Funding Agreement example** is provided in Appendix 8 and is intended to support discussions regarding the proposed funding, commissioning and governance arrangements. This template illustrates a potential framework for partnership working between the parties and demonstrates how investment, outcomes, accountability and programme governance could be structured. However, it remains an indicative document and will require further development, review and negotiation to reflect the agreed service model, financial arrangements, outcomes methodology, governance framework and local legal requirements. As such, the document should be regarded as a working draft rather than a final or binding agreement. Any future agreement will be subject to detailed refinement and approval by all relevant stakeholders and participating organisations.



Monitoring and Evaluation

The programme aims to empower individuals to take greater control of their health, improve quality of life, support long-term condition management, and contribute to a more sustainable and prevention-focused healthcare system. A key part of the Health Coaching programme will be a robust evaluation of the programme each year. The evaluation will be carried out by the Tritech and Innovation team within the Research and Innovation department. The purpose of the of the evaluation will be to understand its impact and value as well as inform funding decisions for each year of the programme running from 2027-2032. A robust health coaching evaluation methodology combines programme design standards, process monitoring, and outcome measures such as patient activation, self-reported changes, and clinical indicators.

Aims and Objectives of the Evaluation

The aims of this evaluation are to understand the effectiveness, impact, feasibility, acceptability, value and cost-consequences of the HDdUHB health coaching programme within HDdUHB, with the following specific objectives:

- Does the programme improve participants' health and wellbeing?
- Does it support sustained behaviour change and self-management?
- Does it increase patient activation and confidence?
- Does it reduce avoidable healthcare utilisation?
- What aspects of the programme are most effective, and for whom?
- Does the programme deliver value for money/ROI?

Health Coaching Evaluation Methodology

The evaluation methodology will be structured around the NHS Quality Framework's four domains, integrate process and outcome measures, and use a mixed-methods approach to capture both measurable changes and lived experiences. This will ensure the evaluation is comprehensive, evidence-informed, and actionable for improving future practice.

Mixed-Methods Approach

The evaluation will adopt a mixed-methods approach, combining quantitative outcome measures with qualitative insights from participants, coaches, and stakeholders. This approach will provide robust evidence of both programme effectiveness and implementation learning to support future service development and commissioning decisions.

Quantitative: Standardised tools for activation, PROMS/PREMS, clinical metrics, service usage, NHS demand factors and satisfaction surveys.

Qualitative: Focus groups, interviews, and open-ended feedback to capture context and lived experience.



Data collection & Assessment criteria

Evidence-based evaluations in health and wellbeing coaching use a multi-indicator approach and will involve several different data sources, such as:

Assessment of Service KPIs- to be developed as part of the service process (focusing on the key deliverables expected by the programme – number of participants etc.

Assessment of Clinical Indicators – assessment of changes in specific clinical outcomes impacted by the service - such as weight, HbA1c, blood pressure, or other relevant clinical data (to be determined by service protocol).

Assessment of general health Indicators – assessment of changes in general health outcomes including patient recorded outcome measures such as PAM score etc. (to be determined by service protocol).

Assessment of Patient Activation – Pre- and post-coaching questionnaires to measure confidence, knowledge, and self-management ability. Measure in place to assess compliance over time with the service.

Assessment of Service utilisation- Measures reflecting the use of a range of different health services over a particular timeframe e.g. over the 12-months after starting to receive health coaching, such as: Number of visits to A&E; Number of inpatient appointments; Number of outpatient appointments; Attendance at appointments; Medication required.

Assessment of service user experience and outcomes – Interviews and focus groups with all service users to understand service effectiveness and experiences of service users.

Assessment of staff experience and outcomes – Interviews, focus groups and regular surveys to help determine - Job-satisfaction, Employee well-being, Absence rate (percentage of working days lost due to sickness or other absence),

Assessment of other key considerations – assessment of the effects of the service on other considerations such as – social and economic benefits, carbon footprint etc.

Assessment of System Outcomes – assessment of changes in service use, adherence, or cost efficiency. This will include staff workflows, process mapping, NHS service utilisation.

Project deliverables

Findings will be reported through quarterly interim reports and a final evaluation report including outcomes, case studies, lessons learned, and recommendations for future development and wider implementation. An end of year evaluation report will be created for each year of the project. The reports will focus on assessing:



Effectiveness

- Does participation in health coaching improve health and wellbeing?
- Does health coaching increase confidence and self-management capabilities?
- Are participants achieving their personal health goals?

Experience

- How do participants experience the coaching process?
- What aspects of the programme are most valued?
- What barriers or facilitators influence engagement?

System Impact

- Does participation reduce demand on healthcare services?
- Does the programme support more appropriate use of NHS services?
- Costs and value/ROI

Sustainability

- Are improvements maintained after programme completion?
- What improvements could be made to maximise programme effectiveness?

The yearly evaluation report(s) will look at recommendations of the previous year's findings and look at incremental improvements.

Continuous Improvement and Learning

Alongside the formal evaluation undertaken by the TriTech and Innovation Team, the Programme Board will adopt a **Quality Improvement (QI) approach** throughout the life of the programme. Using recognised improvement methodologies such as Plan-Do-Study-Act (PDSA) cycles, the programme will continuously test, learn and refine interventions based on emerging evidence, stakeholder feedback and operational experience.

The five-year programme duration provides a unique opportunity to move beyond traditional project delivery and establish a structured learning system that supports continuous improvement. Regular learning reviews will be undertaken across Health Coaching, Community Health and Wellbeing Workers, Community Capacity initiatives and digital services to identify what is working well, where improvements are required and how successful approaches can be scaled across neighbourhoods.

This combined approach of evaluation and quality improvement will enable the programme to:

- Assess impact and return on investment.
- Refine delivery models based on evidence and learning.
- Improve outcomes for individuals and communities.
- Respond to local needs and emerging challenges.
- Support innovation and shared learning across neighbourhoods.
- Build a robust evidence base for future commissioning and sustainability decisions.



Project Governance

The Project will be governed through a dedicated Programme Board chaired by the Executive Director of Public Health, acting as Senior Responsible Owner. The Programme Board will provide strategic leadership, oversight and decision-making, whilst ensuring effective delivery across all workstreams. Partnership oversight will be provided through the Social Model for Health and Wellbeing Steering Committee, with representation from local authorities, County Voluntary Councils and wider system partners. Assurance and escalation arrangements will operate through the Public Health Directorate Senior Leadership Team and the A healthier Mid and West Wales- Healthier Lives, Well Lived governance structure, with onward reporting to the Strategy and Planning Committee and Board as required. This governance framework provides clear accountability, robust partnership engagement and alignment with Hywel Dda University Health Board's corporate governance arrangements.

Table 11: Programme Governance Structure, Roles and Responsibilities

Governance Level	Primary Role	Key Responsibilities
HDdUHB Executive Board & Macmillan Cancer Support Board	Partner Corporate Oversight	Strategic alignment, senior accountability, receipt of assurance
Strategy & Planning Committee	Corporate Scrutiny	Programme oversight, escalation management, strategic approval
A Healthier Mid & West Wales Governance	Executive Scrutiny	System alignment, cross-programme coordination
Value & Sustainability Group	Executive Scrutiny	Value for money assessment
Public Health Directorate SLT	Directorate Assurance & Escalation	Performance oversight, risk and financial management
Social Model for Health & Wellbeing Steering Committee	Regional Partnership Oversight	Collaborative governance, partner alignment, strategic challenge
Programme Board (SRO)	Programme Delivery	Delivery oversight, benefits realisation, workstream management



Project Milestones and Timelines

The programme will be delivered over a five-year period from October 2026 to September 2031. Following funding approval, mobilisation, recruitment and training will take place between October and December 2026. Delivery will commence in Neighbourhood 1 in January 2027, with subsequent neighbourhoods introduced annually in January 2028 and January 2029. This phased approach enables workforce development, continuous evaluation, refinement of referral pathways and progressive benefits realisation before achieving full programme coverage. Years 4 and 5 will focus on consolidating delivery, demonstrating outcomes, realising system benefits and securing recurrent funding mechanisms. Further information on Project Milestones and Timelines is provided in Appendix 7.

Table 12: High Level Timeline

Phase	Key Activities	Timeline
Programme Set-Up & Approval	Approvals, funding agreement, governance arrangements, programme mobilisation	Q3-Q4 2026
Neighbourhood 1 Implementation	Specification, procurement, mobilisation, go-live and delivery	Q1 2027 - Q1 2028
Gateway Review 1	Evaluation and review of Neighbourhood 1 outcomes	Q1-Q2 2028
Neighbourhood 2 Implementation	Specification refresh, procurement, mobilisation, go-live and delivery	Q2 2028 - Q2 2029
Gateway Review 2	Evaluation and review of Neighbourhood 2 outcomes	Q2-Q3 2029
Neighbourhood 3 Implementation	Specification refresh, procurement, mobilisation, go-live and delivery	Q3 2029 - Q2 2030
Gateway Review 3	Programme review and sustainability planning	Q2-Q3 2030
Programme Evaluation & Commissioning Readiness	Benefits realisation, final evaluation, commissioning business case	Q3 2030 - Q1 2031
Commissioning Decision & Closure	Board decision on future commissioning and programme closure	Q2-Q3 2031



Project Exit Strategy

The programme has been designed as a phased, gateway-controlled initiative to minimise financial, operational and reputational risk. Should the programme fail to achieve agreed outcomes, the Health Board will retain the ability to pause, redesign or discontinue delivery.

At the end of Years 1, 2, 3, 4 and 5, TriTech will undertake formal gateway reviews to evaluate programme performance, benefits realisation and value for money. These reviews will provide the evidence base for decisions to continue, expand, modify, pause or discontinue the programme, thereby limiting financial exposure and ensuring investment remains outcome-driven.

A formal review will be initiated if the programme fails to demonstrate satisfactory progress against agreed performance, financial and outcome measures, including KPI delivery, participant engagement, workforce sustainability, benefits realisation and value for money, or where external funding arrangements materially change.

Should the programme fail to demonstrate sufficient benefit to warrant continuation, the Health Board will implement a phased and managed exit strategy that safeguards participants, preserves organisational learning, supports workforce transition and enables community partners to conclude activities in a planned and sustainable manner.

A comprehensive final evaluation will be undertaken to document programme outcomes, benefits realised, lessons learned and best practice. This evidence will inform future neighbourhood prevention initiatives, support organisational learning, and provide recommendations for the future development, commissioning and scaling of prevention-focused services.

Whilst the availability of external funding presents an opportunity to increase workforce capacity through the creation of new posts, it also introduces significant workforce, financial and organisational risks. The fixed-term nature of the funding does not remove the Health Board's employment obligations. Employees appointed to fixed-term contracts will retain statutory employment rights, including protection against redundancy and unfair dismissal once the relevant qualifying periods have been met.

A further important consideration is the proposed duration of the funding. As funding is expected to continue for between four and five years, employees engaged on successive fixed-term contracts may acquire the right to be treated as permanent employees unless the continued use of fixed-term arrangements can be objectively justified. Consequently, the Health Board would remain liable for all associated employment costs and liabilities arising from these posts, including redeployment obligations, redundancy costs (such as contractual redundancy payments, notice pay, accrued annual leave and, where applicable, pension-related costs), together with the risk of employment claims should the funding cease.



Careful consideration should therefore be given to the long-term affordability and sustainability of these roles, as well as the potential expectation that services and staffing established through the funding will continue beyond the funded period. Alternatively, the Health Board must be satisfied that there is a clearly defined exit strategy, adequate provision for any future redundancy liabilities, and a robust business justification for maintaining fixed-term status throughout the life of the programme. Failure to address these issues could expose the Health Board to significant financial, workforce and legal risks extending beyond the period of external funding.



Recommendation

Approval is sought to secure Macmillan investment and establish a neighbourhood-based prevention model that will improve population health, strengthen communities, reduce inequalities and create a more sustainable and prevention-focused health and care system across Hywel Dda University Health Board.

It is recommended that the Board:

1. Approve the Business Case and enter into the necessary agreements with Macmillan and Social Finance to secure the proposed social investment funding and support programme mobilisation. The social investment funding will support the delivery and evaluation of the programme, while the Anchor Grant funding will be paid directly to third-sector partners to support community capacity building, neighbourhood infrastructure, community development, volunteer programmes, micro-grants and local prevention initiatives.
2. Approve the proposed five-year delivery model, recognising that whilst neighbourhood health programmes are typically funded over a three-year period, an extended five-year approach has been agreed to enable the Health Board and partners to establish the model, test and refine interventions, build evidence, demonstrate outcomes and support long-term sustainability.
3. Approve the phased rollout across neighbourhoods, allowing learning and evaluation to inform future implementation and maximise the opportunity for continuous improvement.
4. Approve the proposed programme model, comprising:
 - Health Coaching
 - Community Health and Wellbeing Workers
 - Community Capacity and Local Infrastructure for Prevention
 - Making Every Contact Count
 - Digital Enablement
5. Approve the governance arrangements, including establishment of a Programme Board chaired by the Executive Director of Public Health as Senior Responsible Owner, supported by partnership, assurance and corporate governance structures.
6. Approve the monitoring, evaluation and benefits realisation framework, ensuring robust measurement of population health outcomes, patient experience, healthcare utilisation, financial benefits and return on investment throughout the programme.
7. Request the development of a sustainability and commissioning plan during the lifetime of the programme to identify opportunities for recurrent funding and long-term integration of successful interventions into mainstream services.



8. The Programme Board should work with local authorities, primary care, community services and third-sector partners to explore and develop **pooled budgeting and joint commissioning arrangements** to support the long-term sustainability of the Community Health and Wellbeing Worker model. The programme should utilise the five-year implementation period to evaluate impact, demonstrate value across organisational boundaries, and build the evidence base required to support recurrent investment and integrated funding arrangements beyond the initial Macmillan investment period.

In the event that pooled funding arrangements cannot be agreed, the Health Board would need to identify recurrent core funding to sustain the model beyond the initial investment period. This would include funding for all three programme components, with ongoing responsibility for these costs resting with the Health Board to ensure continuity of service and delivery of anticipated benefits.

Decision Required

The Board is asked to:

- Approve the Business Case.
- Approve progression to formal funding negotiations with Macmillan and Social Finance.
- Approve the proposed five-year implementation and evaluation period.
- Approve the governance, evaluation and benefits realisation arrangements.
- Support the direct allocation of Anchor Grant funding to third-sector delivery partners to strengthen community assets, neighbourhood infrastructure and prevention capacity.

Appendices

Appendix 1. Health Coaching Current Position and Evaluation Findings

Supports: *Programme Component 1: Health Coaching*

Appendix 2. Brazil Community Health and Wellbeing Worker Model

Supports: *Programme Component 2: Community Health and Wellbeing Workers*

Appendix 3. Community Health and Wellbeing Worker (CHWW) Business Case

Supports: *Programme Component 2: Community Health and Wellbeing Workers*

Appendix 4. Demonstrator Project Evaluation Report

Supports: *Programme Component 2: Community Health and Wellbeing Workers*

Appendix 5. Communications and Engagement Plan

Supports: *Programme Governance, Community Capacity and Local Infrastructure for Prevention* and stakeholder engagement arrangements.



Appendix 6. Evidence Reviews and Literature Summary

Supports the evidence base for all three programme components and the assumptions used within the benefits realisation modelling.

Appendix 7. Macmillan GANTT Chart

Supports: *Project milestones and timelines.*

Appendix 8. Macmillan Commissioning Template

Supports: Commissioning arrangements, funding agreements, governance structures, and outcomes-based contracting frameworks.

Appendix 9: Quality Impact Assessment

Appendix 10: Health Equity Impact Assessment

Appendix 11: Equality Impact Assessment

Approval Section

Role	Name	Signature	Date
Senior Responsible Officer	Bruce Bolam	Bruce Bolam	7/08/2026
Digital Lead	Anthony Tracey		31/07/2026
Primary Care Lead	Rhian Bond	Rhian Bond	31/07/2026
Executive Sponsor	Ardiana Gjini		
Patient Representative	Abi Morgan	A Morgan	23/07/2026



Macmillan proposal: Modelling and benefits realisation

1. Executive summary

The modelling was undertaken to test whether the proposed investment could plausibly improve outcomes and moderate future healthcare demand sufficiently to support the Health Board's decision to invest. It is a rapid business-case assessment, not a prediction of guaranteed savings. The quantified results focus narrowly on avoided healthcare utilisation and do not capture the full value of improved health, wellbeing, reduced inequalities or stronger communities.

Two complementary independent modelling perspectives have been used:

- TriTech has modelled a relatively intensive Health Coaching offer, using local service assumptions and principally quantifying smoking- and alcohol-related effects on healthcare utilisation.
- The National Association of Primary Care (NAPC) has modelled a broader mixed-intensity approach across Health Coaching and Community Health and Wellbeing Workers, with intensive support for higher-need groups and lower-cost group, workforce-enabled, community and digital support for a much larger lower-intensity cohort.

The models should not be interpreted as lower and upper confidence limits for the same intervention. They examine different workforce envelopes, service designs, populations and benefit pathways. Their combined value is to identify the conditions under which the programme is more or less likely to deliver value and to inform targeting, delivery design, evaluation, gateway decisions and repayment arrangements.

Headline conclusion

Material demand avoidance is plausible, but its scale depends heavily on who is reached, the intensity and cost of support, sustained behaviour change, and the extent to which delivery evolves from predominantly one-to-one support towards proportionate one-to-few and one-to-many approaches. The modelling therefore supports approval of a controlled learning programme, not reliance on a single financial forecast.

2. Purpose, scope and decision context

The purpose of the modelling is to support an executive decision under uncertainty. It asks whether investment in Health Coaching and Community Health and Wellbeing Workers could plausibly avert future demand and influence future Health Board resource allocation. It does not attempt to value every programme outcome or substitute for prospective evaluation.

The decision standard applied is deliberately demanding. A requirement that prevention should offset its own cost is a higher threshold than the normal NHS cost-effectiveness question of whether the health gained justifies the additional expenditure. Some preventive interventions may be cost avoiding, while many others may still represent good value because they generate more health for the resources used.

For this programme, modelling is most useful when treated as part of a wider assurance system comprising service design, clinical governance, continuous improvement, evaluation, financial monitoring, gateway reviews and contractual controls.

3. The modelling approaches

3.1 Comparison of the two principal models

Feature	TriTech: intensive Health Coaching	NAPC: mixed-intensity strategy
Primary purpose	Estimate healthcare utilisation effects from a locally informed Health Coaching model.	Explore how the intensity mix and breadth of reach could affect overall return.
Modelled input	Health Coaching workforce: 17 WTE Health Coaches, supported by a clinical lead and assistants; modelled workforce cost approximately £4.6m over five years.	32 WTE across Health Coaching and CHWWs; modelled workforce cost approximately £6.8m over five years.
Delivery design	Predominantly intensive, individual Health Coaching, based on current local service assumptions.	Need-stratified Red, Amber and Green cohorts, combining one-to-one, group, MECC, community and digital approaches.
Population reach	Approximately 3,060 completed interventions a year is the core local capacity assumption.	Approximately 3,300 Red, 4,400 Amber and 55,000 Green contacts a year in the broad mixed-intensity scenario.
Benefit pathway	Principally smoking cessation and reduction in harmful alcohol use, translated into estimated healthcare activity avoided.	Top-down estimate of modifiable expenditure and potential impact across a broader programme envelope.
Reported five-year result	Avoided demand equivalent to approximately £5.2m from the reported smoking and alcohol pathways.	Avoided demand equivalent to approximately £13.2m, with substantial dependence on the assumed reach and effectiveness of the Green offer.
Key strength	Uses local pilot and service assumptions; explicitly considers high/low scenarios and relapse.	Shows how targeting and delivery intensity can materially change reach, cost and potential value.
Key limitation	Does not model CHWWs or several wider benefit pathways; some activity categories and disease-event calculations require further assurance.	The broad Green reach and effect assumptions may be optimistic; Health Coaches and CHWWs are treated too interchangeably for final evaluation.

4. What the modelling supports

- Substantial demand avoidance is plausible if the programme reaches the intended populations, achieves sustained behaviour change and adapts delivery to improve reach and productivity.
- Targeting matters. Greater short-term healthcare utilisation effects are more plausible where support reaches people with substantial, modifiable risk and potentially avoidable use of services.
- Intensity matters. A service based entirely on intensive one-to-one support limits reach; a proportionate model can preserve intensive support for higher-need groups while using lower-cost group, workforce-enabled, community and digital approaches for others.
- The quantified estimates understate the full programme value because they do not comprehensively value improved wellbeing, equity, social connection, community capacity or several clinical benefit pathways.
- Evaluation and gateway decisions are more important than apparent precision in the initial modelling because the programme is upstream, multi-component and materially different from business as usual.

Issue	Assurance implication
Attribution	Observed improvements may reflect existing services, wider policy or secular trends as well as the programme. A credible comparator and predefined attribution rules are required.
Target population	The scale of demand avoided depends on who is reached. The same behaviour change may have a different near-term utilisation effect in populations with different baseline risks and service use.
Avoided demand versus cash	Avoided activity may release capacity or moderate future pressure without creating an equivalent cash reduction in existing budgets.
Double counting	Healthcare activity categories and disease-event pathways must not be added together unless overlap has been excluded.
Breadth of benefit	TriTech principally quantifies smoking and alcohol pathways; NAPC applies broader top-down assumptions. Neither fully captures all health, equity and social outcomes.
Green cohort	The NAPC result depends materially on reaching a very large lower-intensity cohort and achieving a meaningful effect at low cost. Both reach and effect require testing.

6. How uncertainty is managed through programme design

The business case does not attempt to eliminate the uncertainty inherent in prevention. It converts uncertainty into a managed delivery and learning process. The programme will not implement a fixed service model for five years. Instead, it will use phased implementation, Plan-Do-Study-Act cycles, embedded evaluation and formal gateways to refine the target population, intervention intensity, workforce allocation and delivery channels.

It is hypothesized that Health Coaching capacity will initially provide personalised support where need and capacity to benefit are greatest. The funded Making Every Contact Count, group and digital components will progressively extend appropriate support from one-to-one towards one-to-few and one-to-many delivery. This is intended to improve reach and productivity while retaining intensive support for people with complex needs and protecting equitable access for people who cannot use digital routes.

This adaptive approach is a direct response to the modelling. It seeks to move delivery over time towards the more favourable combinations of targeting, intensity, reach and unit cost, rather than if the initial service configuration will automatically generate the modelled benefits.

7. Evaluation and gateway framework

The evaluation framework should establish a clear evidence chain from programme reach to longer-term population and service effects. Different evidence should carry different weight as the programme matures.

1. **Reach and equity:** Who is reached, from which neighbourhoods and priority groups, and whether access is equitable.
2. **Delivery and dose:** The type, intensity, duration and completion of one-to-one, group, MECC, community and digital support.
3. **Behaviour and activation:** Changes in smoking, alcohol use, diet, physical activity, patient activation, confidence and self-management.
4. **Clinical and wellbeing outcomes:** Relevant clinical indicators, wellbeing, social connection and outcomes important to participants.

5. **Healthcare utilisation:** Changes in primary, outpatient, urgent, emergency and inpatient use compared with an appropriate matched comparator.
6. **Population and system impact:** Whether intervention neighbourhoods improve relative to comparison areas, together with cost consequences, capacity effects and wider system value.

Early gateway reviews should place greater weight on mobilisation, reach, equity, delivery fidelity, behaviour change and early clinical outcomes. Later gateway and repayment decisions should place increasing weight on independently validated healthcare utilisation and financial effects. Before-and-after comparisons alone should not be treated as sufficient evidence of attribution; matched comparison methods should be used where feasible.

1. BACKGROUND

The Global Burden of Disease study estimates that around one-third of the chronic disease burden is driven by modifiable behavioural risk factors. According to the 2025 National Survey for Wales, among adults aged 16 and over: 10% are current smokers; 62% live with overweight or obesity; 41% do not meet recommended physical activity levels, and 15% consume alcohol at hazardous or harmful levels.

Without further intervention, projections from Public Health Wales suggest these behaviours will significantly contribute to a rise in non-communicable diseases over the next decade, including: a 22% increase in diabetes; a 15% increase in cancer; a 46% increase in heart failure, and a 12% increase in COPD.

Hywel Dda UHB faces rising demand, acute financial pressures, and workforce constraints that challenge the sustainability of reactive, treatment-focused models. In this context, more effective prevention and improved chronic disease management are essential to long-term system sustainability.

This direction is supported nationally. Population health and prevention are core priorities for Welsh Government and NHS Wales, alongside a growing emphasis on digital enablement and the shift to integrated, primary care-led models of care. Regionally, Hywel Dda is committed to earlier intervention, care closer to home, and a shift away from hospital-centric care towards a whole-system model, integrating health and social care and embedding the social model of health in partnership with communities, local authorities, and the third sector.

Achieving this shift will require structured, scalable approaches to behaviour change that are embedded into routine clinical care and effectively connected with existing community infrastructure – a stepped care model of behaviour change.

This proposal builds on a successful local multi-year pilot that emerged from our smoking cessation service in 2022/23 and has evolved to address a broader range of lifestyle behaviours. It provides a practical pathway to further embed prevention into and across services, teams, and communities, while aligning with regional health and wellbeing priorities.

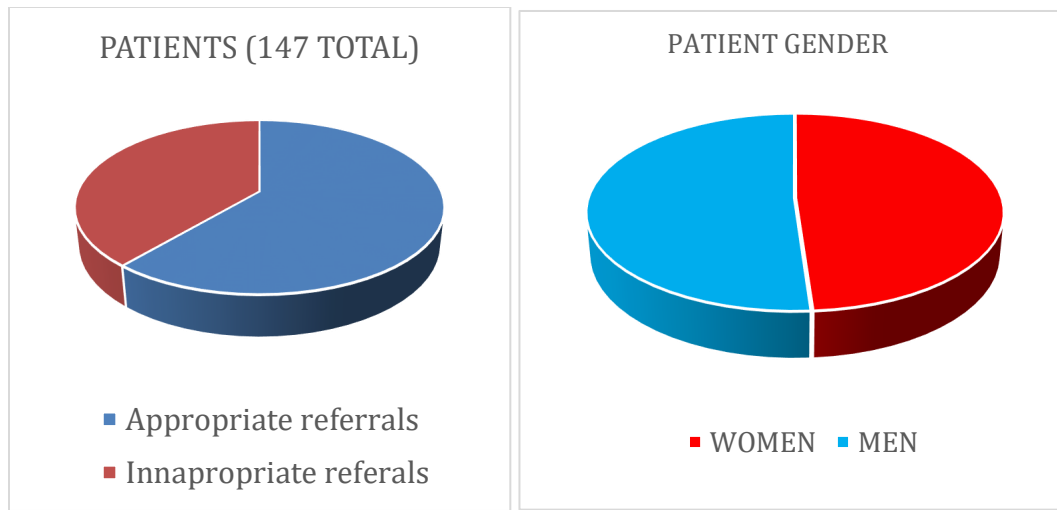
The proposed expansion will build on the successful Lifestyle Medicine pilots delivered within Hywel Dda UHB, which demonstrated the value of combining education across the six pillars of lifestyle medicine with group health coaching, community connector support, peer networks and practical workshops delivered in partnership with community and third-sector organisations. The evaluation showed that participants valued not only the education provided, but also the accountability, confidence, social connection and ongoing support generated through group-based delivery, with many forming lasting peer support networks that continued beyond the programme itself. Participants reported sustained improvements in physical and mental wellbeing,

increased confidence to engage in exercise and community activities, and meaningful behaviour change, while case studies demonstrated improvements in weight, blood pressure, blood sugar control and quality of life. The proposed model will therefore combine evidence-based lifestyle medicine education, personalised health coaching, community connector support and partnerships with local organisations to create scalable, sustainable pathways that help people move beyond managing illness towards living healthier, more connected and fulfilling lives.

2. CURRENT SITUATION

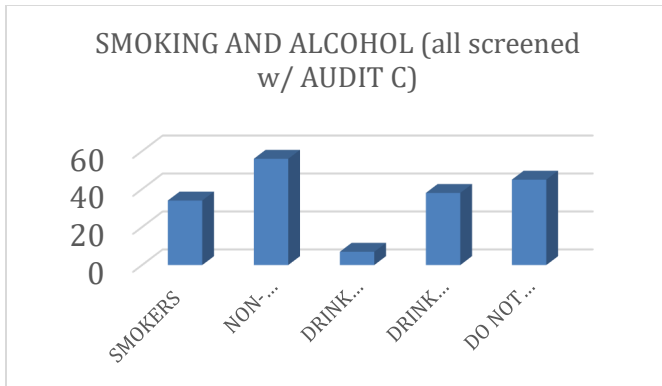
Building on the early Health Coach model, a further pilot was established in Ceredigion to begin in January 2026. This re-pilot focused on 5 GP surgeries in Ceredigion over 3 months to establish demand both from Primary Care and from the public.

During this time, there were 147 referrals into the service, of which 90 people appropriate and these individuals joined the programme.

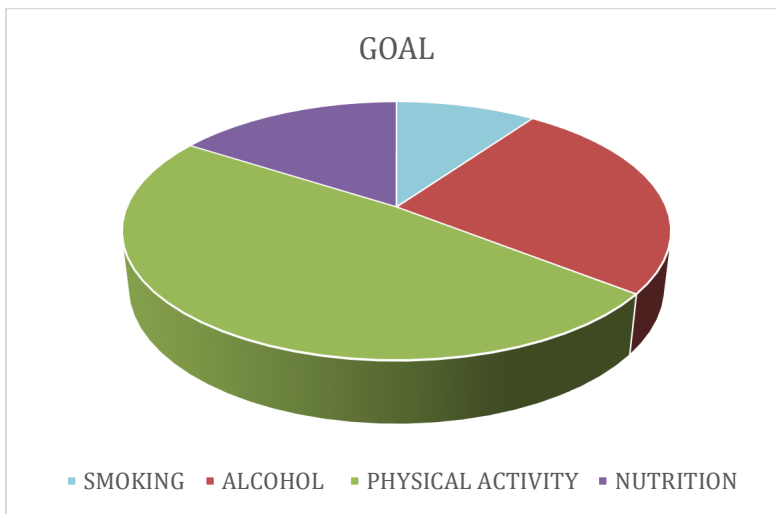


Each person received an initial assessment followed by 11 sessions of support and a range of data was collected as part of the pre-assessment including:

- Smoking status
- Alcohol Consumption (AUDIT C)
- Portions of fruit and vegetables consumed daily (also need to consider frailty markers going forward)
- Minutes of physical activity
- BMI
- Blood Pressure



Further, each individual's key motivator was explored and appropriate support and onward signposting embedded into their care plan.



The Health Coach programme, having shown the impact that it can have in such a short space of time, has been granted funding to expand. Hywel Dda has 7 primary care clusters spread across three counties. The funding will provide a country lead, a health coach for each cluster as well as project support for data, partnership working and training.

3. HEALTH COACH ENHANCEMENT BASED ON DEPRIVATION

GP Surgeries in the Most Deprived Areas of Hywel Dda University Health Board

There is not a single official Hywel Dda list ranking GP surgeries by deprivation, but it is possible can get a good proxy by mapping surgeries onto the Welsh Index of Multiple Deprivation (WIMD) areas in Carmarthenshire, Pembrokeshire and Ceredigion. The most deprived communities within Hywel Dda tend to cluster around Llanelli, parts of the Amman Valley, Pembroke Dock and Milford Haven.

Reference: <https://www.gov.wales/welsh-index-multiple-deprivation-2025>

Llanelli area (largest concentration of deprivation in Hywel Dda)

These practices serve wards/LSOAs such as Llwynhendy, Glanymor, Tyisha and parts of central Llanelli, which regularly score among the most deprived in west Wales.

- Llwynhendy Health Centre
- Ty Elli Group Practice
- Fairfield Surgery
- Avenue Villa Surgery
- Ash Grove Medical Centre

The Llanelli cluster is recognised by Hywel Dda itself as a major urban primary care locality.

Reference: <https://www.tyellihealth.co.uk/about-the-cluster>

Amman Gwendraeth

Parts of Ammanford, Garnant and Gwaun-Cae-Gurwen contain significant deprivation, particularly around former mining communities.

- Margaret Street Surgery
- Brynteg Surgery
- Amman Tawe Partnership (GCG) ((3 HC required))

Pembrokeshire

These towns contain some of Pembrokeshire's most deprived LSOAs, especially around dockland and post-industrial neighbourhoods.

- Argyle Medical Group (2 HC)
- Barlow House Surgery
- Robert Street Surgery
- Neyland

Ceredigion

Ceredigion generally has much lower overall deprivation than Carmarthenshire or Pembrokeshire. GOV.WALES notes that Ceredigion had no areas in the most deprived 10% nationally on the WIMD health domain.

- Cardigan Health Centre (GP Surgery)
- Padarn GP Surgery (Aberystwyth, student heavy population)

Reference: <https://www.gov.wales/welsh-index-multiple-deprivation-wimd-2025-results-report-health-domain-results.html>

Key caveat: Rural deprivation in Hywel Dda is often about poor access to services rather than the classic urban poverty seen in Llanelli. For example, Ceredigion and rural Carmarthenshire score very highly on access to services deprivation despite lower income deprivation.

Reference: <https://www.gov.wales/welsh-index-multiple-deprivation-wimd-2025-results-report-access-services-domain-results-html>

Enhanced Health Coach Proposal

Based on the data above, it is suggested that each of the surgeries listed has its own Health Coach, working full time in a prevention and early intervention capacity to address the lifestyle issues that contribute to chronic diseases. This would require a further 17 health coaches and, given the demand notably in Carmarthenshire, a second county lead in that area to manage the staff and workload.

Development of health coaches with an RDC setting and Dietetics

The Rapid Diagnostic Clinic for Hywel Dda (held on a Wednesday in Prince Philip Hospital) sees an average of 5 patients per week who are referred directly via their GP when they have vague symptoms of cancer which don't fit specific cancer site pathways. In clinic, patients receive a consultation, examination, scan and diagnosis (aimed to be delivered on the same day). Of these patients, less than 1 in 10 have a confirmed cancer.

Attendance at the RDC clinic and a following negative diagnosis provides a unique and timely opportunity/trigger for patients to consider adopting healthier lifestyles and reduce their risk of developing cancer. The Lead CNS for this service has observed that patients who present at the RDC often have lifestyle habits which could include smoking, alcohol, nutrition, physical activity, emotional wellbeing, sleep, etc. and could indicate that development of a future cancer diagnosis.

Lifestyle issues could be initially identified via the RDC nursing team during triage and the first clinic appointment. The 2 Clinical Nurse Specialists running the service would be trained as motivational health coaches. The Macmillan eHNA tool could be used to help record some of this information. 10 further specialist cancer nurses would also be trained to gradually implement the model across other disciplines.

Where a patient has received a negative diagnosis, the nursing team would contact the patient shortly after the diagnosis to consider addressing some of the lifestyle factors identified via the first clinic appointment. The nursing team would then signpost/refer onto suitable services (third sector, community, etc.).

This approach could also be used where a patient has received a positive cancer diagnosis. Noting that lifestyle change is harder during uncertainty, fear and stress there needs to be clear messaging “This is about supporting you while we find out what’s going on” not “you’ve caused this or done something wrong”.

A key consideration is the role of dietetics both alongside Health Coaches but more specifically, alongside those patients identified via the RDC. Whilst some patients will require weight management services, the more vulnerable may be underweight (likely if they have been referred for a cancer screening process) and appropriate support from a clinician would be vital. With this in mind, there may be more allied health professionals who might be suited to the role (such as physiotherapy or occupational therapy) so some funding should be allocated for their training in the same way as the cancer nursing staff.

TOTAL COSTS

TYPE OF EXPENDITURE	COST
Health Coaching Staff	
1 WTE Band 6 Clinical Lead	<i>£4,634,116</i>
17 WTE Band 5 Health Coaches	
3 Band 4 Health Coach Assistants	
Training Costs	<i>£92,500</i>
TOTAL	<i>£4,726,616</i>

EXPECTED PATIENT NUMBERS

This projection assumes that staff productivity, referral behaviour and patient demand remain consistent as the service expands.

Original Pilot Data

- 2 members of staff
- 147 total referrals over a 3-month period
- 90 patients seen
- 57 referrals did not progress (e.g. inappropriate, declined or otherwise not suitable)

Calculation

- Referrals per staff member over 3 months = $147 \div 2 = 73.5$
- Annual referrals per staff member = $73.5 \times 4 = 294$
- Annual referrals for 17 staff = $294 \times 17 = 4,998$

Projected Annual Activity

Measure	Calculation	Projected Annual Total
Total referrals	294×17	4,998
Patients seen	61.2% of referrals	$\approx 3,060$
Referrals not progressing	38.8% of referrals	$\approx 1,938$



OECD Reviews of Health Systems

Primary Health Care in Brazil



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Foreword

Primary health care in Brazil has a well-organised approach which is the result of sustained commitment to providing high quality primary health care for the whole population. The Family Health Strategy, launched in 1994, has been a key pillar of the efforts to reorganise and strengthen primary health care in Brazil. Since its implementation, a growing share of the Brazilian population can benefit from free access to a community-based approach to preventive and primary health care services. An extensive body of academic research has provided evidence of the key contributions of the Family Health Strategy to the reduction of almost 18 deaths per 1 000 birth in infant mortality rate observed between 1990 and 2019 in Brazil. The programme has also been associated with reduction in mortality and hospitalisations for chronic disease. Recent estimations also show that the implementation of the Family Health Strategy was associated with a reduction of 45% of the standardised hospitalisation rates per 10 000 inhabitants between 2001 and 2016, mostly for primary care conditions such as asthma, gastroenteritis, cardiovascular and cerebrovascular diseases. However, as some challenges are overcome, new ones approach, as Brazilian population ages, population risk factors such as obesity are on the rise, and emerging pandemic threats require resilience and adaptability.

Many features of the primary health care sector in Brazil deserve to be recognised as examples for OECD countries, including the use of multidisciplinary primary care teams called family health teams, and the extensive use of community health agents, which are frontline public health workers. Brazil has also implemented a set of key reforms to improve access to high quality primary health care over the past decades, including the National Programme for Improving Access and Quality of Primary Health Care (PMAQ) and the More Doctors Programme (MDP).

Nevertheless, Brazil still needs to address a number of challenges. Most critically, only 65% of the Brazilian population are covered by family health teams and too many patients bypass primary health care and directly seek care in outpatient specialties and hospitals, notably for conditions that could be more effectively treated at community level. The expansion of primary health care has also been marked by stark disparities across states and municipalities, linked mainly to workforce shortages and imbalances in the distribution of medical doctors. There are also been a rise in the number of elderly patients and an increasing prevalence of chronic non-communicable diseases – including cancer, diabetes and hypertension, which call for effective prevention, early detection and treatment. Lastly, while Brazil has a long history of developing and investing in networks, data, interoperability and skills, progress towards effective use of digital technologies in primary health care has been slow, and fundamentally unequal.

The recent reform agenda for the primary health care sector in Brazil – the *Previnha Brasil* programme – has ambitious goals toward improving access, fostering longitudinal and co-ordinated care. While primary health care is mainly a responsibility of municipalities in a decentralised health system, stronger oversight and steering from the federal government, and regional co-ordination and support from state governments will be needed to both improving access and quality, while reducing inequalities. Furthermore, as the programme is just a couple of years short of its 30th anniversary, it is in high need of innovation to get it ready for the challenges of the next 30 years, with better use of digital technology, better use of skills, modern integrated care models, backed by better performance measurements and financial incentives.

This review was prepared by the OECD Secretariat to draw on evidence and best practices from across OECD health care systems to support Brazil in strengthening its primary health care sector, and guarantee it is responsive to people's changing needs, capable of offering preventive, continuous and co-ordinated care equally across the country.

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Acronyms and abbreviations

ADAPS	Agency for the development of primary health care
APS	Atenção Primária à Saúde
ANVISA	National Agency of Sanitary Surveillance
BMI	Body mass index
BRL	Brazilian Real
CAD	Canadian Dollar
CaSAPS	Primary health care service portfolio
CFM	Brazilian Federal Council of Medicine
CGTIC	General Co-ordination of Information and Communications Technology
CHW	Community health worker
CIB	Comissão Intergestores Bipartite
CIT	Comissão Intergestores Tripartite
CQC	Care Quality Commission
CLT	Consolidation of Labor Law
CNS	National Council of Health
CME	Continuous medical education
CNES	National Register of health facilities
CONASS	National Council of State Health Secretaries
CONASEMS	National Council of the Municipal Health Secretaries
COSEMS	State Council of Municipal Health Secretariats
CPG	Clinical practice guideline
DATASUS	Informatics Department of the Sistema Único de Saúde
DENASUS	National Audit Department of Sistema Único de Saúde
DGMP	DigiSUS Gestor – Módulo Planejamento
EHR	Electronic health record
ENADE	National Exam of Student Performance
ESD 28	The Brazilian National Digital Health Strategy 2020-28
ESF	Family Health Strategy
FBG	Fasting blood glucose
FCC	Federal Communications Commission
FCM	Family and community medicine
FHT	Family health team
GESAC	Electronic Government – Citizen Assistance Service
GP	General practitioner
HPV	Human papillomavirus
HSBP	High systolic blood pressure
IBGE	Brazilian Institute of Geography and Statistics
ICT	Information and communication technologies
INCA	National Cancer Institute
INEP	Institute of Educational Studies and Research Anísio Teixeira
LAC	Latin America and the Caribbean

LDCT	Low-dose computed tomography
MACC	Modelo de Atenção às Condições Crônicas
MCTIC	Ministry of Science, Technology and Innovation
MDP	More Doctor Programme
MLA	Medical Licensing Assessment
NASF	Family health support centres
NCD	Non-communicable disease
NHS	National Health Service
OECD	Organisation for Economic Co-operation and Development
PaRIS	Patient-Reported Indicator Surveys
PACS	Community Health Agents Programme
PAHO	Pan American Health Organisation
PAPS	Planificação da Atenção Primária à Saúde
PCAT	Primary health care assessment tool
PHC	Primary health care
PMAQ	National Programme for Improving Primary Health Care Access and Quality
PLAB	Professional and Linguistic Assessments Board
PNAB	Política Nacional de Atenção Básica
P4P	Pay-for-performance
PNIS	National Policy for Health Information and Informatics
PNS	National Health Plan
QICH	National Programme for Quality Indicators in Community Healthcare
RAS	Health care networks
RNDS	National Health Data Network
RNP	National School and Research Network
RCS	Rural Clinical School
RUTE	Brazilian Telemedicine University Network
SAEM	Accreditation System of Medical Schools
SAPS	Primary Health Care Secretariat
SMART	Sistema de Monitoramento e Avaliação de Resultados do Programa Telessaúde
SBMFC	Brazilian Society of Family and Community Medicine
SINAES	National System for the Evaluation of Higher Education
SISAB	Primary Health Care Information System
SISCAN	Cancer Information System
SISCOLO	Cervical Cancer Information System
SISMAMA	Breast Cancer Information System
SOBRAMFA	Brazilian Society for Family Medicine
SUS	Unified Health System
UBS	Basic health care units
USD	United States Dollar
WHO	World Health Organization

Executive summary

The introduction of the *Sistema Único de Saúde* (SUS) in 1990 has been a major achievement for Brazil in increasing access to health care services and reducing health inequalities. The implementation of the Family Health Strategy, starting in 1994 and aimed at the reorganisation and strengthening of primary health care, has been a key component in this success. Since its implementation, the Brazilian population can benefit from free access to preventive and primary health care services delivered by multi-disciplinary family health teams (FHTs). Over the past two decades, reforms have sought to rebalance service delivery to move away from a health system that was historically very hospital-centric. Remarkably, reforms have focussed on developing modern models of care, introducing a range of quality initiatives and tools for monitoring the activities and quality of care. The expansion of the Family Health Strategy has contributed to measurable improvements in terms of infant mortality rates, maternal health, immunisation uptake and avoidable hospitalisation for chronic conditions. Life expectancy at birth increased by 5.7 years, from 70.2 years in 2000 to 75.9 years in 2019. Infant mortality rate has decreased by 60%, from 30.3 deaths per 1 000 live births in 2000 to 12.4 deaths per 1 000 live births in 2019. The implementation of the Family Health Strategy was associated with a reduction of 45% in hospitalisation rates per 10 000 inhabitants between 2001 and 2016, mostly for primary care conditions such as asthma, gastroenteritis, cardiovascular and cerebrovascular diseases. Brazil also makes spending on primary health care a high priority. In 2019, Brazil dedicated around 16% of its financial resources to primary health care, similar to OECD countries.

Despite this progress, key indicators suggest that Brazil primary health care is not working as effectively. Brazil is facing a dual challenge with still a primary health care system with marked inequalities in access and care quality, and with acute workforce shortages. The growing burden of chronic non-communicable diseases associated with the expansion in exposure to risk factors and rapid population ageing will compound existing challenges. In this context, the Review identifies scope for Brazil to building a stronger primary health care system, and to strengthen existing policies and practices to improve both access and care quality. It provides in-depth review of four priority areas: screening and prevention for major chronic non-communicable diseases, quality of primary health care provision, workforce shortages, and the digital transformation of primary health care in Brazil.

In Brazil, some cancers, hypertension and diabetes have **screening and prevention strategies**, but more could be done to improve depth and scope of such strategies. Breast and cervical cancer screening is opportunistic and does not happen within a population-based programme designed and managed at federal level. This contributes to low and uneven screening coverage. While breast cancer screening among the target group of women between 50 to 69 years of age increased from 15.2% to 24.2% between 2014 and 2019, it stands below all OECD countries and well below the 58% average in 2019. At the same time, mortality from breast and cervical cancer remains high and has been on the rise over the past decades, signalling shortcomings in terms of prevention, early detection and treatment at early stages. Between 2008 and 2019, breast cancer mortality increased from 19.2 to 20.8 deaths per 100 000 women in Brazil, compared to a decrease across OECD countries. Brazil should thus implement a strong national strategy for cancer prevention and screening, notably by moving towards population-based programmes, with a personalised approach to invite target populations. Improving data systems and health literacy are complementary strategies not to be underestimated to support stronger cancer prevention and screening.

When it comes to screening for hypertension and diabetes, Brazil should further develop disease management pathways with a people centred perspective, integrating all health care providers across different sectors. Family health teams will need to have the right tools, capacities, and incentives to undertake these responsibilities.

There are also clear shortcomings with regards to access to **high quality primary health care**. Critically, only 65% of the population are covered by FHTs and too many patients bypass primary health care and directly seek care in outpatient specialties and hospitals. These give strong argument to strengthen the gatekeeping system in Brazil with a systematic registration system with FHTs, which will control and orient the patient's into specialist care. At the same time, the federal government will need to play a more prescriptive role to strengthen comprehensiveness of primary health care evenly across the country. To embed a larger range of activities and effective interventions in FHTs, it will be critical to ensure that all municipalities are able to deliver actions and services properly. Introducing economic incentives, alongside appropriate educational programme in prevention, detection, and treatment are key approaches to consider. There are also several opportunities for extending further quality strategies toward primary health care. Particular attention should be given to transparency and the collection of richer performance data. A robust accreditation system that would apply uniformly across the country could help standardise primary health care quality, and to identify areas that may require greater financial and organisational support. The new Agency for the Development of Primary Health Care could act as an inspectorate for primary health care, for example to provide independent and external verification that standards are being met, identify good practice and support weaker centres to improve their standards.

When it comes to **workforce challenges**, Brazil primary health care sector faces acute shortages and imbalance in the distribution of primary health care doctors with a North-South gradient, where North and Northeast regions show a lower density of medical doctors. In addition, little attention has been devoted to building a strong credibility of the primary health care specialty, and to promote workforce quality. In 2018, only 5 486 medical doctors had a specialty training in family and community medicine, representing 1.4% of all specialists in Brazil. Going forward, Brazil needs to implement a coherent workforce planning based on an objective assessment of present and future needs to govern health care human resources, which is currently absent. In tandem, Brazil could look at the experiences of OECD countries to train more rural doctors and to provide both wage and non-wage related financial incentives linked with a return of service obligation. Brazil should take steps to expand the role of nurses and community health workers and explore the potential of task delegation. Perhaps more crucially, there are opportunities to make the primary health care specialty training compulsory for all doctors wishing to practise primary health care, and to implement stronger requirements around continuous medical education.

Brazil's decentralised government creates challenges to digitalising primary health care. Brazil has made strides towards a **digital transformation** of primary health care, building on more than a decade of policies to digitalise health care and make better use of health data, and with key investments in networks, data, interoperability and skills. Yet, progress towards effective use of digital primary health care has been slow, and fundamentally unequal, with significant inequalities in the use of digital technologies and tools among health workers and citizens. In 2019, an estimated 78% of PHC units had electronic health record (EHR) systems. Adoption of EHR systems was highest in the South and Southeast regions (at 90% and 83% of facilities in those regions, respectively) compared to the Northeast and North regions (77% and 74% respectively). Other major barriers include human and technical capacities in municipalities, with potential diseconomies of scale resulting from setting responsibilities for digital health at municipal level. Tackling these challenges requires to digitalising all primary health care units and teams, while promoting inclusive connectivity for all Brazilian citizens, especially the most vulnerable. Brazil should also consider establishing a governance structure with clear well-funded mandates at the right levels of government.

Across all these areas, there is scope to strengthen oversight from the federal government and regional co-ordination and support from state governments. Better and stronger co-ordination mechanisms between levels of government have the potential to lessen regional inequalities, and prepare the health system

against systemic shocks, such as the one observed in the COVID-19 pandemic. The Brazilian unified health system (SUS) has had high aspirations since its introduction by the Federal Constitution of 1988, which was the result of a society-wide mobilisation. Primary health care has been a cornerstone of the extension of health care coverage since the beginning of SUS. Continuing to strengthen and modernise primary health care remains the essential lever for the Brazilian health system to realise its potential of effectively achieving universal coverage of high quality services for all Brazilians.

1

Primary health care in Brazil: Assessment and recommendations

Primary health care (PHC) in Brazil has a well-organised approach, the result of sustained commitment to providing high-quality PHC for the entire population. Brazil has introduced a comprehensive package of policies designed to modernise and strengthen PHC. However, a number of key challenges remain, including the increasing prevalence of chronic non-communicable diseases, large disparities across regions in the supply and quality of PHC, and a low and unequal use of digital technologies. This chapter assesses Brazil's PHC system. It considers four topics in detail: screening and prevention for major chronic non-communicable diseases, enhancing the quality of PHC provision, tackling workforce shortages and promoting the digital transformation of PHC in Brazil. It provides a set of recommendations on improving PHC in the country.

Brazil has a relatively well-functioning and well-organised primary health care (PHC) system, the result of sustained commitment to providing high-quality PHC for the whole population. The Family Health Strategy (Estratégia de Saúde da Família [ESF]), one of the largest community-based PHC programmes in the world, has successfully increased population coverage and improved key health outcomes. Since the strategy's launch in 1994, the Brazilian population has enjoyed free access to preventive and PHC services, delivered by multidisciplinary family health teams (FHTs) (Box 1). The expansion of the Family Health Strategy from 1994 has contributed to measurable improvements in infant mortality rates, maternal health, immunisation uptake and avoidable hospitalisation for chronic conditions. For example, infant mortality rates decreased from 30.3 deaths per 1 000 live births in 2000 to 12.4 deaths per 1 000 live births in 2019. Life expectancy at birth increased by 5.7 years between 2000 and 2019, from 70.2 years to 75.9 years. In addition, the Requalifica programme launched in 2011 has led to major public investment to expand the supply and equipment of PHC facilities. Still today, investment in PHC is a high priority for the federal government: in 2020, it allocated a budget of BRL 20.9 billion (Brazilian real) to PHC (USD 3.8 billion), compared to BRL 17.5 billion in 2019 (USD 3.2 billion) (CONASEMS, 2020^[1]). In the context of the COVID-19 pandemic, the federal government provided punctual financial and organisational assistance to FHTs to manage COVID-19 cases, and non-COVID-19 patients. These included for example the development of special COVID-19 care centres, the development of safety standards and protocols to manage high-risk patients.

Despite this progress, important challenges lie ahead for Brazil. Only 65% of the population is covered by FHTs. Too many patients bypass PHC, directly seeking care in outpatient specialised clinics and hospitals, notably for conditions that could be more effectively treated at the community level. This is partly the result of structural challenges that make it difficult for FHTs in some municipalities to deliver PHC functions. PHC has traditionally provided few low-complexity procedures and surgeries. In addition, the expansion of PHC has been marked by stark disparities across regions and municipalities. The most vulnerable and remote municipalities of the North and Northeast regions consistently present poorer health outcomes and lower health care quality than the national average. The proportion of hospitalisations owing to conditions that could be more effectively treated in the PHC setting ranges from 24% in the South region to 40% in the North region.. Such disparities stem in part from workforce shortages and imbalances in the distribution of medical doctors in rural and remote areas. Not only does Brazil have too few PHC physicians, but they are also highly concentrated in the South and Southeast regions. Last but not least, as in all OECD countries, Brazil is experiencing a rise in the number of elderly patients and an increasing prevalence of chronic non-communicable diseases, including cancer, diabetes and hypertension (see Box 1). These call for more effective prevention and stronger PHC to manage evolving health needs, and ensure a healthy and active population across the country.

Box 1. Key features of the primary health care sector in Brazil

The Brazilian *Sistema Único de Saúde* (SUS), instituted in 1990 after the approval of the new Brazilian Constitution, is decentralised. The Ministry of Health is responsible for the central management of the health system, and **the 5570 municipalities are responsible for the organisation, management and delivery of PHC services**. With federal funding, municipalities have the freedom to hire family health team members according to the population needs and geographic characteristics.

The federal government finances one-third of total PHC costs (33%), with the remaining part being funded predominantly by municipalities (61%). The participation of states is the smallest. **Brazil dedicated around 16.3% of its current health expenditure to PHC**, slightly lower than OECD average (16.6%).

The Brazilian SUS offers preventive services and PHC free of charge to its population. The private sector is supplemental as health plans mainly offer specialist and hospital services, although private health plans have begun to offer access to family doctors.

Multi-professional family health teams (FHTs) are responsible for the delivery of PHC. Each FHT is assigned a geographic area, covering up to 4 000 individuals located in non-overlapping areas. FHTs are responsible for registering every family in their geographic area. As of 2019, there were 43 234 Family Health Teams in Brazil. The FHT includes physicians, nurses, and up to 12 community health workers. Other PHC workers can provide additional supports depending on local needs.

FHTs are expected to be the initial contact point for patients having new health needs, although it does not work as a “gatekeeper” and many Brazilian have direct access to outpatient specialties and hospitals. In addition, only about 65% of Brazilians are currently registered with a FHT.

In 2020, there were around 60 000 medical doctors working in PHC, alongside 50 000 nurses and 35 000 dentists in practice in Brazil. Practicing doctors per 1 000 population in 2019 were lower compared to the OECD average, at 2.3 compared to 3.5. While the specialty of family and community medicine was introduced as a clinical and licensed specialty, PHC is mostly delivered by physicians who do not have any specialty or by physicians with a specialty in internal medicine. In 2018, only 5 486 medical doctors had a specialty training in family and community medicine (1.4% of all specialists).

There is a marked disparity in the concentration of doctors in different regions and states, and between state capitals and other parts of states. The Southeast is the region with the highest medical density per inhabitant (with 2.81 per 1 000 population), against 1.16 per 1 000 population in the North and 1.41 in the Northeast.

Table 1.1. Key performance indicators in Brazil and OECD, 2019 (or latest year available)

	Brazil	OECD average
Spending in PHC (% of current health spending)	16.3%	16.6%
Doctors (per 1 000 pop)	2.3	3.5
Nurses (per 1 000 pop)	8	9.1
Life expectancy at birth (years)	75.9 years	81 years
Infant mortality (deaths per 1 000 live births)	12.4	4.2
Smoking (daily smokers, % population aged 15+)	9.8%	16.5%
Alcohol (litres consumed per capita, population aged 15+)	6.1	8.7
Overweight prevalence (age-standardized %)	56.5%	58.4%
Raised blood pressure (% of the pop)	23.3%	20.8%
Raised fasting blood glucose (% of the pop)	8.3%	6.7%
Vaccination rates for DTP 3 (% of children)	83%	94.8%
Breast cancer screening (% of women aged 50-59)	24%	58%
Cervical cancer screening (% of women aged 20-69)	37%	59%
Hospital admission due to CHF and hypertension (Age-sex standardized rate)	167 per 100 000 pop	301 per 100 000 pop
Hospital admission due to diabetes (Age-sex standardized rate)	92 per 100 000 pop	130 per 100 000 pop

Faced with these significant challenges, Brazil has introduced a comprehensive package of policies to modernise and strengthen PHC. Implemented in 2013, the More Doctors Programme (MDP) is considered the world’s largest government-led initiative recruiting domestic and foreign physicians to work in PHC settings within municipalities facing difficulties in recruitment and retention. By November 2020, the MDP was responsible for the presence of 16 426 doctors in 3 837 Brazilian municipalities. It has also yielded positive results in terms of investment in PHC facilities, physician availability and health outcomes (Hone et al., 2020^[2]; Netto et al., 2018^[3]).

Reforms have also focused on developing new models of care, primarily through family health support centres (*Núcleo Ampliado de Saúde da Família* [NASF]), which function as integrated multidisciplinary teams providing a broad range of services to the community. Health care networks (*Redes de Atenção à Saúde* [RAS]) have been developed to integrate health care across different sectors and improve patient-centred care in the country. The National Programme for Improving Primary Health Care Access and Quality (PMAQ) has been a strong tool for monitoring and improving the performance of PHC facilities. Finally, the *Previne Brasil* programme, introduced in 2019, has set ambitious goals, such as introducing a new financing scheme based on a weighted capitation model, a pay-for-performance component and incentives for strategic actions. The weighted capitation payment and associated patient registration system are welcome steps in building a profile of population health needs. The *Previne Brasil* programme has the potential to improve access to PHC and promote longitudinal, co-ordinated care while reducing inequalities between regions. But Brazil still has a way to go before the fruits of this reform are truly felt across the PHC system. First, the country must address several key persisting challenges across the PHC sector:

- Brazil's ambitious reform agenda must be accompanied by a stronger central oversight and a focus on improving health care quality and supporting the most vulnerable municipalities. Central agencies need to play a greater role in this respect, as poorer municipalities have weaker infrastructures and reduced capacities to deliver care than the best-performing areas. At the same time, Brazil needs to underpin quality monitoring and improvement with adequate mechanisms to ensure implementation at all municipal levels; this requires cross-governmental collaboration and co-ordination.
- Brazil should explicitly recognise the central role of FHTs in the prevention, treatment and management of chronic non-communicable diseases, such as diabetes and hypertension. Patients' ability to access specialist care without prior consultation with a FHT suggests that the potential of PHC to lead in health promotion, disease prevention, treatment and follow-up is not being fulfilled.
- Some key quality strategies are underdeveloped or absent from Brazil's PHC sector, including establishing requirements for accreditation, continuous medical education (CME) and compliance with clinical guidelines. Currently, payment systems only timidly reward improvements in the quality of care. Overall, Brazil lacks a support system to ensure continuous improvement in health care quality at the team or facility level.
- Despite the existence of numerous data sources, Brazil's health information systems are insufficient to provide a full picture of the effectiveness and safety of PHC. Most indicators are input and process indicators, such as the number of FHTs, the number of people registered with a FHT, the number of community health workers, or the number of pregnant women receiving prenatal care services. Regarding cancer care, there is no integrated information system based on a registry that allows assessing the effectiveness of current cancer screening protocols and cancer care delivery.
- Little attention has been devoted to enhance the credibility and recognition of the PHC specialty, as well as promote workforce quality. Within PHC, the specialty of PHC (called family and community medicine [FCM]) is the least preferred specialty among physicians. In addition, geographical distribution of medical doctors is inequitable, with acute workforce shortages in rural and remote areas.
- Brazil has made significant inroads in building a foundation for digital PHC, including essential investments in networks, data, interoperability and skills. However, progress in implementing digital technologies in PHC has been slow and fundamentally uneven across the country.

This chapter makes a detailed assessment of Brazil's PHC system and proposes a set of recommendations for improving it. It considers four topics: 1) enhancing the quality of PHC provision,

2) increasing the screening and prevention of major chronic non-communicable diseases, 3) resolving workforce shortages, and 4) advancing the digital transformation of PHC in Brazil.

Policy recommendations for improving primary health care in Brazil

Improve the quality of PHC provision

- Strengthen the gatekeeping system with systematic registration of patients with FHTs, which guide patients into specialist care as needed through a referral system.
- Encourage greater comprehensiveness of PHC by ensuring that all municipalities are able to deliver the actions and health care services defined in the Primary Health Care Portfolio, and encourage compliance with clinical guidelines. Financial incentives could be introduced through existing contracting schemes or the *Previne Brasil* programme.
- Develop educational programmes in disease prevention and early diagnosis through formal frameworks and requirements to undertake CME programmes.
- Collect a richer set of quality indicators on a wider array of preventive activities and management of chronic conditions (such as alcohol consumption, obesity, cancer screening and mental health), including quality of life and patient experiences with PHC; implement a public reporting system to allow patients to monitor the performance of individual FHTs.
- Build accreditation systems around standards of care for all FHTs in order to assess their performance, identify areas that may require improvements and provide support for such improvements. The new Agency for the Development of Primary Health Care could carry out accreditation and provide independent verification that standards are being met, ensuring continuous improvement in health care quality.
- Expand health care networks to prioritise the provision of integrated care for patients with multiple needs. Additional resources, guidelines on establishing and running community care services, and a better use of digital technology are all areas that would benefit from greater steering and oversight by the federal government.

Improve cancer screening coverage and promote early detection of cancer

- Move from opportunistic screening programmes toward population-based screening programmes for breast and cervical cancer, implementing a more personalised approach (e.g. by sending personalised screening invitations to high-risk populations).
- Develop more effective communication strategies and education programmes to improve health literacy and promote the importance of early detection among people with different socio-economic and cultural backgrounds.
- Develop and expand national registries for breast and cervical cancer in order to identify and invite people with a high risk of cancer; monitor and evaluate the effectiveness of screening programmes and cancer care delivery.
- Facilitate deeper analysis of diabetes and hypertension care through a stronger data infrastructure allowing a more comprehensive set of quality measures and feeding back data to FHTs to improve the quality of care.
- Promote the development of disease-management pathways with a people-centred perspective, integrating all health care providers across different sectors; ensure that FHTs have the skills and capacity for screening and management of hypertension and diabetes through expanding the role of other health workers, including nurses and pharmacists.
- Collect and report information on cancer screening at the individual patient level to improve the effectiveness of existing cancer screening programmes and protocols.

Address workforce shortages and reduce imbalance in the distribution of primary care workforce

- Define a clearer workforce management policy to govern human resources for health, based on an objective assessment of present and future needs.
- Increase the number of trained doctors in rural areas to encourage recruitment and retention of medical doctors in underserved areas; provide both wage- and non-wage related financial incentives linked to a service obligation.
- Address workforce gaps by changing the scope of practice for nurses and community health workers so that they help meet health care needs in more remote areas; establish a governance structure that better supports municipalities with insufficient capacities.
- Support the creation of academic FCM departments in all Brazilian medical schools to undertake research in PHC, conveying the specialty's value and promoting it among students.
- Improve the attractiveness of the PHC profession by standardising contractual arrangements for primary care workers. Ensure a more consistent salary scale across the country through stronger oversight and regulation by the federal government, notably using the new Agency for the Development of Primary Health Care
- Introduce a national test at the end of the university period to help standardise the medical curriculum in Brazil, complementing the current accreditations.
- Make specialty training in FCM compulsory for all medical doctors wishing to practise PHC; establish formal requirements for CME and recertification to secure a high-quality workforce. CME should be directed towards key challenges, such as the prevention of risk factors for health and management of chronic conditions.

Advance the digital transformation of PHC

- Set ambitious and clear targets for both reliability and speed of connections as part of existing and future collaborations between the Ministry of Health, other ministries and the National School and Research Network (Rede Nacional de Ensino e Pesquisa [RNP]) to increase Internet connectivity in PHC units.
- Expand the financial payments included in the Connect SUS pilot project in Alagoas to the entire Brazilian territory; condition receipt of financial support to participation by municipal and state officials in capacity-building workshops; develop training specifically targeting senior executives.
- Set ambitious and clear quantitative targets for adoption and use of EHR systems (such as e-SUS AB). Such targets should include the effective use of electronic drug prescribing and the exchange of clinical and administrative data across units.
- Strengthen the role of telecentres as training providers, especially in rural and remote areas, and ensure appropriate funding and technical assistance at all levels of government; develop mobile access to reach those vulnerable groups that are most likely to need PHC.
- Consider aggregating shared services and responsibilities related to digitalisation of PHC at less decentralised levels; Set up dynamic purchasing systems at the federal level, with appropriately funded agencies at the state or regional levels providing municipalities with specialised human resources, purchasing and management services for digital PHC.

1.1. Improving Brazil's primary health care quality and outcomes

1.1.1. Brazil has introduced a number of initiatives to improve the quality of care

Brazil has implemented a set of key reforms over the past decades to improve access to high-quality PHC and develop new models of PHC, primarily through family health support centres. These centres give FHTs greater flexibility in managing their practice according to population health needs, ensuring more co-ordinated and integrated care. The introduction of the health care networks (*Redes de Atenção à Saúde* [RAS]) is also an innovative approach to delivering patient-centred care. The overarching objective is to integrate the organisation, planning and provision of health care services (from PHC, mental health, public health and emergency care, to specialised care and hospital care) at the regional level. RAS are agreed by municipalities and states through interagency commissions (*Comissões Intergestores*) according to population health needs, and local capacities.

A range of quality initiatives have also been introduced since the implementation of the Family Health Strategy in 1994, including the Requalifica programme to expand the availability and equipment of PHC facilities, the PMAQ, and the MDP to recruit and retain PHC professionals. Among the new tools for monitoring the activities and quality of care, the Primary Health Care Information System (SISAB) gathers an impressive number of indicators underpinning PHC, but mostly focuses on input and process indicators such as the number of FHTs, the number of people registered with a FHT, the number of community health workers and the number of pregnant women receiving prenatal care services.

Some clinical guidelines exist. The Ministry of Health has issued several “primary health care notebooks” (*Cadernos de Atenção Básica em Atenção Primária*) to help FHTs improve the delivery of high-quality PHC (Ministério de Saúde, 2021^[4]). The notebooks address specific diseases or risk factors, such as care for smokers, arterial hypertension or mental health. While such guidelines are useful, the PHC notebooks are still too narrowly defined and do not meet patients’ expectations for integrated health care. Furthermore, information about their effective use is lacking. Implementation is the responsibility of the municipalities, but there is no mechanism at the federal level to monitor compliance, and no systematic incentives for guideline uptake. Given inadequate follow-up observation of diabetes control and blood pressure control for people with hypertension, compliance with clinical practice guidelines is arguably unsatisfactory (Leite Simão et al., 2017^[5]; da Silva Rêgo and Radovanovic, 2018^[6]). In general, Brazil lacks system-wide support for ensuring continuous improvement in the quality of health care at the team or facility level.

Brazil has also recently committed significant efforts to reforming its federal financing model. The new PHC funding model, *Previne Brasil*, came into effect in 2020. It includes a weighted capitation component, a pay-for-performance programme, and some incentives for strategic actions and priority areas. The new capitation formula, which allocates resources based on needs, is well-aligned with OECD member countries’ policies in this area (e.g. in Chile, Israel, Portugal, New Zealand, the United Kingdom and the Netherlands). The single capitation amount will be applicable to the population registered with FHTs and adjusted for socio-economic, demographic and geographical factors, providing an incentive for FHTs to register the 50 million people who are not yet registered. This is a welcome step to better identify the people linked to each FHT, and encourage more appropriate longitudinal and co-ordinated care. Importantly, the weighted capitation component will ensure that municipalities with higher population needs owing to older populations and less favourable socio-economic situations receive the necessary resources to provide care to everyone. This will limit the risk of increasing inequalities in access to and quality of PHC services.

1.1.2. Despite progress, some concerns about quality persist, and primary health care struggles to deliver its essential functions

Prevention and treatment at the PHC level can still improve. International figures show that breast and cervical cancer screening can progress, and immunisation rates could be higher (Table 1.1). In 2018, 84% of Brazilian children around one year old were vaccinated against measles, below the OECD average (95%). At the same time, simple procedures and surgeries need to be made more widely available in PHC settings. This is both a matter of striving for better safety and care quality, and an economic necessity to avoid unnecessary use of expensive hospital services. Available data show that only 60% of FHTs performed low-complexity procedures or small surgeries – too few if Brazil wants to expand and strengthen the role of PHC.

Inappropriate preventive care and management of chronic diseases has also been highlighted by previous researches. A number of worrying indicators suggest an urgent need for better prevention, especially at a time when the burden of disease is shifting towards chronic diseases. According to the last National Health Survey, 21% of all Brazilians with diagnosed diabetes did not receive medical care in the 12 months prior to the survey. In addition, only 22% of SUS users indicated they had received some recommendations on healthy lifestyles from their family doctor, and less than 40% of public users had a preventive visit in the past two years (Guanais et al., 2019^[7]). Significant differences in health care quality also exist across regions, with a north-south gradient where North and Northeast regions show higher hospitalisation rates for chronic conditions. The 2019 National Health Survey shows for example that diabetic patients have higher hospitalisation rates for chronic conditions in the Northeast than in the Southeast, raising some concerns about the effectiveness of local PHC in the North and Northeast regions.

1.1.3. Renew efforts to strengthen key functions of PHC, particularly through stronger gatekeeping and more comprehensive care

Despite clear investment in PHC, too many patients use public speciality units or emergency departments to receive health care services. This is a strong argument for strengthening the gatekeeping system in Brazil, both by enforcing systematic registration with FHTs and establishing a referral system to better control and orient patients into specialist care. This approach is taken internationally, including in Chile, Portugal, Italy and Norway, where strong gatekeeping systems are seen as a way to ensure that patients receive the best possible care for their conditions, achieving greater appropriateness and co-ordination of care. Brazil may consider such a reform in the future to expand the role and expansion of FHTs. This would help build accurate patient registers recognising the central co-ordinating role of FHTs. The implementation of the new *Previne Brasil* programme is a step in the direction of increasing patients' registration with FHTs.

In addition, the scope of PHC services needs to expand to ensure that FHTs are able to deliver essential functions, including continuous and comprehensive care focused on prevention, treatment, diagnostic follow-up and management of chronic conditions. A large range of activities and interventions should be systematically embedded in FHTs. This more comprehensive approach should be underpinned by the delivery of actions defined in the Primary Health Care Portfolio (called CaSAPS), while ensuring that all municipalities are able to deliver such actions and services properly. Incentives for primary care providers to deliver these actions could be introduced through existing contracting schemes with the municipality or the incentive structures outlined in the strategic actions of the *Previne Brasil* programme. The Ministry of Health could also use the *Previne Brasil* pay-for-performance programme to monitor and encourage more systematic compliance with clinical guidelines, in order to embed more effective prevention in FHT practice. At the same time, educational programmes covering the prevention, detection, treatment and management of diseases are key instruments to encourage primary care professionals to fully implement these tasks. Several bodies, such as the National Council of Health Secretaries (CONASS), the National Council of Municipal Secretaries of Health (CONASEMS), the Brazilian Medical Association and the Brazilian Society of Family Medicine, organise trainings, conferences and online learning.

However, a more formal framework for CME that clarifies expectations and helps FHTs meet objectives is required to provide more comprehensive care, in line with the best available evidence.

1.1.4. *There exist several opportunities to extend quality strategies targeting PHC*

A robust accreditation system, which would apply uniformly and consistently to the primary health care sector nationwide, could be a tool for improving the quality and performance of primary health care in Brazil. Other federalised health systems in the OECD (e.g. in Australia, Canada and the United States) have established PHC accreditation systems. Building an accreditation system for all FHTs in Brazil would make it possible to assess their performance, identify areas for improvement and provide support for such improvement. Since Brazil does not have a National Inspectorate for Health to provide accreditation and independent verification that standards are being met, the new Agency for the Development of Primary Health Care could take on this role. It could, for example, undertake quality monitoring and improvement, and review current practice and performance, and setting standards for performance and reporting. It could also develop tools such as evaluation frameworks, deploy teams to visit and support municipalities with special needs, and redistribute resources when needed. The agency could then be responsible for ensuring continuous improvement in health care quality at the local level, which is currently not – or only timidly – in place.

Perhaps more crucially, a greater transparency and richer set of performance data are needed to drive quality improvement. Brazil already collects a wide range of inputs and process indicators. As part of the new *Previne Brasil* pay-for-performance programme, the country also monitors several indicators on access to PHC, risk factors and quality covering maternal health, child immunisation, breast cancer screening, and management of hypertension and diabetes. While this is an important initiative for quality monitoring, five quality indicators are process indicators, and two indicators are intermediate outcomes measures. This is insufficient. A richer set of quality indicators on a wider array of preventive activities and management of chronic conditions, such as alcohol consumption, obesity, cancer screening, mental health and patients' experiences with PHC, should be a priority and will be critical to ensure that ongoing reforms do not adversely affect quality and outcomes of care. In this respect, Portugal and Israel offer models of comprehensive, actionable indicators supporting quality improvement in primary care.

Brazil might also want to join the OECD Patient-Reported Indicator Surveys (PaRIS) initiative to understand how the outcomes and experiences of care in the country compare with OECD countries. Under PaRIS, countries work together on developing, standardising and implementing a new generation of indicators that measure the outcomes and experiences of health care most important to people. In this regard, the International Survey of People Living with Chronic Conditions under PaRIS will be the first of its kind to assess patients managed in PHC across countries.

1.1.5. *Integrated care needs better central support and direction*

Given the challenges brought by demographic and epidemiological changes, Brazil urgently needs to shift the focus of health care services to addressing longer episodes of health care needs. While the country is on the right track with the development of the RAS health care networks, this integrated care model is still unevenly implemented nationwide. The challenge should not be underestimated: interactions between different health care providers about patient cases and transitions from one service to another need to be timely, safe and seamless for patients and families. PHC play a key role in achieving this goal.

A lack of guidance and national leadership has resulted in a low and uneven diffusion of RAS across the country. The federal government should consider playing a greater role in steering a more consistent development of RAS, providing additional resources, training programmes, and guidelines on developing and running community care services, and better using information and communication technology (ICT).

Exchanging good experiences and learning from the top-performing regions or facilities is another way to encourage more extensive and ambitious development of these networks.

Better information-sharing and data linkage across care levels, as well as smarter provider-payment mechanisms will be key to ensure a successful rollout of RAS. There exist good examples of payment mechanisms – such as add-on payments, bundled payments and population-based payments – in some OECD countries that incentivise providers from different health sectors to work together effectively. While it may be too early to institute such mechanisms at the moment, Brazil could certainly consider introducing add-on payments or bundled payments in the near future.

1.2. Health screening for chronic non-communicable diseases in Brazil

In Brazil, as in OECD countries, chronic non-communicable diseases (NCDs) such as cancer and cardiovascular diseases are major public health concerns. Cancer mortality is the second cause of death in Brazil. The leading cause of cancer mortality is breast, lung, colorectal and cervical cancer among women, and lung, prostate and colorectal cancer among men. In keeping with trends across OECD countries, cancer mortality rates in Brazil are higher among men than among women, which can be explained at least in part by a higher prevalence of risk factors among men. Many cancers associated with high mortality rates in Brazil – such as prostate, breast, colorectal and lung cancer – feature common risk factors, including obesity, poor lifestyle and physical inactivity.

While the incidence of cancer in Brazil is below the OECD average, cancer mortality has not decreased in recent years. Between 2009 and 2019, Brazil's cancer mortality oscillated between 161 and 162 deaths per 100 000 people, while the OECD average decreased from 211 to 196 over the same period. The decrease in cancer mortality across OECD countries is related to early diagnosis, which makes a strong argument for building more effective screening schemes in Brazil.

The fifth cause of mortality in Brazil is diabetes, which accounts for 5% of all deaths compared to 3% in OECD countries. Diabetes mortality in Brazil reached 50 deaths per 100 000 people in 2019 – higher than the OECD average of 22 deaths per 100 000 people, and only lower than Mexico and Costa Rica. Fasting blood glucose levels significantly increased over the past decade, suggesting that the burden of diabetes will be significant in the near future. Hypertension, which contributes to the development of more serious cardiovascular problems, is a major risk factor for poor health in Brazil, if not the main one. In 2015, the average prevalence of raised blood pressure in Brazil was 23.3% – higher than the OECD average of 20.8% and the Latin America and the Caribbean average of 21.5%. While Brazil reduced this prevalence by 5.3% between 2005 and 2015, OECD countries decreased it by an average of 16.4% over the same period.

1.2.1. Improving cancer screening and prevention in Brazil

While Brazil has expanded its cancer care system, population coverage remains low

To tackle the burden of cancer, Brazil has developed its cancer care system incrementally over the past decades. It launched the first National Oncology Care Policy in 2005 and set its Strategic Action Plan for Coping with Chronic Non-Communicable Diseases 2011-22, demonstrating its commitment to reducing the burden of cancer. It introduced the National Policy for the Prevention and Control of Cancer in the Health Care Network of People with Chronic Diseases in 2013 to update the national cancer policy and improve access to cancer care. Brazil has also sought to implement quality improvements in its cancer care system by establishing in 2013 the Cancer Information System (SISCAN), which integrates the Cervical Cancer Information Systems (SISCOLO) and the Breast Cancer Information Systems (SISMAMA). Clinical guidelines for early detection of breast cancer were updated in 2015, and national

guidelines for cervical cancer screening in 2016. More importantly, Brazil introduced the PAINEL-Oncology instrument, which helps monitor Law No. 12.732 of 22 November 2012 establishing a 60-day maximum waiting time for the start of treatment for patients with proven malignant neoplasia. While PAINEL-Oncology is key to guarantee access to high-quality cancer care in Brazil, the data presented refer exclusively to users who have a national health card (DATASUS, 2021^[8]). Brazil has also developed 17 population-based cancer registries, which provide crucial information about incidence, trends and survival (INCA, 2013^[9]).

However, cancer screening in Brazil is opportunistic, managed by each FHT as a result of a recommendation made by a health care provider during a routine medical consultation, or through individual self-referral. Brazil needs to step up its efforts to increase breast cancer screening coverage, notably to improve early diagnosis and reduce mortality. Unlike in many OECD countries, cancer screening does not occur within a population-based programme designed and managed at the central level to reach the bulk of the population at risk. As a result, while coverage for breast cancer screening in Brazil increased from 15.2% in 2014 to 24.2% in 2019 among the target group of women aged 50 to 69 years, it remains very low compared to the OECD average of 58%. Between 2008 and 2019, breast cancer mortality increased by 8.3% in Brazil, whereas it decreased by 10% on average across OECD countries. Breast cancer takes the greatest toll on Brazilian women, with 20.8 deaths per 100 000 females, representing 16.4% of all women's cancer deaths. Moreover, breast cancer survival dropped by more than two years in Brazil between the 2005-09 and 2010-14 periods, whereas it increased by one year in the OECD region. The lack of availability of mammographs, and their uneven distribution across the country, are major problems contributing to late diagnosis of breast cancer in Brazil (Da Costa Vieira, Formenton and Bertolini, 2017^[10]). In 2020, Brazil had 13 mammographs per million people, almost half the OECD average of 24.

In a similar vein, cervical cancer screening in Brazil has substantially risen (from 20.5% in 2014 to 37% in 2019), yet it remains well below the OECD average coverage (57%), and below all OECD countries except Costa Rica and Hungary. Worrying evidence points to large social inequalities: access to cervical cancer screening is particularly low among women living in the North, Northeast and Central-West regions, and those with low socio-economic backgrounds (including poorer, less educated, non-white and younger women) (Costa et al., 2018^[11]). Cervical cancer mortality in 2019, for its part, was substantially higher in Brazil than the OECD average (7.4 deaths vs. 3.9 deaths per 100 000 women), and cervical cancer survival in Brazil decreased by 11.3% between 2000-04 and 2010-14.

These worrisome trends suggest large room for improvement in Brazil, including through a national strategy for cancer prevention and screening.

Towards population-based screening programmes using a more systematic and personalised approach

In the area of prevention of cervical cancer, the National Immunisation Programme instituted in 2014 the human papillomavirus (HPV) vaccination campaign for girls aged 9 to 13, then expanded to boys aged 11 to 14 in 2017. Following the same trend as in OECD countries, the vaccine is offered under Brazil's Unified Health System (SUS) in PHC facilities and in the context of school-based vaccination campaigns. Cervical cancer screening is performed through a cytopathological examination (Pap smear), offered to women aged 25 to 64 years and repeated every three years. While the three-year periodicity is aligned with World Health Organization (WHO) recommendations, Brazil could learn from OECD countries and move towards a population-based screening programme while introducing HPV-DNA primary testing. This approach is taken internationally (e.g. in Italy and Finland) and recommended by international guidelines, including the European guidelines and American Cancer Society 2020 guidelines. Its implementation would require use of reliable, validated HPV tests in qualified laboratories, accredited by authorised bodies (such as the National Agency of Sanitary Surveillance [ANVISA]) and complying with international standards. Training for health workers is crucial, along with communication strategies for women in target populations. Such a

national strategy should include quality assurance for monitoring cervical cancer screening performance, which the European guidelines define as achieving: (1) an invitation coverage of at least 95% of targeted women; (2) an examination coverage of at least 70% (85% is desirable); and (3) a participation rate of at least 70% (85% is desirable) (von Karsa et al., 2015^[12]).

In the area of breast cancer, the Brazilian Ministry of Health recommends a biennial mammogram for women aged 50 to 69 years old as a screening strategy in PHC. Brazil could go beyond this recommendation and develop a population-based breast cancer screening programme, as implemented in several OECD countries (e.g. Australia, France or Portugal). In practice, this would require implementing an effective call-recall structure, devising a screening registry and safeguarding robust quality assurance at all levels. The window of opportunity is large today with the implementation of the *Previne Brasil* strategy, which will improve patients' registration with FHTs. Brazil could take advantage of the patient registration system to develop organised screening programmes that allow identifying the eligible target population in each municipality. The establishment of an organised breast cancer screening programme will require building capacity in primary care teams, including providing health professionals with guidelines and protocols on cancer signs and symptoms to improve diagnostic accuracy, giving them sufficient time to evaluate patients, and training practitioners in how to diagnose cancer and develop reliable referral mechanisms. At the same time, the distribution of mammographs across the country will need to be revisited to achieve a balanced supply matching population needs.

Population-based programmes can go hand in hand with personalised invitations. In this regard, Brazil could adopt a more systematic and personalised approach to inviting target populations. A growing number of OECD countries send a personal invitation letter for cancer screening, systematically issued through a registry, to each individual in the target group. Brazil has no such national initiative, and invitations are only issued at some local levels, depending on the initiative of individual municipalities or FHTs. By contrast, Denmark, Germany and Italy send invitation letters with a fixed appointment date for a mammography. Based on the available evidence, the most effective interventions to increase participation in organised screening programmes include postal and telephone reminders, the primary care physician's signature on the invitation letter and the provision of a scheduled appointment instead of an open appointment (Camilloni et al., 2013^[13]). A cost-effectiveness analysis would need to be performed to determine the appropriate interventions in the Brazilian context.

Brazil should develop more systematic monitoring of cancer while using digital health to improve public awareness and health literacy

While Brazil has expanded its cancer information infrastructure (notably with SISCAN, PAINEL-Oncologia and the population-based cancer registry), the main challenges are low coverage, low interconnectedness and lack of feedback. Brazil needs a more comprehensive registry-based information system capable of following each patient individually, to ensure efficient management of screening programmes and cancer care delivery. Such a system is key to identifying target populations and sending personalised invitations and reminders for cancer screening. At the same time, the effectiveness of existing cancer screening protocols in Brazil – such as target groups, screening frequency, methods and cross-population groups – needs to be more systematically assessed. This entails developing integrated information systems to improve the quality of screening programmes, notably through performance benchmarking and feedback to health providers. In fact, provider assessment and feedback have been found to increase coverage for cervical, breast and colorectal cancer screening (Sabatino et al., 2012^[14]).

Brazil should also develop further digital health strategies to improve public awareness of cancer prevention, cancer screening and health literacy. The Ministry of Health has implemented some actions since 2002, such as the October Pink Campaign aiming to promote prevention and healthy lifestyles, and overcome the fear or stigma associated with cancer. At the local level, municipalities promote screening through campaigns using local media and social networks. Some good initiatives are also implemented at

municipal levels through the deployment of units to reach very remote and vulnerable communities, and increase access to screening, diagnosis and treatment.

While all these strategies are valuable and should be maintained, Brazil could also develop more effective communication strategies and education programmes to improve health literacy. Local stakeholders, such as health care providers and civil society representatives, should also be consulted to identify specific barriers to screening and address specific needs, particularly among disadvantaged populations (i.e. remote, poor and less educated people). As systematic sharing of information related to cancer is limited – particularly for cancers that are not subject to screening programmes – Brazil could also develop information-sharing strategies to help patients seek care, including cancer diagnostic services. Brazil could learn from Denmark, England and Estonia, which have developed website or e-health platforms that share evidence-based information on cancer, to promote health literacy and help patients seek appropriate and timely health care, including cancer screening, diagnosis and treatment.

1.2.2. Improving screening for hypertension and diabetes

Faced with the rising prevalence of diabetes and hypertension, which are complex and costly diseases, Brazil must strengthen prevention, early detection and treatment. It should prioritise a shift towards disease-management programmes, building capacity for PHC, and improving the data infrastructure underpinning diabetes and hypertension.

Diabetes and hypertension care are improving

In Brazil, as in many OECD countries, diabetes and hypertension are leading causes of morbidity, associated with significant co-morbidities and expenditures. The total mortality of patients with diabetes and hypertension has been falling in Brazil: diabetes mortality decreased by 9.8% between 2007 and 2019, and hypertension prevalence decreased by 5.3% between 2005 and 2015. Low rates of avoidable admissions for diabetes and hypertension in Brazil also suggest relatively good management of these conditions in PHC: in 2019, the rate of hospital admissions for diabetes was 92 patients per 100 000 people (below the OECD average of 130) and 48 patients per 100 000 people for hypertension (less than half the OECD average).

These are all welcome signs of improved prevention and care for diabetes and hypertension. Nevertheless, the available evidence indicates that the diabetes mortality burden will increase by 144% by 2040, becoming the third leading cause of death (Duncan et al., 2020^[15]). Hypertension also became the leading risk factor for deaths in 2017 (Nascimento et al., 2020^[16]). Given these trends, more efforts are needed to strengthen diabetes and hypertension prevention in the context of PHC. In Brazil, basic health checks and risk screening for chronic diseases rely on FHTs and are predominantly done through opportunistic screenings during health visits. The Ministry of Health has issued several guidelines relating to chronic non-communicable diseases. Screening for hypertension is recommended for adults without the knowledge that they are hypertensive. It suggests screening every two years for people with blood pressure below 120/80 millimetres of mercury (mmHg) and annual screening if systolic blood pressure is between 120 and 139 mmHg or diastolic blood pressure between 80 and 90 mmHg. For diabetes, the guideline recommends screening for asymptomatic adults with sustained blood pressure greater than 135/80 mmHg, but does not apply to other screening criteria, such as obesity, family history or age range. Specific booklets (*Cadernos de atenção básica*) for the management of hypertension and a recent protocol for diabetes also provide guidance for PHC practitioners.

Brazil has sought to improve the quality of diabetes and hypertension care, notably through the PMAQ pay-for-performance system, launched in 2011. As part of PMAQ, the evaluation of FHTs included topics such as early detection of hypertension, laboratory exams for diabetes and prescription refills for users on continued care (e.g. for hypertension and diabetes), without the need to make medical appointments. However, the pay-for-performance component of the new *Previne Brasil* programme only includes two

indicators related to hypertension and diabetes; these only cover follow-up of patients who already have the diseases, and do not address prevention in PHC. It will therefore be paramount in the future to monitor how the new programme influences screening activities for hypertension and diabetes in PHC.

Brazil could learn from OECD countries such as Australia, Estonia, Korea and England that have instigated population-based screening programmes targeting high-risk populations. In Australia, for example, PHC physicians can provide a single health assessment for people aged 45-49 who present at least one risk factor (lifestyle habit or family history) for developing a chronic disease, such as type 2 diabetes or heart disease. The WHO Regional Office for Europe also recommends targeting screening in PHC for patients at risk for developing diabetes or hypertension. If Brazil wants to move in this direction, it will be important to ensure that programmes target high-risk population groups, based on the country's national epidemiology, the characteristics of the health system and a cost-effectiveness analysis.

Improved disease-management pathways and increased workforce capacity in primary health care are essential to better screening and disease management

Chronic disease-management pathways are critical to delivering seamless co-ordinated care for chronic patients. Widely used in some OECD countries to improve care integration and co-ordination, they are associated with cost reductions and better quality of care. In Brazil, the development of two clinical guidelines with explicit pathways of care to guide health professionals in PHC are the only attempts to promote care co-ordination. However, these guidelines do not integrate other care providers or stakeholders (e.g. specialists or patient-support groups) and are not produced in a patient-friendly format.

Brazil should promote disease-management pathways with a people-centred perspective to clearly establish the responsibilities of health professionals at different stages of disease, set quality expectations for chronic diseases such as diabetes and cardiovascular disease, and standardise the quality of care across Brazil. As seen in OECD countries, co-ordinated management approaches enable PHC teams and individual patients to take appropriate actions to manage diabetes and hypertension. Brazil should also ensure that pathways integrate all health care providers across different sectors.

PHC professionals should have the tools and capacities to undertake these responsibilities and tasks. There is scope in Brazil to build capacity for screening and management of hypertension and diabetes through task-shifting or expanding the role of other health workers, including nurses, nutritionists and pharmacists. Nurses could have the capacity and training to conduct an initial patient assessment, including blood-sugar testing in all practices, and collaborating with doctors and other team members in follow-up activities. The benefit of nurse-led-care has already been demonstrated (Maier, Aiken and Busse, 2017^[17]; Martínez-González et al., 2015^[18]).

Last but not least, integrated patient pathways require adequate payment systems to incentivise multiple health professionals to work together. Brazil could consider experimenting the use of bundled payments for chronic conditions. As implemented in several OECD countries (i.e. the Netherlands, Australia and Canada), bundled payments consist of one payment per patient with a chronic illness to cover the cost of all health care services extended by the full range of providers during a specific time period. Such a system has the potential to incentivise co-ordination of care between providers and ensure a broader range of care. While Brazil has already taken steps to improve PHC funding through weighted capitation, that payment model is not likely to spur care co-ordination and integration for diabetes and hypertension.

Improving the data infrastructure underpinning diabetic and hypertension care should be a priority

Brazil has developed Hiperdia, a specific registration and monitoring system for hypertensive and diabetic patients, which is managed by the Ministry of Health in conjunction with the State and Municipal Health Secretariats. Through Hiperdia, municipalities that are members of the National Pharmaceutical

Assistance Programme for Hypertension and Diabetes Mellitus send information on the registration and monitoring of patients with these diseases. Remarkably, all this information is processed and made available by DATASUS, the health data infrastructure and information system of the Ministry of Health. However, Hiperdia does not collect information on screening, nor is it capable of capturing the pathway of care for each patient (referrals, waiting times, consultations with specialists, laboratory tests and medications). It is also difficult for FHTs to review and act on this information as they do not receive regular feedback, and patients do not have access to their personal health data. There are also some problems linking data with other databases, such as hospitalisation and death registries.

Improving the data infrastructure by linking different data sources should therefore be a priority in Brazil, to help patients and health professionals identify shortcomings and further fine-tune prevention strategies. Health information systems should be expanded so that they provide more and better information to both PHC teams and patients, thereby improving clinical management decisions in primary care and empowering patients to take a leading role in their own prevention and control actions. Data should be systematically fed back to practitioners, and should be used to identify risks or shortcomings. Brazil could learn from OECD countries in developing a strong health information system within a data-governance framework that protects the privacy of patients' health information, as laid out in OECD (2019^[19]). A stronger information system would also be crucial to streamlining efforts to identify and invite target populations for hypertension and diabetes screening.

1.3. Addressing workforce challenges in the primary health care sector

Brazil is struggling with a shortage and uneven distribution of medical doctors across regions. The ratio of practising doctors in 2019 was lower in Brazil (2.3 per 1 000 population) than the OECD average (3.5 per 1 000 population). The distribution of medical doctors followed a north-south gradient, where the North and Northeast regions show a lower density of medical doctors. At the same time, FCM is undervalued as a specialty in Brazil. In 2018, only 5 486 medical doctors (1.4% of all specialists in Brazil) had specialty training in FCM. While residents of the North and Northeast have the highest health care needs, less than 20% of all FCM specialists work in these regions. Brazil therefore needs to meet two main challenges: 1) securing a greater number and distribution of primary care doctors; and 2) securing a high-quality workforce.

1.3.1. Securing a greater number and distribution of PHC doctors across the country

The MDP has been successful in strengthening the provision of PHC services in underserved communities

Established in 2013, the MDP is a large-scale health system intervention aiming to strengthen the provision of PHC services in underserved communities. The MDP was designed by the Ministry of Health as a supply-side intervention featuring three objectives: 1) transferring funds to municipalities to strengthen the PHC infrastructure; 2) improving access to and the quality of medical school training; and 3) recruiting Brazilian and foreign physicians to municipalities struggling to recruit and retain doctors. Based on the available evidence, the programme has been successful. Good progress has been made with regard to investing in and renovating the PHC infrastructure, expanding the number of new medical undergraduate vacancies in PHC, and recruiting more than 16 000 physicians from both Brazil and abroad to work exclusively within FHTs. By its fifth year of implementation, the MDP was associated with a 12.2% increase in the number of PHC physicians, with more pronounced effects in municipalities featuring lower levels of physician availability at the start of the programme (Hone et al., 2020^[2]). The evidence also mostly suggests that the MDP led to improved access to care and quality of care, as measured by reductions in avoidable hospitalisations for some chronic conditions (Fontes, Conceição and Jacinto, 2018^[20]).

While these results are remarkable, the implementation of the MDP has not been without drawbacks. First, there is evidence that the MDP led to the substitution of Brazilian physicians who were already working in communities. Second, only three-quarters of priority municipalities have at least one MDP physician, meaning that community-targeting methods were insufficient to recruit physicians for all priority municipalities. Third, MDP physicians did not have to undergo recertification or revalidation to practise medicine in Brazil. It is also fair to note that despite these positive results, unfilled PHC positions are still the norm in Brazil, with persisting geographic disparities in access to medical training. The North and Northeast regions offer the lowest levels of per capita medical undergraduate positions in the country.

The new *Programa Médicos pelo Brasil*, initiated in 2020 by the federal government to replace the MDP, will continue to increase the provision of medical services in remote or highly vulnerable municipalities as well as promote the training of doctors specialising in FCM. To avoid repeating mistakes, the experience of the MDP can inform the design and implementation of the *Programa Médicos pelo Brasil*. The federal government should focus on community-targeting methods to maximise the recruitment of physicians for all priority municipalities. It should also ensure that the new programme does not lead to a substitution of Brazilian physicians. The introduction of a compulsory revalidation process for all medical licences and the establishment of a two-year specialty programme in FCM as part of the *Programa Médicos pelo Brasil* are certainly positive steps for strengthening PHC.

Complementary solutions to recruit and retain the workforce would make the Programa Médicos pelo Brasil more effective

While Brazil already has many complementary solutions to recruit and retain the PHC workforce, it could learn from international experience on health workforce policies:

- First, as part of the new *Programa Médicos pelo Brasil*, there is scope to train more rural doctors in order to boost numbers and improve retention over the longer term. Extensive international evidence shows that establishing medical education in rural locations leads to increases in the number of medical graduates who will work in rural places over the long term (McGirr, Barnard and Cheek, 2019^[21]). Learning from the experiences of OECD countries such as Australia, Canada, England and Norway, Brazil could establish medical education programmes in rural localities. At the same time, it could encourage medical students to gain experience in rural areas, for example, by including a compulsory rural rotation in any medical internship.
- Second, Brazil could make financial and non-financial incentives for medical doctors contingent on practising for a time in underserved communities. As part of the *Programa Médicos pelo Brasil*, physicians in underserved areas will receive financial incentives ranging from BRL 3 000 to BRL 6 000, depending on the location. As implemented in Chile, Germany and Canada, these incentives could be contingent on a return-of-service obligation of two to four years. The federal government will also need to ensure that the various types of financial incentives are received directly by the medical doctors, and not distributed at the discretion of the municipalities. In Chile, the programme combining financial and non-financial incentives with a return-of-service obligation of three to six years has been successful, with the number of applicants exceeding the number of available positions, high satisfaction rates among applicants and high retention rates among the hired doctors (Pena et al., 2010^[22]).
- The final option to redress the geographical imbalance of doctors would be to restrict the choice of practice location for PHC doctors. In this respect, Brazil could learn from Turkey, Germany, Norway or some Canadian provinces (e.g. New Brunswick and Quebec), which restrict the choice of location for newly qualified medical doctors. Such a regulatory approach requires careful and coherent workforce planning, which is far from being the case today. In Brazil, unlike in many OECD countries, there is a lack of knowledge of present and future needs for doctors and other health care professionals (Oliveira et al., 2017^[23]). Health care workforce planning is not based on

an objective assessment of professionals' characteristics, work processes, the characteristics of the health system in place and the population's health needs.

Boosting the local workforce, changing its scope of practice and making greater use of technology are key levers to cope with health workforce shortages

In many low, middle and high-income countries, task-shifting is a key lever to fill the gap in PHC provision stemming from a lack of PHC doctors (Afobali et al., 2019^[24]). While extensive international evidence supports the transfer of roles traditionally performed by doctors to nurses and other allied health professionals, Brazil has not yet experimented with changing the scope of practice and task-shifting among health workers such as nurses and community health workers. As implemented internationally (e.g. in Australia and France), Brazil should capitalise on its large existing nurse training network to expand and recognise the role of advanced practice nurses, who could help manage the care of patients with mild acute diseases and chronic disorders, complementing the practising PHC doctors. Alternatively, advanced practice nurses can be deployed in remote and underserved areas to enhance access to qualified health professionals. Such a policy would require a thorough analysis of current nursing curriculums to further invest in core competencies, including team practice, care co-ordination between and across health sectors, clinical and professional leadership, and quality and safety management (Cassiani et al., 2018^[25]). In a similar vein, community health workers – who already play a crucial role in PHC provision in Brazil – could play a bigger role, notably by taking a patient's medical history, conducting basic physical exams, ordering tests and providing counselling on preventive care. This would certainly require investments in training, as well as improved communication between community health workers and other FHT staff.

Improved use of alternatives to face-to-face consultations could be another avenue for coping with PHC doctor shortages and geographical imbalances in Brazil. While telehealth has been used for quite some time, its development and use are patchy. For example, just under 3.5 million telehealth services across all levels of care were registered in Brazil between 2016 and 2019. Of these, 50% occurred in the Southeast, 36% in the South and only 1% in the North, the most deprived region in terms of workforce shortages. A lack of funding, training and equipment for FHTs are factors impeding the use of telehealth services to deliver safe, fast and seamless health care services in Brazil. Scaling up telemedicine in Brazil without compounding existing social divides will require new investment and funding to ensure the effective and equal use of digital technologies in PHC. Brazil will also need to establish a governance structure that better supports municipalities with insufficient resources.

1.3.2. Securing a high-quality workforce through medical education

Beyond the need to address the uneven geographic distribution of primary care physicians, Brazil should focus on three broad areas: 1) increasing the credibility and recognition of the primary care sector; 2) making FCM a mandatory requirement to be allowed to practise primary care; and 3) introducing quality measures to medical education.

Increasing the credibility and recognition of the primary care specialty is a huge challenge in Brazil

While the Family Health Strategy has made access to PHC a priority, the country has not invested sufficiently in enhancing the credibility and recognition of FCM. Several worrying indicators suggest that the FCM specialty is undervalued and is the last option chosen by physicians: in 2018, only 1.5% of recent graduates chose the FCM speciality among the 55 specialities available. The low degree of academic incorporation and low funding in this research area are factors impeding recognition of the specialty (Wenceslau, Sarti and Trindade, 2020^[26]; Fontenelle et al., 2020^[27]). And yet they are critical to promoting FCM and encouraging students to choose this specialty as their future career, notably by proving that PHC is not a profession that does not require many qualifications. If Brazil wishes to fill the gap in the

PHC workforce, it needs to create academic FCM departments in all medical schools, promote interdisciplinary collaboration with other specialists and encourage mutual respect.

At the same time, making specialty training in FCM compulsory for all medical doctors wishing to practise PHC would be paramount to promote excellence in Brazil, and deliver more effective and patient-centred primary care. The benefits of specialised PHC have already been demonstrated elsewhere (OECD, 2020^[28]), notably to address the bulk of patient needs, and provide a stronger focus on prevention and early management of health conditions. Post-training requirements to practise PHC are currently implemented in 22 OECD countries; there is no doubt that such a professionalisation would bring benefits to Brazil.

Last but not least, standardising the contractual arrangements for primary care workers is key to improve the attractiveness of PHC as a profession. Since each municipality defines the level of salaries, bonuses and other payment mechanisms, these arrangements are widely heterogeneous across municipalities. Stronger oversight and regulation by the federal government is warranted to ensure consistent salary scales across the country, and it could be important to ensure that smaller municipalities offer similar contracts as larger municipalities. The new Agency for the Development of Primary Health Care could play a role in this regard. Key examples for learning are available internationally. In Turkey, for example, the implementation of the Health Transformation Programme led to family physicians being paid according to national terms and conditions, with more generous salaries and improved working conditions.

Quality measures for medical education should be introduced without delay

While Brazil's Ministry of Health (in partnership with Ministry of Education) regulates medical education nationwide, more could be done to standardise the medical curriculum. The inadequate academic programme, and the gap in the quality of medical education among medical schools, are areas of concern. To achieve greater standardisation of the medical curriculum and teaching, Brazil could introduce a national test at the end of the university period. The national licensing examination would complement the accreditation performed by the National Institute of Educational Studies and Research Anísio Teixeira (INEP) or the Federal Council of Medicine. The national examination has been increasingly used across OECD countries. The United Kingdom is introducing a nationwide Medical Licensing Assessment, which is overseen and regulated by the General Medical Council and will test the core knowledge, skills and behaviours needed to practise medicine. Brazil could learn from the United Kingdom's example.

At the same time, Brazil could take further step to promote workforce quality through stronger requirements for CME. Unlike in many OECD countries, CME is voluntary in Brazil. Not only is this insufficient to guarantee a high standard of competencies for PHC doctors, but it also does not ensure their fitness to practise throughout their career. More needs to be done to help medical doctors maintain their competency and acquire new skills. While a number of bodies, such as CONASS, CONASEM, the Brazilian Medical Association or the Brazilian Society of Family and Community Medicine organise trainings to maintain and update competencies, it is impossible to know whether – and how many – PHC medical doctors have undertaken CME. A stronger regulatory approach is required, for example through a formal CME framework that clarifies the expectations placed on PHC doctors and helps them meet these expectations. Brazil could learn from the United Kingdom and the Netherlands, where CME for PHC doctors is compulsory and linked to recertification: PHC physicians need to demonstrate they have regularly participated in CME activities, and undergo peer evaluation of their professional skills. At the federal level, the Agency for the Development of Primary Health Care could set such requirements and monitor providers' CME compliance. Municipalities could also use the contracting arrangement to actively push PHC doctors to comply with CME requirements and encourage quality improvement. CME requirements should be directed towards the key challenges of the health care system, such as the prevention of risk factors for health (e.g. obesity) and management of chronic conditions (including diabetes).

1.4. Promoting the digital transformation of primary health care in Brazil

1.4.1. Brazil is on the right track for an effecting digital transformation of PHC

Brazil has established a solid foundation for digital PHC by developing strategies at various levels of government, and making key investments in networks, data, interoperability and skills. As a result of the country's use of digital tools and data to enhance the efficiency, transparency and responsiveness of its public institutions to citizen and business demands, Brazil scored above the OECD average on the 2019 Digital Government Index (OECD, 2020^[29]).

The Brazilian National Digital Health Strategy 2020-28 (ESD28) builds on more than a decade of policies to digitalise health care and make better use of health data (Ministério da Saúde, 2020^[30]), going as far back as the National Policy for Health Information and Informatics of 2004 (PNIIS). Actions within ESD28 focus on three axes: 1) build on existing programmes and initiatives to digitalise health care in Brazil, including the National Health Data Network (RNDS), the Connect SUS (*Conecte SUS*) and Primary Health Unit Computerisation (Informatiza APS) programmes; 2) develop the necessary conditions (e.g. organisational, legal, regulatory) to spur effective collaboration in digital health; and 3) establish a conceptual, normative, educational and technological forum (*espaço de colaboração*) to operationalise collaboration.

A component of the *Previne Brasil* programme, Informatiza APS provides federal funding to FHTs that collect and send data from electronic health record (EHR) systems (Harzheim et al., 2020^[31]). The programme also supports training in ICT and data analysis for employees in municipal health departments. In 2020, Connect SUS ran a pilot project in the state of Alagoas that provided additional funding for digitalisation to PHC units that had been unable to digitalise, disbursing more than BRL 5.5 million (USD 1.07 million [United States dollars]) to 97 municipalities (Ministério da Saúde, 2020^[32]).

Connect SUS is also seeking to expand the use of health information systems in PHC through its e-SUS APS strategy (also known as e-SUS AB), which seeks to better manage individual health and care, optimise data collection and improve health information (Ministério da Saúde, 2020^[33]). The informatics department of SUS (DATASUS) has made various applications available to citizens, frontline care workers and managers. Meanwhile, municipalities are increasingly providing digital services to citizens, with 25% of prefectures allowing patients to book appointments online and consult with a doctor online. Start-ups in telemedicine are flourishing, including Brasil Telemedicina, which produces around 60 000 reports per month and has clients in more than 700 Brazilian cities, and Telelaudo, which has provided more than 4.7 million teleradiology reports to more than 450 hospitals.

As elsewhere, the COVID-19 pandemic has provided further impetus to the digital transformation of health in Brazil. On 15 April 2020, the federal government authorised the use of telemedicine during the COVID-19 pandemic (Law No. 13.989/2020) to ensure continuity of care and protect health care workers. To help PHC units access digital technologies, the Ministry of Health also instituted an exceptional incentive to be paid to municipalities and states for each health team that has not yet been digitalised, budgeting over USD 83.8 million (BRL 432 million) for 2020. The Brazilian Telemedicine University Network (RUTE) also created a special interest group on COVID-19.

1.4.2. An effective digital PHC hinges on addressing three sets of challenges

There is no doubt that Brazil has a solid basis on which to build an effective, equitable and efficient digital PHC. Achieving a successful digital transformation of PHC now hinges on tackling three sets of interconnected challenges: 1) digitalising all PHC units and teams; 2) promoting inclusive connectivity for all Brazilian citizens, especially the most vulnerable; and 3) establishing a governance structure with clear, well-funded mandates at the right levels of government.

Focus on fundamental enablers to increase the digitalisation of PHC

While pockets of excellence exist, progress towards effective use of digital technologies in PHC has been slow and fundamentally unequal: 3 600 PHC units across the nation currently have no computers, and 7 200 have no Internet access. Only 17% of PHC units have Internet speeds over 10 megabytes per second (Mbps), limiting simultaneous use of EHR systems and high-quality video consultations, not to mention remote patient monitoring. Additionally, around 18 000 health care facilities (18% of all facilities) do not have an EHR system, with large regional differences in adoption: in 2019, one in four PHC units still maintained clinical records and patient data in paper format only (CGI.br, 2020^[34]). Only 40% of primary care managers consider their ICT equipment to be new and up-to-date, and only 32% believe their Internet connection suits their needs. Around one-third of nurses and physicians reported having completed capacity-building courses in health informatics. COVID-19 has likely accelerated digitalisation, but its true impact remains unclear.

A real digital transformation of PHC in Brazil will not happen without setting two foundations. First, the country must ensure a reliable energy supply and fast connectivity for all PHC units, tackling one of the main barriers to digitalisation reported by municipalities during the recent pilot project in Alagoas. Although previous programmes (e.g. Connected Brazil) have been instrumental in expanding broadband connectivity, it is crucial that these initiatives set ambitious and clear targets for both the reliability and speed of connections in PHC units. It is also important to improve co-operation among governmental entities and across the different levels of government (federal, state and municipal) in implementing broadband connectivity initiatives, building on good practices in OECD countries and beyond.

Second, financial payments included in the Connect SUS pilot project in Alagoas should be consolidated and expanded to the entire Brazilian territory. To ensure the funds are used for their intended purpose, receipt of financial support should be tied to participation by municipal and state officials in capacity-building workshops, especially targeting senior executives. Funding should be linked to explicit targets for the adoption and use of EHR systems (like e-SUS AB), including effective electronic drug prescribing and exchange of clinical and administrative data across units. As PHC units go digital, it will be vital to safeguard operational dependence on digital systems by enforcing appropriate cybersecurity.

Ensuring digital uptake and effective use among the most vulnerable and digitally excluded

Even if Brazil were to break down all barriers to the connectivity and digitalisation of PHC units, significant inequalities among citizens in the use of digital technologies and tools would persist. Nearly 47 million Brazilians did not use the Internet in 2019 (more than 20% of the population). Digital divides are clearly associated with social inequalities, with older Brazilians, those from rural areas, those with lower incomes and those with lower education levels less likely to use the Internet, and more likely to use mobile phones only. As a result, there exists a serious risk that a digital transformation of PHC will compound existing social divides.

An effective digital PHC that reaches all Brazilians is only possible if every citizen is not only connected, but also considered in the design of digital PHC services. Brazil should focus on eliminating digital divides as a foundation for effective digital PHC, strengthening the role of telecentres as providers of training – especially in rural and remote areas – and ensuring appropriate funding and technical assistance at all levels of government. Given the importance of mobile phones as a means of accessing the Internet, it should prioritise a multichannel approach focusing on mobile access among vulnerable groups. Although digital PHC content, services and applications should meet the needs of both health workers and patients, they should specifically target citizens with low digital uptake, with continued efforts to expand patients' access to their personal health information.

Moving from digital PHC strategies to implementation and monitoring

Brazil's decentralised government creates challenges to digitalising PHC. The country's single federal district, 26 states, well over 5 000 municipalities (around 44% of which have under 10 000 inhabitants) and 438 health regions have highly heterogeneous demographics, socio-economic indicators and governance structures. With only around 800 inhabitants, the municipality of Serra da Saudade has the same responsibilities in managing the digitalisation of PHC as the municipality of São Paulo, which numbers over 12 million inhabitants. Decentralisation has many benefits, but it is also creating barriers to the digital transformation of PHC in Brazil to meet the targets set in ESD28.

A major barrier to digital PHC in Brazil is human and technical capacity in municipalities. Setting responsibilities for the digitalisation of PHC at the municipal level across Brazil is likely leading to diseconomies of scale and creating challenges for small municipalities. Brazil could look to examples from New Zealand, England and Portugal for a new model of shared services and responsibilities related to the digitalisation of PHC at less decentralised levels. Building on existing institutions (e.g. Co-ordenação-Geral de Tecnologia da Informação e Comunicações/General Co-ordination of Information and Communications Technology) and experiences, Brazil could explore setting up dynamic purchasing systems at the federal level, with appropriately funded agencies at the state or regional levels providing municipalities with specialised human resources, purchasing and management services for digital PHC.

Brazil has invested significantly in connectivity (e.g. through *Governo Eletrônico – Serviço de Atendimento ao Cidadão*) and software (e.g. e-SUS AB), as well as provided municipalities with support for hardware and training (e.g. Informatiza APS). Yet the ambitious vision set out in ESD28 will likely require both larger and smarter investments, tying financial support to effective and equitable use. Taking into account the magnitude of investments made in more digitally advanced countries, Brazil could review its current investments and consider providing more funding for change management, capacity-building, and monitoring and evaluation. Financial support could be tied to specific milestones in the adoption and use of digital PHC functionalities, starting with key functionalities like ePrescribing, referral management, appointment scheduling and patient access to EHR.

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2 Health care needs and organisation of primary health care in Brazil

The expansion of PHC in Brazil has been remarkable, contributing to significant improvements in many measures of population health over the past decades. SUS prioritisation on PHC permitted to offer services free of charge to the Brazilian population, allowing an easier contact with the health system at the community level through the development of modern multi-disciplinary family health teams. But gains have not been equal across the country, with North and Northeast regions faring worse than South and Southeast regions in terms of health improvements and health outcomes. There has also been a rise in the number of elderly patients and an increasing prevalence of risk factors for health and chronic non-communicable diseases. Tackling these challenges call for more effective PHC, responsive to people's changing needs, capable of offering preventive, continuous and co-ordinated care.

2.1. Introduction

Brazil is a high-middle income country, with a population of 212 million inhabitants in 2020. The country is divided into 26 states and one federal district, with 5 570 municipalities. The states are further organised into five geopolitical regions: North, Northeast, Centre-West, Southeast and South. Health is a universal right for the whole population and a state responsibility since the approval of the new Brazilian constitution in 1988, which established the Unified Health System (Sistema Único de Saúde, SUS) in 1990. The management and financing of SUS is shared between the Federal Ministry of Health, state departments and municipal health departments. In the decentralised health care system, municipalities have the freedom to organise PHC family health teams according to the population needs and geographic characteristics.

Brazil has made investments in PHC a priority to address disparities in access to care and improve health outcomes. The Family Health Strategy, one of the largest community-based PHC programme worldwide, launched in 1994 was successful to increase population coverage and improve health outcomes. The expansion of the Family Health Strategy has contributed to measurable improvements in terms of infant mortality rates, maternal health, immunisation uptake and avoidable hospitalisation. However, PHC in Brazil faces a number of challenges resulting from its vast territory and complex geography, which have made difficult to meet the needs of population living in vulnerable and remote communities. The expansion of PHC has been marked by stark disparities across the states and municipalities, linked mainly to workforce shortages and imbalances in the distribution of medical doctors. There has also been a rise in the number of elderly patients and an increasing prevalence of chronic non-communicable diseases, which call for effective prevention and stronger PHC to better manage the burden of chronic conditions.

This chapter describes the PHC system in Brazil. It starts by analysing the socio-economic context in which the system operates, as well as describing the governance of the PHC sector. The chapter then examines the health care needs of the Brazilian population, and lastly presents the organisation of the PHC sector.

2.2. The socio-economic context for PHC in Brazil

Brazil's government is divided into 26 states and the federal district, and 5 570 municipalities. The states and the federal district are also grouped into five regions: North, Northeast, Central-West, Southeast and South, which are used mainly for statistical purposes and to define the distribution of federal funds. Municipalities are granted the status of federal entities, at the same level as the states. While there is great heterogeneity in the geographical area and population, as well as social and economic indicators, among the subnational jurisdictions, all Brazilian municipalities enjoy the same legal status (OECD/UCLG, 2019^[1]). In 2015, 17 municipalities had more than 1 million inhabitants (22% of population). About 44% of municipalities had less than 10 000 inhabitants. Brazil also has 438 health regions, which are made up of municipalities, and are very heterogeneous in their demographics, socio-economic indicators and governance structures (Ministério da Saúde, 2020^[2]).

In the recent past, strong economic growth and income distribution programmes were the driving forces for generating significant inequality and poverty reduction in Brazil. However, income inequalities have been rising again in recent years, and there are large regional disparities in prosperity and growth across the country, with North and Northeast regions faring worse than South and Southeast regions. Large inequalities along several dimensions affect health status.

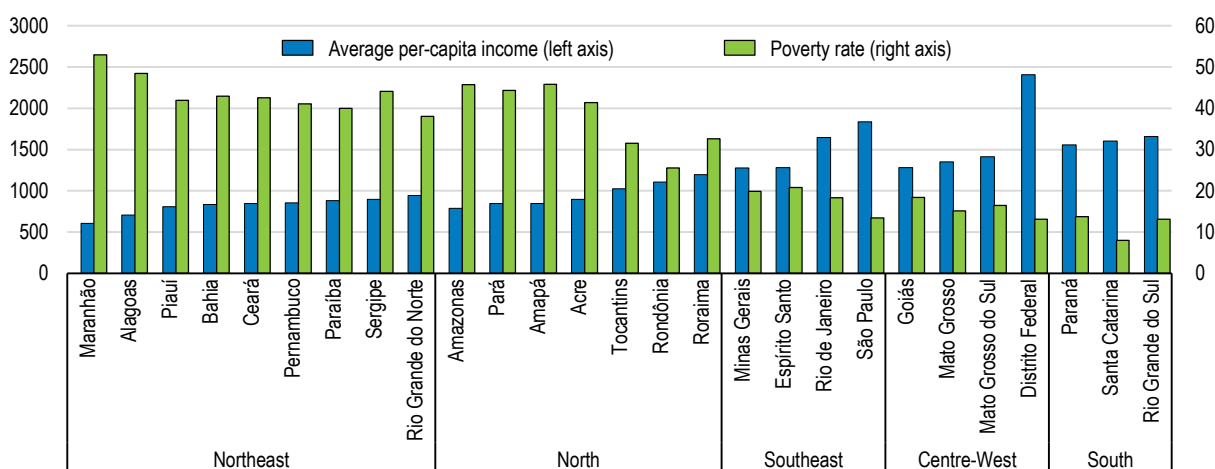
2.2.1. Despite progress, socio-economic inequalities remain a significant problem in Brazil

Brazil has had strong economic growth combined with remarkable social progress over the past two decades. In 2010, Brazil GDP grew by 7.5%, compared to around 3% across OECD countries. Currently social benefits amount to over 15% of GDP, and are characterised by poor targeting if pensions are also included. As a result, poverty rates have been reduced from 41% of the population in 2001 to 19% in 2019.¹ Public programmes have contributed to the decline in poverty rates (OECD, 2020^[3]). Recent evidence shows that the *Bolsa Família* programme alone has reduced poverty rates by 15% (Ferreira de Souza, Osorio and Paiva e Sergei Soares, 2019^[4]).

These progresses have also generated reduction in income inequality. From 2001 to 2011, the GINI of income inequality in Brazil dropped continuously from around 58 to 52. However, it has been rising again in recent years, at around 53, and remains a significant problem in Brazil. Currently, the richest 10% of the population earn more than four times as much as the bottom 40% (OECD, 2020^[3]).

Regional inequalities are another crucial challenge in Brazil. The distribution of income across Brazilian regions is highly unequal. Average incomes in the Northeast state of Maranhão are less than half of the national average. The opposite holds true for poverty. The poverty rates in the North and Northeast regions are between three to four times higher than the poverty rates in the Southeast, Central-west and South regions (Figure 2.1). Large regional disparities also exist with regards to labour market informality and illiteracy, which are three to four times more common in the poorer North and Northeast regions than in the relatively affluent South and Southeast regions (OECD, 2020^[3]). As the social and economic context strongly influences health outcomes, economic disparities across regions are mirrored in health status and health outcomes.

Figure 2.1. Socioeconomic inequalities across regions, 2019



Source: IBGE (2019^[5]), Síntese de Indicadores Sociais: uma análise das condições de vida da população.

2.2.2. The management of the Brazilian Unified Health System is decentralised

The principles and structure of Brazil's health care system were conceived in 1988 after the approval of the new Brazilian Constitution that established health as a universal right for the whole population. It became a state responsibility with the establishment of the Unified Health System (Sistema Único de Saúde, SUS). The SUS is structured around three main priorities (Massuda et al., 2020^[6]):

- The universal right to comprehensive health care at all levels of complexity (primary, secondary, and tertiary).
- Decentralisation with responsibilities given to the three levels of government: federal, state, and municipal
- Social participation in formulating and monitoring the implementation of health policies through federal, state, and municipal health councils.

The management of SUS is shared across three different levels of governance: Federal Ministry of Health, state departments and municipal health departments. At federal level, the Ministry of Health defines the general objectives of the SUS and sets policy development, planning financing, auditing and control of health policies and health services linked to SUS. It is responsible for the overall co-ordination of the national priorities and actions defines as part of the Multi-Annual Plan 2020-23 and the National Health Plan (Plano Nacional de Saúde, [PNS]) 2020-23. The health related objectives for the Multi-Annual Plan 2020-23 is the expansion of the coverage and resolution of PHC, improving prevention and strengthening of integration between health care services. The PNS 2020-23 elaborates on SUS health policy objectives, with the definition of seven programmes (Ministério da Saúde, 2020^[7]):

- Pharmaceutical services in SUS
- Specialised health care services
- PHC
- Scientific, technological and productive development in health
- SUS management and organisation
- Protection, promotion and recovery of indigenous health
- Health surveillance

In the area of PHC, the PNS 2020-23 is structured around three main priorities. First, the expansion of PHC access and of its infrastructure. Second, improvement in the digitalisation of PHC units, with better-integrated data and use of electronic medical records in North and Northeast regions. Third, increasing health promotion and prevention at PHC levels to address risk factors for health (for example nutrition and smoking) and encourage uptake of screening exams (for example cytopathology and mammography exams). The PNS 2020-23 defines 20 objectives in the area of PHC, with specific targets to achieve by 2023 around for example the number of Family Health Teams (both for the general and incarcerated populations), dental care coverage, pregnant women's prenatal consultations, cytopathology and mammography exams coverage, and quality of mental health (Table 2.1).

Table 2.1. The PNS 2020-23 objectives in the area of PHC

	Goals	Indicators	Reference Level	2023 Target
1	Increase to 72% the population coverage of PHC	Population coverage with PHC	63,5%	72,7%
2	Increase to 185 the number of "Doctor's Office at the Street" Teams	Number of "Doctor's Office at the Street" Teams	154	185
3	Increase to 2000 the number of Family Health Units with extended working hours financed by <i>Saúde na Hora</i> Programme	Number of Family Health Units functioning on extended working hours financed through the <i>Saúde na Hora</i> Programme	264	2 000
4	Increase to 155 the number of reference PHC team targeted at teenagers serving socio-educational measures	Number of PHC reference teams targeted at teenagers serving socio-educational measures	119	155
5	Increase to 677 the number of prison-based health teams	Number of prison-based health teams	357	677
6	Increase the occupation rate of more than 85% for placements offered through federal medical programmes (<i>Mais Médicos</i> project + <i>Médicos pelo Brasil</i> project)	Occupation rate of offered placements through federal medical programmes	82%	>85%
7	Increase to 80% the share of pregnant women with at least	Percentage of pregnant women with at least 6	40%	80%

	Goals	Indicators	Reference Level	2023 Target
	6 prenatal consultations taking place during the first 20 th week of pregnancy	prenatal consultations, taking place during the first 20 th week of pregnancy		
8	Increase to 49% the population coverage for Dental Specialties Centres	Percentage of population covered by Dental Specialties Centres	45%	49%
9	Increase to 689 Dental Specialties Centres that have adhered to the Network of care for People with Deficiencies	Number of Dental Specialties Centres that have adhered to the Network of Care for People with Deficiencies	579	689
10	Increase to 46% the population coverage of Dental Health Teams	Percentage of population covered by Dental Health Teams	42%	46%
11	Increase to 3 118 the number of municipalities with Dental Prosthesis Regional Laboratories services	Number of municipalities with Dental Prosthesis Regional Laboratories services	2468	3118
12	Increase to 60% the share of pregnant women with prenatal consultations at PHC who had individual dental appointments with Dental Health Teams	Proportion of pregnant women with dental appointments performed	25%	60%
13	Implement 39 River-based Family Health Units	Number of River-based Family Health Units implemented	19	39
14	Implement out-of-hospital psychosocial service focused at acute situations in 14 capitals	Number of capitals with out-of-hospital psychosocial service focused at acute situations	0	14
15	Reduce to 26 days the average time of psychiatric hospitalisations in medical institutions	Average time of psychiatric hospitalisation	45	26
16	Increase to 1.2% the proportion of alcohol dependents' appointments at the Centres of Psychosocial Care	Percentage of alcohol dependents' appointments at the Centres of Psychosocial Care	0,6%	1,2%
17	Increase to 80% the share of beneficiaries of the <i>Bolsa Família</i> Programme who have their health conditions monitored at PHC	Percentage of the <i>Bolsa Família</i> beneficiaries with health conditions monitored at PHC level	76.08%	80%
18	Increase to 48% the share of women aged between 25 and 64 years old that have done a cytopathology exam in the last 3 years	Percentage of women aged between 25 and 64 that did a cytopathology exam in the last 3 years	36%	48%
19	Increase to 35% the share of women aged between 50 and 69 years old that have done a mammography exam in the last 2 years	Percentage of women aged between 50 to 69 years old that have done a mammography exam in the last 2 years	23%	35%
20	Reduce the proportion of caesarean sections to 52.4%	Percentage of caesarean sections	55.7%	52.4%

Source: (Ministério da Saúde, 2020^[7]), Plano Nacional De Saúde 2020-2023.

The 26 state governments, through states secretary of health, are responsible for regional governance and the regional health system organisation, co-ordination of strategic programmes, and delivery of specialised health care services that have not been decentralised to municipalities. At local level, 5 570 municipalities, through municipal secretary of health, handle the management of SUS, including co-financing, co-ordination of health programmes, and direct delivery and contracting of health services, including PHC services (Massuda et al., 2020^[6]).

The Interfederative Pact of Executive Management allows SUS to have a dynamic functioning through agreements between each federative entity, with the establishment of the following councils or commissions:

- The *Conselho Nacional de Secretários de Saúde* (CONASS) brings together the health secretaries of the states and the Federal District with the purpose of implementing SUS principles, legislation and guidelines into health actions and services. CONASS also allows the exchange of experiences and information among States Secretaries.
- The *Conselho Nacional das Secretarias Municipais de Saúde* (CONASEMS) represents municipal health departments, with the objective of guiding health managers for the provision of PHC services.

- The *Conselho Estadual de Secretarias Municipais de Saúde* (COSEMS) brings together municipal secretaries of health to discuss and co-ordinates strategic health issues.

Box 2.1. The Governance of PHC in Brazil

The *Política Nacional de Atenção Básica* (PNAB) describes the governance of PHC in Brazil, the roles of the main actors and other strategic aspects of the PHC policy. The federal government, the state governments and municipalities are the main actors involved in the governance of PHC in Brazil. The specific competencies of each level of government is foreseen in the Federal Constitution and the PNAB (art. 7, Ord. 2 436).

The formulation of policy priorities and actions for the PHC sector is agreed and approved during the CIT, which is made of representatives from the Ministry of Health, CONASS and CONASEMS. Decisions on the operational, financial and administrative aspects for the management of PHC are taken by consensus.

Within the Ministry of health, the PHC Secretariat (SAPS) (established in 2019 – Decreto nº 9 795) is responsible for defining and co-ordinating the PNAB. It is also responsible for allocating federal resources to finance PHC (on a monthly, regular and automatic basis), providing integrated support to the state and municipal managers, formulating strategies for monitoring and improving PHC quality, establishing national guidelines and providing technical and educational supports for PHC managers and professionals.

The states define decentralised PHC actions, strategies, guidelines, as well as rules for the implementation of the PNAB during the CIB, which is made of representatives from the state secretaries of health and COSEMs. States are expected to allocate financial resources for PHC, monitor PHC actions and care quality in municipalities, verify the quality and consistency of information sent by municipalities. They might also provide institutional support to the municipalities in the process of implementing and monitoring PHC and expanding the Family Health Strategy. States are expected to provide technical and educational instruments to the municipalities to facilitate training and continuous education of PHC managers and professionals.

Municipalities have the main attribution concerning the provision and financing of PHC services. They are responsible for organising, managing and providing services and actions within each territory. Municipalities have to provide support to PHC teams and units in the process of monitoring and expanding the Family Health Strategy. To this end, municipalities select, hire and remunerate the PHC professionals in accordance with the current legislation, and guarantee that PHC teams and units have sufficient material resources, equipment and supplies to deliver PHC services. They might organise effective patient flow within SUS and adopt referral mechanisms to ensure effective care co-ordination. Municipalities should keep the registration of teams, professionals, workload, available services and equipment on a monthly basis in the National Registry System of Health Establishments according to specific regulations. Community participation should also be guaranteed through social control. Lastly, municipalities are required to analyse and verify the quality and consistency of the data collected in the national health information systems to be sent to other spheres of management, use them in the health care planning and disseminate results.

Source: Information provided by Brazilian Ministry of Health.

The *Comissão Intergestores Tripartite* (CIT) and the *Comissão Intergestores Bipartite* (CIB) are important bodies for the development of SUS strategies and resource allocation through collective consensus from the different spheres of government (see Box 2.1). The National Health Council, a collegiate body

composed of government representatives, service providers, health professionals and users, is also involved in the formulation of PHC priorities and actions. The OECD health system review provides a detailed description of these governance bodies (OECD, 2021^[8]).

Since the establishment of SUS, Brazil has made substantial progress toward achieving universal health coverage. Virtually all of the population is covered for equal benefits and equal financial protection within the public health sector. Since its inception, a strong focus of SUS was to strengthen the role of PHC to reduce health inequalities and improve access to care, but also to move away from a health system that was historically very hospital-centred.

2.2.3. Brazil makes spending on PHC a priority

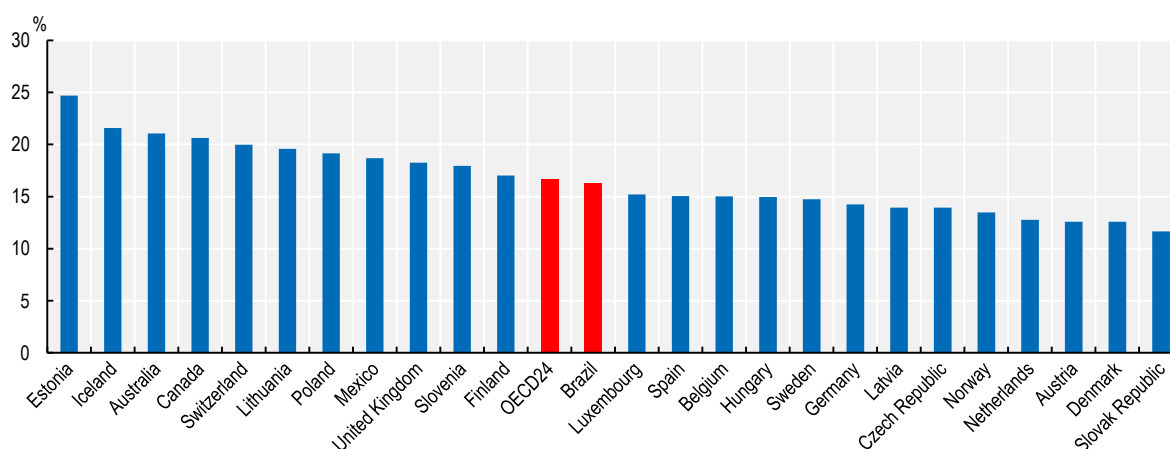
The public system is financed by tax revenues and social contributions from all three levels of government. By law, the federal government is required to spend 15% of net current government income on health, the states are required to spend 12% of total revenue on health, and municipalities are required to spend 15% of total revenue on health (Massuda et al., 2020^[9]). It is reported that over the past 30 years, the share of municipal funding has increased to reach nearly one-fourth of their own total revenues on health (Massuda et al., 2020^[9]).

Under SUS, the federal government has traditionally financed roughly one-third of total PHC costs (33%), with the remaining part being funded predominantly by municipalities (61%) (Ministério da Saúde/Fundação Oswaldo Cruz, 2018^[10]). The participation of states is the smallest, with very few cases where states co-finance policies for PHC.

Analysing the composition of overall health spending in an international context, Brazil seems to make spending on PHC a higher priority than many OECD countries. In 2019, Brazil dedicated around 16% of its financial resources to general outpatient care, dental care, home-based curative care and preventive activities (Figure 2.2).

In 2020, the budget allocated to PHC from the federal government was BRL 20.9 billion, compared to BRL 17.5 billion in 2019 (CONASEMS, 2020^[11]). In total, the federal government increased its PHC budget by around BRL 3.4 billion between 2019 and 2020.

Figure 2.2. Spending in PHC as a share of current health spending, 2019 or nearest year



Note: Basic health services combine general outpatient care, dental care, home-based curative care and preventive activities provided regardless of the setting. It can be used as a proxy for PHC for international comparisons.

Source: OECD (2021^[12]), *Health Statistics Database*, <http://stats.oecd.org>; Brazilian Ministry of Health.

2.3. Health care needs of the Brazilian population

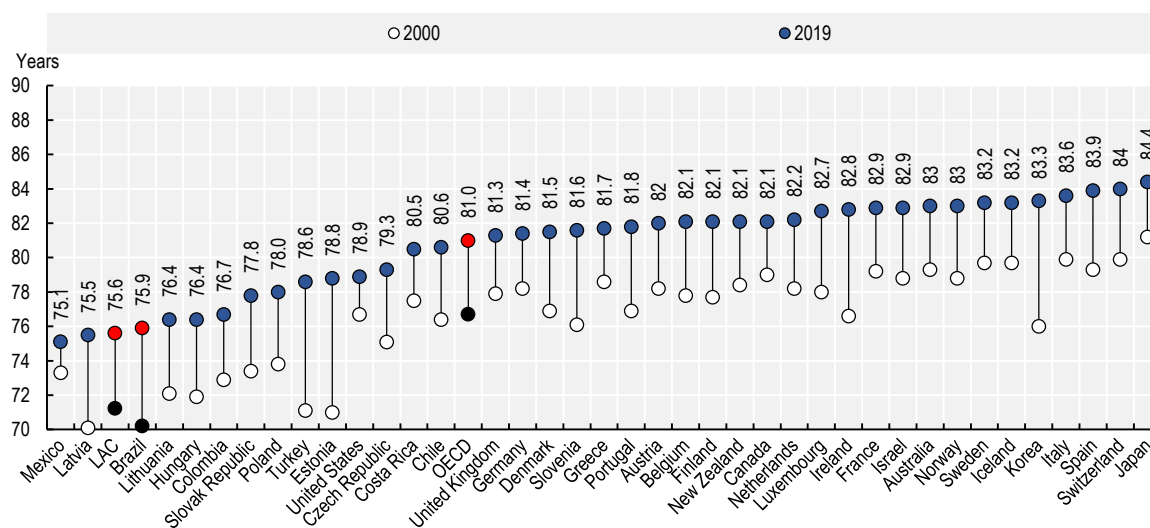
Since SUS inception, Brazil has made significant developments by improving in most of the general population health indicators. Brazil is however undergoing a profound demographic transition. As a result of decrease in fertility and mortality rates, the Brazilian population is rapidly ageing. The rapidly ageing population and increasing risk factors for health in Brazil (notably growing overweight rates) goes hand in hand with an increased prevalence of chronic non-communicable diseases, which call for effective prevention and stronger PHC to better manage the burden of chronic conditions.

2.3.1. Life expectancy at birth has improved in recent years, but is still falling behind other OECD countries

Life expectancy at birth in Brazil increased rapidly over the past decades, but is still five years below the OECD average of 81 years but above the LAC average of 75.6 years (see Figure 2.3). In Brazil, life expectancy at birth increased by more than five years between 2000 and 2019 (rising from 70.2 to 75.9 years), compared to four years across OECD countries. The gap in longevity between Brazil and other OECD countries has shortened from 6.5 years to 5.1 years.

Figure 2.3. Life expectancy at birth in Brazil has increased over the past decades

Life expectancy at birth, among OECD and LAC countries, 2000 and 2019 (or nearest years)



Source: OECD (2021^[12]), *Health Statistics Database*, <http://stats.oecd.org>; World Bank World Development Indicators Online 2021.

Infant mortality rates have decreased from 30.3 deaths per 1 000 live births in 2000 to 12.4 deaths per 1 000 live births in 2019. Despite this decrease, the infant mortality rate in Brazil is still above the OECD average of 4.2 deaths per 1 000 live births. The same is true for maternal mortality rates in Brazil, which have decreased to 60 women per 100 000 live births in 2017 (13 percentage points reduction since 2000), although still higher than the OECD average of 8 per 100 000 live births but lower than the LAC average of 83 (OECD/The World Bank, 2020^[13]).

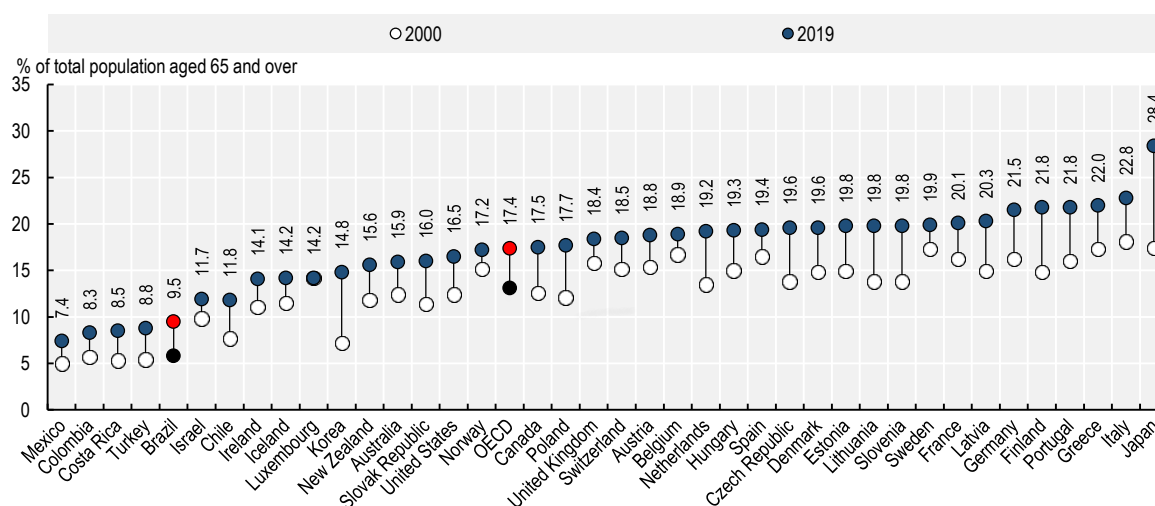
2.3.2. Brazil faces rapid ageing of its population

Brazil, similarly to other OECD countries, is experiencing a demographic transition characterised by a shift from high levels of mortality and fertility to lower levels. The fertility rate fell from 2.9 births per woman in

1990 to 1.7 births per woman in 2019 (OECD, 2021^[14]). The combination of falling mortality rates (resulting in increasing life expectancy) and fertility rates causes rapid population ageing. The share of the population aged over 65 years has increased by 65% over the past decade, rising from 5.8% in 2000 to 9.5% in 2019 (Figure 2.4). Across the OECD, the share has increased by 32% over the same period, rising from 13.1 in 2000 to 17.4 in 2019.

Therefore, the dependency ratio (the number of people of working age for every person aged 65 years and more) is projected to substantially decrease by 2050, close to the OECD level. The rapid ageing population has an important impact on the health of the population in Brazil, putting pressure on both the health care systems and the economy.

Figure 2.4. Share of the population aged over 65 years, 2000 and 2019



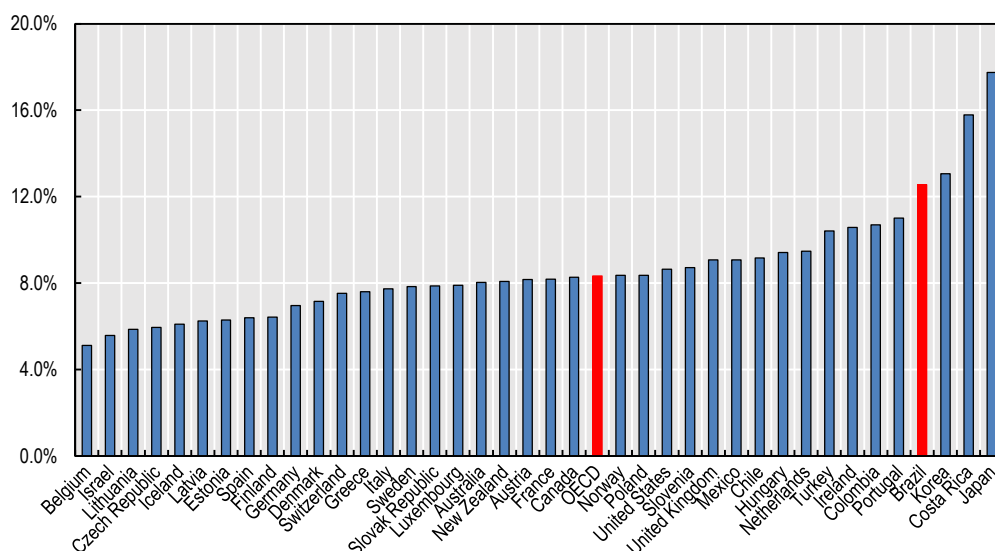
Source: OECD (2021^[12]), *Health Statistics Database*, <http://stats.oecd.org>.

2.3.3. Rapidly increasing risk factors for health and associated ill-health are a major concern

Several risk factors for health give cause for concern in Brazil, including particularly overweight. Being overweight is a major risk factor for various chronic non-communicable diseases including diabetes, cardiovascular diseases and certain cancers. While overweight in Brazil is currently below OECD average, the trend is increasingly upwards. In 2016, estimations show that 56.5% of adults in Brazil were overweight, close to the OECD average of 58.4%. However, Brazil showed the fourth largest change in overweight between 2006 and 2016 with a growth of 13% in the past decade, compared to an increase of 8% across the OECD. This is the third highest growth rates behind Japan, Costa Rica and Korea (Figure 2.5).

Children overweight rates in Brazil are also increasing at a fast rate. In 2016, Brazil had a children overweight rate of 28%, very close to the 28.5% in average amongst OECD countries. Nevertheless, Brazil experienced an increase of 27% between 2006 and 2016, higher than the 15% increase in the OECD. Increasing children overweight rates means higher risk of cardiovascular diseases or diabetes during adult age, putting further pressure on the PHC sector.

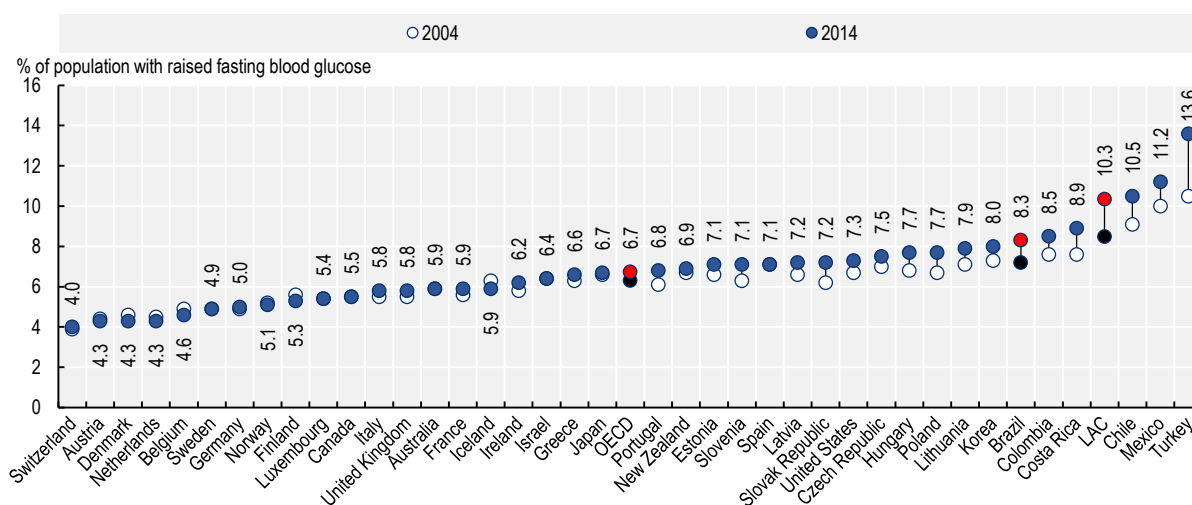
Figure 2.5. Changes in overweight rates among adults between 2016 and 2006



Source: OECD (2021^[12]), *Health Statistics Database*, <http://stats.oecd.org>.

Already, when compared to OECD countries, diabetes represents a higher burden of disease in Brazil. Raised levels of blood sugar can lead to the development of diabetes. Fasting blood glucose (FBG) contributes to diagnose and monitor diabetes. In Brazil, 8.3% of the population had raised FBG in 2014, above the 6.7% average in OECD countries. Importantly, between 2004 and 2014, Brazil recorded one of the largest increase in its population with raised FBG, increasing by 15.3% (Figure 2.6). Only Turkey, Mexico, Chile, Costa Rica and Colombia had larger increases amongst OECD countries, evidencing a worrying trend in Brazil.

Figure 2.6. Raised fasting blood glucose among adults in Brazil and OECD, 2004 and 2014



Source: WHO Global Health Observatory (2017^[15]), <https://www.who.int/data/gho>.

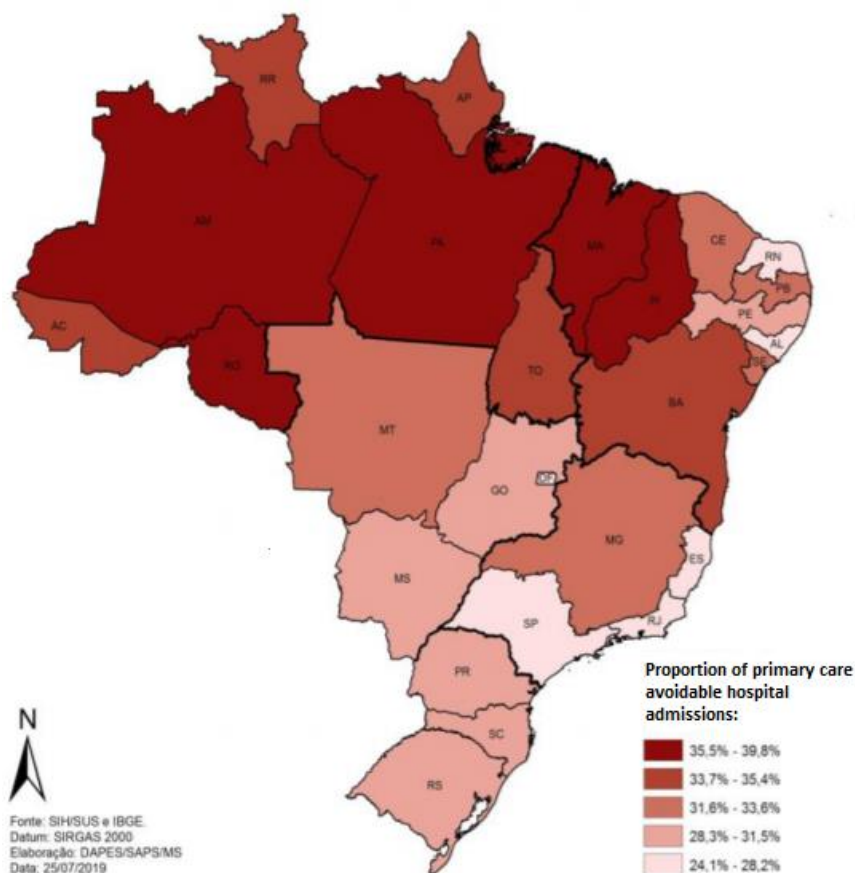
Overall, with increasing risk factors for health, the predominance of chronic non-communicable diseases is growing in Brazil. In 2017, four chronic non-communicable diseases dominated the causes of mortality in Brazil: circulatory system diseases (27%), neoplasms (17%), chronic respiratory diseases (12%), external causes (12%) and diabetes (5%), compared to only two in 1990 (cardiovascular diseases with 27.8%, and neoplasms with 11.4%).

2.3.4. There are large regional differences in health

There are large regional inequalities in health status across Brazil, where the most vulnerable municipalities consistently present poorer health outcomes than national average. While infant mortality rate in Brazil has decreased from 47.1 to 12.4 per 1 000 live births between 1990 and 2019, there are large differences in the magnitude of this reduction across Brazil's regions. The Northeast region showed the most significant decline of 80% in the period, while the lowest degree of reduction occurred in the Centre-west and North regions (reduction of 62%) (Szwarcwald et al., 2020^[16]).

Another illustration of regional differences in health can be seen in terms of avoidable hospitalisation for conditions that could be treated in the PHC setting. The proportion of hospitalisations due to conditions that could be more effectively treated in the PHC setting ranges from 24% in the South region to 40% in the North region (Figure 2.7). Such regional differences point out that some remote and vulnerable areas require more attention than others to improve quality of PHC.

Figure 2.7. Share of hospital admissions due to conditions that are treatable in PHC, population aged 60 or more by region, 2019



Source: Ministério da Saúde (2020^[2]), Plano Nacional de Saúde 2020-23.

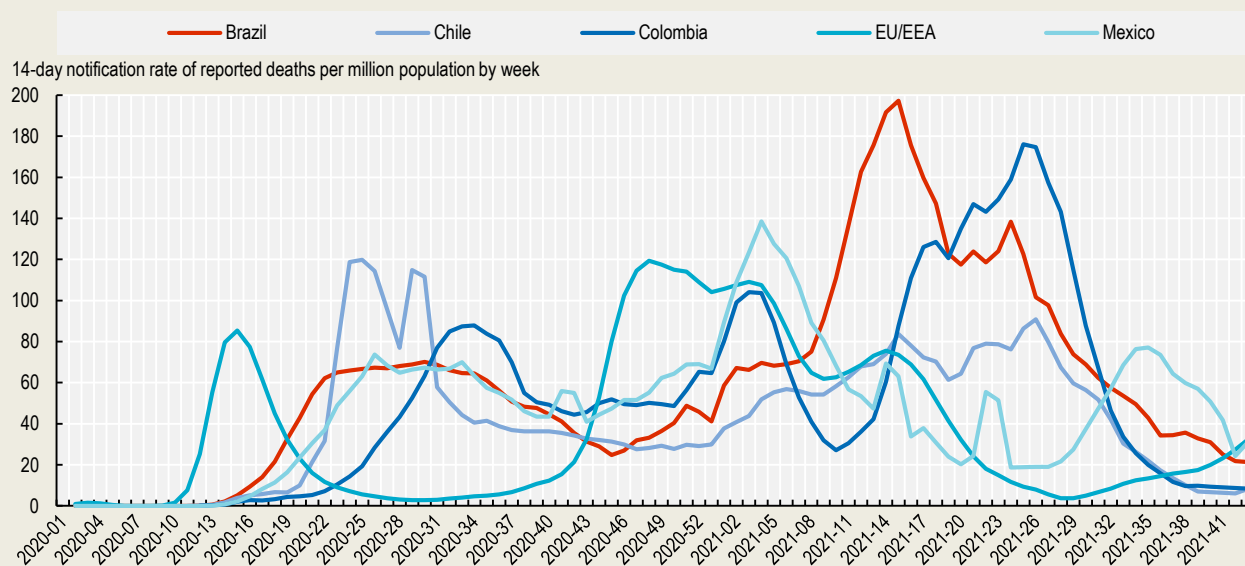
There are also significant inequalities in the rates of vaccination against yellow fever in Brazilian states where vaccination has been recommended, ranging, in 2018, from 21.8% in Sergipe to 100% in the Federal District and Roraima (the national average for all states where vaccination is recommended is 64.1%).

Regional inequalities in health have aggravated the impact of the COVID-19 pandemic and will likely exacerbate health inequalities in the future. The most precarious and vulnerable population are more likely to be affected by the COVID-19 virus, not only because of poor living and working conditions, but also because they are more likely to suffer from comorbidities and risk factors for COVID-19 severity. Mortality rates in the State of Amazonas have been found four-times higher than the national average, particularly among indigenous people (Pires, Carvalho and Rawet, 2020^[17]). Box 2.2 presents the impact of COVID-19 on health in Brazil.

Box 2.2. The impact of COVID-19 on health in Brazil

The first case of COVID-19 in Brazil was reported on the 25th of February 2020. Since then, the impact on population health has been large. The total number of registered deaths is above 611 000 as of 17 November 2021 with 41 000 average monthly registered deaths in 2021. This situates COVID-19 as the first cause of death during the pandemic when compared to the average monthly figures for 2015-19 of deaths attributed to other conditions, the highest ones being 30 000 for circulatory diseases, 18 500 for cancers, and 13 000 for diseases of the respiratory system. However, there are scientific basis showing an underreporting of around 22% due to limitations to perform the SARS-CoV-2 screen by RT-PCR test (Carvalho, Boschiero and Marson, 2021^[18]).

Figure 2.8. COVID-19 mortality rate evolution in Brazil and selected OECD countries between March 2020 and October 2021



Source: ECDC (2021^[19]), COVID-19 situation updates.

The mortality epidemic curve in Brazil experienced an increase in cases by early April 2020, reaching the first peak during July 2020, later than what Europe experienced in the early months of the pandemic (Figure 2.8). Then, cases started to reduce but still with high level of virus circulation and mortality rates, similar to other Latin American countries such as Chile, Colombia and Mexico but dissimilar to Europe where mortality was substantially lowered between July and September. By mid-November, cases in Brazil began to rise once again. By the beginning of April 2021, mortality rate in Brazil reached its highest levels since the start of the pandemic and then started to decline.

One of the most important problems in handling the COVID-19 pandemic in Brazil was weaknesses in governance of the health sector. At the federal level, in the first year of the pandemic, four different ministers of health were in office, limiting continuity in the management of the response. Risk communication from federal authorities has also been conflicting and confusing for the population. Brazil is not unique, in that other OECD countries have pursued similar approaches.

As highlighted in Chapter 3, the COVID-19 crisis demonstrates the importance of placing PHC at the core of health systems, both to reduce pressure on health systems and minimise complications and direct deaths from the COVID-19, as well as to maintain ongoing routine care to patients who do not have COVID-19.

2.4. The organisation and structure of the PHC sector

In Brazil, the 5 570 municipalities handle the management, organisation and delivery of PHC. The Family Health Strategy programme launched in 1994 was remarkable. Based on multi-professionals PHC teams (called Family Health Teams), the expansion of the Family Health Strategy led to improvement in health care coverage and health outcomes. SUS prioritisation on PHC permitted an easier contact with health services at the community level.

2.4.1. Municipalities have primary responsibility for managing and delivering PHC

Since the implementation in 1994 of the federal programme called the Family Health Strategy (Estratégia de Saúde da Família – ESF), also known as the Family Health Programme (Programa Saúde da Família), the 5 570 municipalities have primary responsibility for managing and delivering PHC. With federal funding, municipalities have the freedom to organise family health teams according to the population needs and geographic characteristics.

The ESF is a decentralised programme aimed at expanding the PHC system in Brazil toward more co-ordinated, comprehensive and continuous care. The overarching objective was to ensure universal access to health care services for all Brazilians under the overall organisation of SUS (Macinko and Harris, 2015^[20]; Castro et al., 2019^[21]). The ESF became the main mechanism for expanding coverage to PHC services in Brazil. The ESF resulted in two major shifts. First a shift in priorities, from treating patients with diseases to delivering comprehensive care and preventing diseases through active health promotion. Second a shift in governance, from states to municipalities to allow for greater accountability and more collaboration between the private and public sector.

The ESF is funded through federal transfers but also includes financial contributions from municipalities. As earlier mentioned, the municipalities is the main financing agent of PHC. Smallest municipalities (up to 20 000 inhabitants), where there is no other supply of health care services, spend around 60% of their health expenditure directed towards PHC.

2.4.2. Public PHC is free of charge and is expected to be the initial contact point to SUS

SUS offers preventive services and PHC free of charge. Services that are publicly covered are based on health conditions, clinical and nonclinical procedures, and target population groups. The list of services has been gradually adapted to reflect changes in population health. Treatment and control of major chronic conditions (such as hypertension and diabetes) have been added in the list of services covered (OECD, 2015^[22]). The federal Ministry of Health negotiates with state and municipal health secretariats to decide new health care services to be covered by SUS.

PHC provided by the ESF is expected to be the initial contact point for patients having new health needs, although it does not work as a “gatekeeper” and many Brazilian have direct access to outpatient specialties and hospital. For example, the latest Health Information Survey found that around 50% of the diabetic population identified basic health units as their last contact with the health system, 11% used public speciality units in hospitals, 6% public urgent care units and 30% private practices or clinics (IGBE, 2020^[23]).

This is unlike many other OECD countries such as Australia, Canada, or the United Kingdom where general practitioners operate as gatekeeper to the health system as a referral is required and compulsory to access to specialist and hospital care (OECD, 2020^[24]).

In PHC, private sector is more supplemental as health plans mainly offer specialist and hospital services. However, health plans in the private sector have begun to offer access to family doctors with no co-payment as an alternative to specialist visits (Massuda et al., 2020^[9]).

2.4.3. PHC in Brazil is provided by multidisciplinary Family Health Teams

Prior to 1994, the role of PHC and health promotion was limited and services were predominantly delivered in a range of hospitals. As a community-based, large-scale PHC programme, the ESF relies on multi-professional teams called Family Health Teams (FHTs). Each FHT is assigned a geographic area, covering up to 4 000 individuals located in non-overlapping areas. This helps FHTs to consider social, economic and environmental factors of the catchment area. FHT are responsible for registering every family in their geographic area, and are thus designed to facilitate access to the health care system (Massuda et al., 2020^[9]).

A range of different health professionals are involved in PHC delivery. The FHT include physicians, nurses, and up to 12 community health workers. Other PHC workers, including nutritionists, pharmacists, social workers, psychologists, obstetricians and gynaecologists, and public health workers can provide additional support depending on local needs. Oral health teams have been integrated to the FHT in recent years to provide universal and free oral health services to the population. By 2019, there were 43 234 Family Health Teams and 27 564 Oral Health Teams in operation throughout Brazil (Figure 2.9). Solo practice in Brazil is thus becoming scarce, although there is no official registration and reporting of this practice.

This model of service delivery – based on multidisciplinary team – is well aligned with modern approaches to PHC which house multiple professionals with advanced skills working in teams (OECD, 2020^[24]). Such models of PHC based on teams or networks of providers were reported by 18 OECD countries in 2018, including Australia, Canada, France, Switzerland or the United States.

FHTs are working in basic health care units (*Unidades Básicas de Saúde* – UBS) or other PHC facilities called health clinics, health centres or health posts, and can also receive support from the Family Health Support Centers (*Núcleos de Apoio à Saúde da Família* – NASFs). NASFs were introduced in 2008 to provide support to between one and nine FHTs. NASF is composed of health and social workers included for example acupuncturists, social workers, physical education professionals, pharmaceuticals, physiotherapists, gynaecologists and obstetricians; homeopathic doctors; nutritionists; paediatricians or psychologists. By 2019, there were 4 487 NASFs in operation throughout Brazil (see also Chapter 4 for a detailed description of NASFs). As of May 2021, there is a total of 50 029 PHC facilities in Brazil, including among other UBS, Health Post and NASFs.

In 2020, the government introduced a new classification of 12 health teams categories (Table 2.2 summarises the various family health teams in Brazil).

Table 2.2. PHC teams in the National Register of Health Facilities CNES, 2020

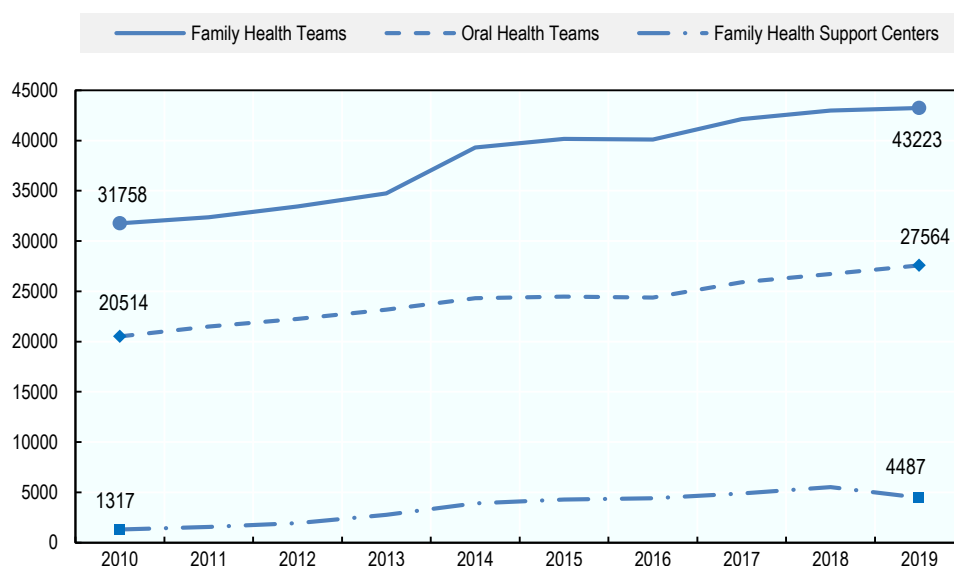
Acronym	Official name	PHC team
EMSI	<i>Equipe Multidisciplinar de Saúde Indígena</i>	Multidisciplinary Indigenous Health Team
EMAD I	<i>Equipe Multidisciplinar de Atenção Domiciliar Tipo I</i>	Multidisciplinary Team of Home Care type 1
EMAD II	<i>Equipe Multidisciplinar de Apoio</i>	Multidisciplinary Team of Home Care type 2
EMAP	<i>Equipe Multidisciplinar de Atenção Domiciliar Tipo II</i>	Multidisciplinary Support Team
EAD	<i>Equipe de Cuidados Domiciliares</i>	Home Care Team
eSF	<i>Equipe de Saúde da Família</i>	Family Health Team
eSB	<i>Equipe de Saúde Bucal</i>	Oral Health Team
eNASF-AP	<i>Equipe do Núcleo Ampliado de Saúde da Família e Atenção Primária</i>	Extended Family Health and PHC Team
ECR	<i>Equipe dos Consultórios na Rua</i>	Outdoors clinics Team
eAPP	<i>Equipe de Atenção Primária Prisional</i>	Prison PHC Team
eMAESM	<i>Equipe Multiprofissional Atenção Especializada em Saúde Mental</i>	Multi-professional Team Specialised Attention in Mental Health
eAP	<i>Equipe de Atenção Primária</i>	PHC Team

The scope of practice of the FHT has gradually increased over the past years to provide proactive preventive care and deliver public health intervention including for instance immunisation programmes and screening programmes (see Chapter 3). Services include not only maternal and child health care services, but also entail preventive care and control and management of chronic non-communicable diseases (Özçelik et al., 2021^[25]). The FHT also link patients with social services (such as conditional cash transfer programmes), water and sanitation services, law enforcement, and school programmes (Massuda et al., 2020^[9]).

2.4.4. The expansion of the Family Health Strategy has been remarkable, improving health care coverage and health outcomes

In 2019, 98% of the municipalities had adopted the Family Health Strategy model. In 2019, there were 43 223 family health teams, 27 564 oral health teams and 4 487 family health support centres (Figure 2.9), providing care to more than 127 million people (61% of the population). The number of family health teams and oral health teams has increased by respectively 36% and 34% over the past decade. At the same time, the number of NASFs has more than tripled over the past decade (Figure 2.9). As of May 2021, there are more than 50 000 PHC facilities in Brazil where teams can deliver services.

Figure 2.9. The number of Family Health Teams has increased by nearly 40% over the last decade



Source: Information provided by the Brazilian Ministry of Health.

There are strong evidence suggesting that the FHT provides better access and quality than other traditional health posts and centres (Massuda et al., 2020^[9]). In a systematic review of the literature examining the impact of the ESF in Brazil, Bastos et al. (2017) show that increasing health care coverage through the expansion of the ESF was consistently associated with improvement in child mortality (Bastos et al., 2017^[26]). A longitudinal analysis using panel data show that a 10% increase in FHT coverage was associated with a 4.5% decrease in infant mortality rate, all other things being equal (Macinko, Guanais and Marinho De Souza, 2006^[27]).

Among adults, expansion of the ESF was associated with reductions in hospitalisations for diabetes mellitus and respiratory problems (Guanais and MacInko, 2009^[28]). Estimations show that the ESF may have contributed to an estimated 126 000 fewer hospitalisations between 1999 and 2002. More recently,

Pinto and Giovanella (2018) show that the implementation of the ESF was associated with a reduction of 45% of the standardised hospitalisation rates per 10 000 inhabitants between 2001 and 2016. The decline was particularly pronounced for hospitalisation due to asthma, gastroenteritis and cardiovascular and cerebrovascular diseases (Pinto and Giovanella, 2018^[29]). The implementation of the ESF is also positively associated with a significant reduction in the number of hospitalisations for heart failure and stroke per 10 000 inhabitants from 1998 to 2013 (De Fátima Barros Cavalcante et al., 2018^[30]). Better prevention of PHC sensitive conditions, greater follow-up for chronic conditions, improved diagnosis, and easier access to medicines partly explain improvement in health care quality.

In addition to improving health outcomes through better access to care, the SUS made considerable investments to expand the availability and equipment of PHC facilities through the *Requalifica programme*. The programme carries out construction, expansion and refurbishment of the PHC facilities, including the construction of the Basic Fluvial Health Units, providing better conditions for accessing and offering public health actions and services. It also involves actions aimed at the computerisation of PHC, and the qualification of PHC workers (Ministério da Saúde, 2020^[2]).

While these are all welcome news, the ESF expansion has also been marked by stark disparities across the states and municipalities (Andrade et al., 2018^[31]). The ESF expansion has been heterogeneous across municipalities mainly because of geographic imbalances in the supply of physicians. The lack of PHC physicians in some remote municipalities has been one of the most important barrier to the expansion of the ESF (Andrade et al., 2018^[31]).

2.4.5. The role of community health workers is predominant

Following the implementation of the Community Health Agents Programme (PACS), community health workers (CHWs) started to play a crucial role in the Brazilian PHC system. CHWs are frontline public health workers, often members of the communities in which they work. They have valuable knowledge of people needs and are trustful source of information. The implementation of the ESF gave an expanded role of CHWs, at the core of the country's PHC policy (Wadge et al., 2016^[32]).

CWHs are integrated members of the FHT. They provide comprehensive PHC supports including clinical triage, supporting chronic disease management, screening uptake, immunisations, providing pre- and postnatal care assistance, health promotion, social determinants, community participation and household data (Wadge et al., 2016^[32]). They are expected to resolve many low-level problems such as medication review for chronically ill patients, while referring more complex issues to doctors and nurses of the FHT. At the clinic or health posts, CHWs are also responsible for organising the waiting rooms and appointments for other health workers, and offering educations sessions.

Each CHW is assigned to maximum 150 households in a geographic area, and each household receives at least one visit every month from a dedicated CHW irrespective of needs or demand. Patients are not able to choose their CHW. Monthly visits allow health promotion activities and basic clinical care, checking whether prescriptions have been filled and whether patients have been taking their medications regularly, identifying risk factors for health or other social determinants, but also collecting individual and household level data (Wadge et al., 2016^[32]). CHWs also help co-ordinate PHC services with public health efforts.

In 2020, more than 250 000 CWHs served 61% of the population (see Chapter 5), up from 60 000 in 1998. The minimum requirement to become a CHW is to get secondary education, but around two-thirds of them have a professional diploma. CHWs are most often recruited from within their own community. It is a stable and respectful profession and as a result, turnover is low.

A large body of evidence demonstrates the positive impact of CHWs on patient satisfaction and health literacy, notably for low-income and vulnerable groups (Wadge et al., 2016^[32]; Grossman-Kahn et al., 2018^[33]). CHWs have been key to effectively respond to the Zika virus for instance, providing health advice and incidence reporting. CHWs could thus have played a greater role to support the COVID-19 response.

So far, evidence shows that only an estimated 9% of CHWs have received infection control training and personal protective equipment during the pandemic. They have not been provided with clear guidance about their role in the COVID-19 response. The COVID-19 pandemic has revealed governance issues linked to the health sector, mainly due to a lack of leadership and oversight from the federal government. Risk communication from federal authorities has also been conflicting and confusing for the population and PHC workers, with no overall co-ordination (Lotta et al., 2020^[34]).

2.4.6. Municipalities are responsible for hiring health professionals for PHC, with wide variations across the country

In the decentralised health care system, municipalities are responsible for the organisation, management and delivery of PHC services. In 2020, the federal government provided BRL 20.9 billion to the municipalities, which were then responsible for organising and delivering PHC services, by hiring FHT members as civil servants. As each municipality defines the level of salaries for health team members, there are wide wage variations across the country (Massuda et al., 2020^[6]).

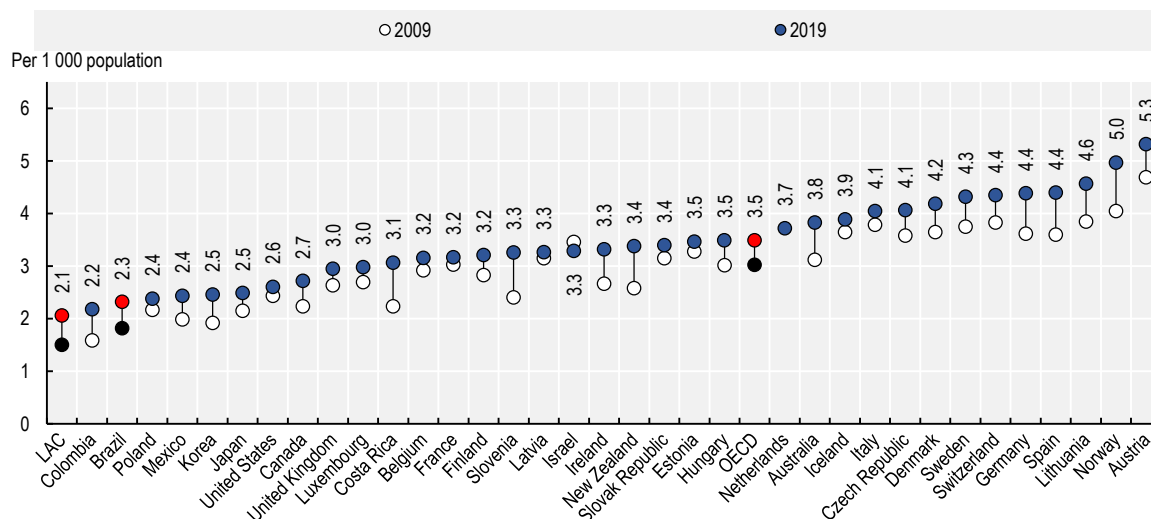
There are three types of possible contractual schemes with the municipality:

- Public officials: there are civil servants hired as statutory workers (there are salaried, with a stable position and there is a ceiling of money to pay them);
- Private workers hired by social organisations, non-profit private NGO (but anecdotal evidence suggests there are forms of corruption under such schemes),
- Private workers hired by Government owned social organisation NGO.

2.4.7. There are too few PHC physicians in Brazil

The number of trained PHC physicians is well below the OECD countries, and of other LAC countries. In Brazil, practicing doctors per 1 000 population in 2019 were very low compared to the OECD average, at 2.3 compared to 3.5 (Figure 2.10). This is among the lowest density, just above Colombia (with 2.2 doctors per 1 000 population) and the LAC average (2.1 doctor per 1 000 population). However, the number of doctors increased by 50% in Brazil over the past decade.

Figure 2.10. Doctors per 1 000 population, 2009 and 2019 (or latest year available)



Source: OECD (2021^[12]), *Health Statistics Database*, <http://stats.oecd.org>; Brazilian Ministry of Health.

At the same time, while the specialty of family and community medicine was introduced as a clinical and licensed specialty, the number of PHC doctors has not increased accordingly. In Brazil, PHC is mostly delivered by physicians who do not have any specialty or by physicians with a specialty in internal medicine (see Chapter 5).

Table 2.3. Distribution of selected medical specialties per regions, 2018

	Physicians with no specialty training	Physicians with specialty training	Internal medicine (<i>Clinica Medica</i>)	Family and Community Medicine (<i>Medicina de Família e comunidade</i>)
North	10118	10766	3.4%	4.6%
North East	34461	46162	18.0%	14.8%
South East	91124	153180	52.3%	43.1%
South	20948	47482	17.3%	31.6%
Central West	12828	24708	8.9%	5.8%
Total number	169479	282298	42728	5486
%	37.5% of all doctors	62.5% of all doctors	11.2% of all specialists	1.4% of all specialists

Source: Scheffer et al. (2018^[35]), Demographia Médica No Brasil 2018.

In 2020, there are around 60 000 medical doctors working in PHC, alongside 50 000 nurses and 35 000 dentists currently in practice in Brazil. In 2018, only 5 486 medical doctors had a specialty training in family and community medicine, representing only 1.4% of all specialists in Brazil (see Table 2.3). Medical doctors specialised in internal medicine who have some competencies in family medicine to provide PHC, represent 11.2% of all specialists in Brazil. Lastly, there are 169 479 physicians, with no specialty training, representing around 37% of all medical doctors.

2.4.8. The uneven distribution of medical doctors is a major concern in Brazil

There is a marked disparity in the concentration of doctors in different regions and states, and between state capitals and other parts of states. Overall, there are 2.3 doctors per thousand population (Figure 2.10) but there are capitals with more than 12 doctors per thousand population – like Vitória, in Espírito Santo – and in the interior part of the Northeast region or in the Amazon with values below one doctor per thousand inhabitants. The Southeast is the region with the highest medical density per inhabitant (with 2.81 per thousand population), against 1.16 per thousand population in the North and 1.41 in the Northeast (Scheffer et al., 2018^[35]).

Among states, the Federal District has the highest density with 4.35 doctors per thousand population, followed by Rio de Janeiro (with 3.55 doctors per thousand population), São Paulo (with 2.81), Rio Grande do South (with 2.56), Espírito Santo (with 2.40), and Minas Gerais (with 2.30). At the other end of the scale, are states of the North and Northeast. Maranhão presents the lowest density of medical doctors with 0.87 doctor per thousand population, followed by Pará with 0.97 doctor per thousand population (Scheffer et al., 2018^[35]).

Geographical inequalities in the distribution of medical doctors hold true when looking at physicians with specialty training. Evidence indicates a clustering of PHC specialists in some regions: the best supplied regions are the South and Southeast Regions, followed by the Central-West (Scheffer et al., 2018^[35]). Workforce imbalance across regions is analysed in further details in Chapter 5 of this report.

Several nationwide initiatives have attempted to address the imbalances in the distribution of health workers. Brazil introduced several programmes to improve access to medical training in underserved communities, including the 2001 Programme of the Interiorisation for Health Work (Programa de Interiorização do Trabalho em Saúde), the 2011 Programme for Valuing PHC Professionals (Programa de

Valorização dos Profissionais da Atenção) and the 2013 More Doctors Programmes (Programa Mais Médicos).

As described by Chapters 4 and 5 of this report, the More Doctor Programme (MDP) is considered as the world's largest government-led initiative that recruited domestic and foreign physicians to work in PHC settings within municipalities that typically faced difficulties in recruitment and retention. By November 2020, the programme was responsible for the presence of 16 426 doctors in 3 837 Brazilian municipalities. This programme has proven to have positive results, for instance, in terms of investment in PHC facilities, doctors availability and health outcomes (Hone et al., 2020^[36]; Netto et al., 2018^[37]).

On a less positive note, unfilled medical positions are still common in Brazil, even after the implementation of the MDP. In April 2019, it is estimated that 3 847 public sector medical positions in almost 3 000 municipalities remained unfilled (Scheffer et al., 2018^[35]). Unfilled medical positions in the PHC sector certainly impede prevention, early diagnoses, the monitoring of children, pregnancies and the continuation of treatments for those having underlying health conditions.

2.5. Conclusion

In Brazil, the 5 570 municipalities handle the management, organisation and delivery of PHC. SUS offers preventive services and PHC free of charge for its population, based on multidisciplinary family health teams. Overall, the organisation of PHC delivery in Brazil is well aligned with modern approaches to PHC, which house multiple professionals with advanced skills working in teams. SUS prioritisation on PHC also permitted an easier contact with health services at the community level. The implementation of the Family Health Strategy, one of the largest community-based PHC programme worldwide, has contributed to increase population coverage, and led to measurable improvements in terms of infant mortality rates, maternal health, and avoidable hospitalisations. The life expectancy at birth in Brazil increased rapidly over the past decades – faster than across the OECD countries.

Despite major investments to improve access to PHC, health continue to be unequally distributed with people in North and Northeast regions suffering from poorer health outcomes. In part, these disparities are linked to workforce shortages and imbalances in the distribution of medical doctors, as well as lower PHC infrastructure and equipment in North and Northeast regions. There are also been a rise in the number of elderly patients and an increasing prevalence of chronic non-communicable diseases, which call for more effective prevention and stronger PHC to better manage the burden of chronic conditions. Brazil will need to continue to invest in a strong PHC system that is responsive to people's changing needs, capable of offering continuous, proactive and preventive care. The next four chapters set out in detail where changes are needed and how it can be achieved, notably to increase screening and prevention for the main chronic non-communicable diseases (Chapter 3), enhance the quality of PHC provision (Chapter 4), tackle workforce challenges in PHC (Chapter 5) and promote the digitalisation of PHC (Chapter 6).

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Notes

¹ The poverty rates is measured by the poverty headcount ratio at USD 5.50 a day (2011 PPP).

3

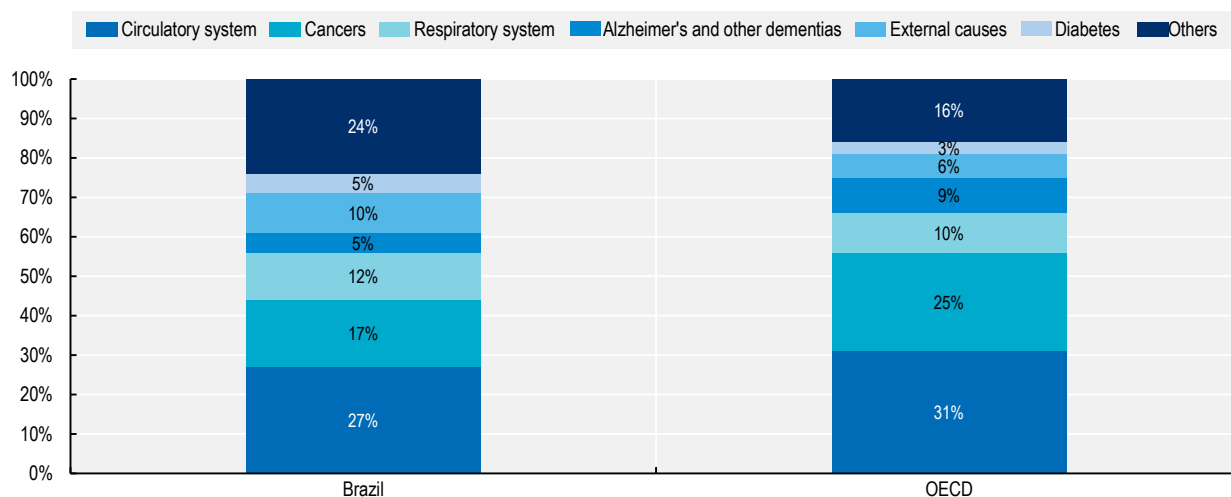
Screening in primary health care for the main chronic non-communicable diseases in Brazil

In Brazil, chronic non-communicable diseases such as cancer, diabetes and hypertension are of high public health importance. Brazil has already built mechanisms in the PHC sector to screen for some of the most epidemiologically relevant diseases. Some cancers, hypertension and diabetes have screening and prevention strategies, but more could be done to improve depth and scope of such strategies. Key priorities are to move towards population-based screening programmes for breast and cervical cancer, with a personalised approach and more communication strategies. In the area of diabetes and hypertension, Brazil will need to further develop disease management pathways with a people-centred perspective, integrating all health care providers across different sectors. Family health teams will need to have the right tools, capacities, and incentives to undertake these responsibilities. Last but not least, a more comprehensive information system based on registries, and allowing linking different data sources will also be important.

3.1. Introduction

In Brazil, as in other OECD countries, chronic non-communicable diseases such as cancer and cardiovascular diseases are of high public health importance. Diseases of the circulatory system was the leading cause of death in both Brazil and OECD countries in 2019 representing 27% and 31% of all deaths, respectively, followed by cancer representing 17% and 25% of all deaths. Diabetes in Brazil stands as the fifth cause of mortality with 5% of all deaths, higher than the 3% in OECD countries (Figure 3.1). This reflects the epidemiological transition from communicable to non-communicable diseases, where population ageing largely explains this mortality pattern but the prevalence of different risk factors and the performance of the health system play a relevant role as well.

Figure 3.1. Main causes of mortality in Brazil and the OECD, 2019



Source: OECD (2021^[1]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

In this scenario, Brazil, based on its strong PHC system, has already built mechanisms to screen for some of the most epidemiologically relevant diseases. Some cancers, hypertension and diabetes are on this list, for which screening and prevention strategies have proven to be effective and cost-effective, becoming a very good public health intervention. However, many challenges remain, for instance, related to escalating coverage and expanding to new diseases.

This chapter begins by setting out the current cancer burden in Brazil, while exploring the strengths and weaknesses of Brazil cancer screening system. Next, it analyses the impact of hypertension and diabetes in Brazil's population health and health system, and addresses hypertension and type 2 diabetes screening in PHC as important interventions to control these medical conditions. Throughout the chapter, a series of recommendations on the policy priorities for Brazil when it comes to enhancing screening coverage and promoting early detection of these conditions are provided.

3.2. Cancer in Brazil has been recognised as a major public health challenge but much can be done in terms of prevention in PHC

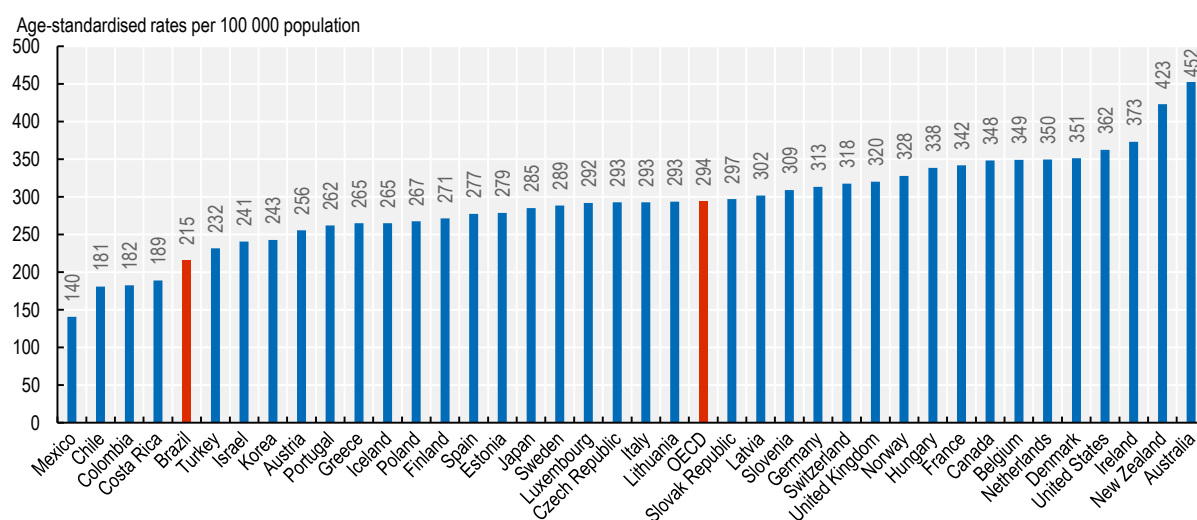
Cancer is the second highest cause of death across OECD countries, but its burden is increasing faster relative to the first cause of death, diseases of the circulatory system. The mortality rate of cardiovascular diseases has declined substantially over recent decades, while cancer mortality has also decreased but at

a slower pace (OECD, 2019^[2]). This section summarises the incidence and mortality rates of cancer and their changes over time in Brazil compared to OECD countries to illustrate the relative importance of public health policies in Brazil.

3.2.1. Brazil has a lower cancer incidence than the OECD but mortality has remained stable when it has decreased in the vast majority of OECD countries

Cancer incidence rates vary across OECD countries from over 400 new cases per 100 000 people in Australia and New Zealand, to less than 200 cases in Mexico, Chile, Colombia and Costa Rica. Brazil stands at 215 new cases per 100 000 people, below the OECD average of 294 (Figure 3.2). It is important to notice that cross-country variations in incidence rates reflect differences not only in new cancers occurring each year but also disparities in national cancer screening policies, quality of cancer surveillance and reporting.

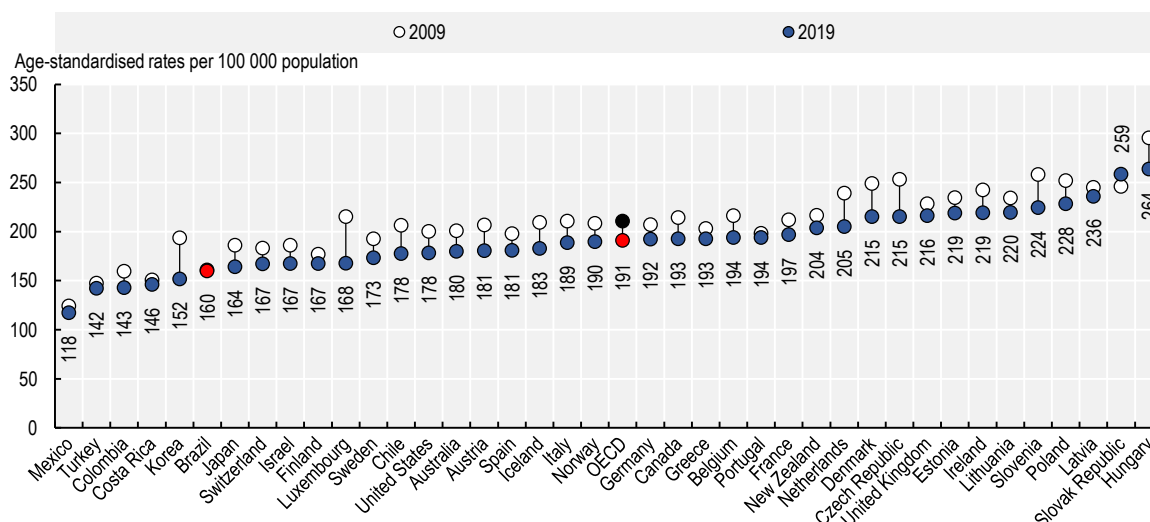
Figure 3.2. All cancer incidence in Brazil and OECD countries, 2020 (estimated)



Source: GLOBOCAN, (2020^[3]), International Agency for Research on Cancer (IARC), <https://gco.iarc.fr/>.

Despite having a relatively low rate of cancer incidence, cancer mortality in Brazil has not been reduced in recent years. Between 2009 and 2019, Brazil's cancer mortality passed from 161 to 162 deaths per 100 000 people, while the OECD average was reduced from 211 to 196. This decline was observed in most OECD countries, except in Chile and Turkey (Figure 3.3). The decrease of cancer mortality relates to early diagnosis, which puts a strong argument in favour of building effective screening schemes.

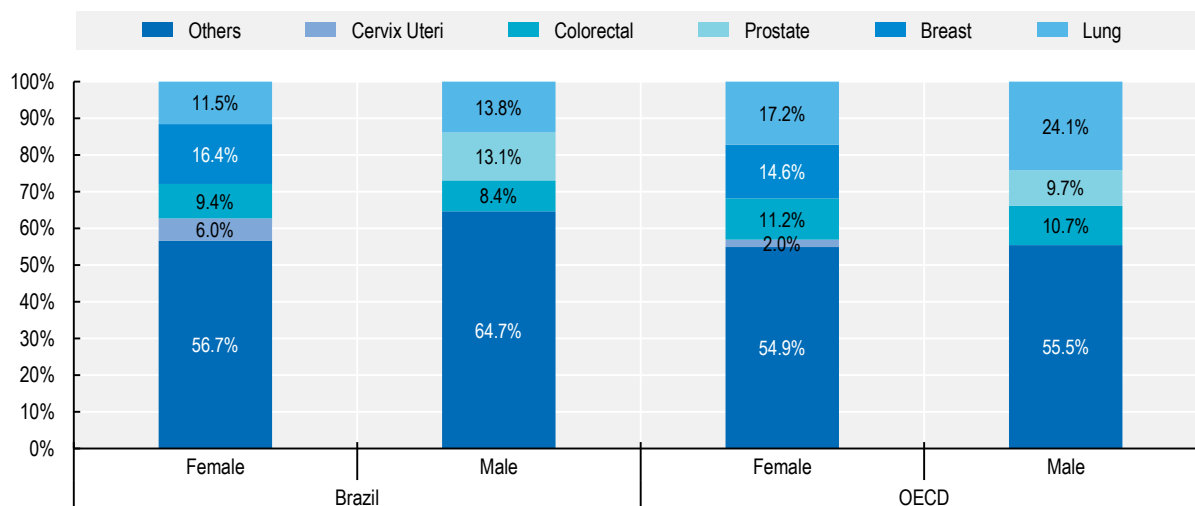
Figure 3.3. Cancer mortality in Brazil and OECD, 2009 and 2019 (or latest available year)



Source: OECD (2021^[1]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

In Brazil, among women, breast cancer takes the largest toll among Brazilian women with 20.6 deaths per 100 000 females, representing 16.4% of all women's cancer deaths in 2019. It is followed by trachea, bronchus and lung cancer with 15.7 deaths per 100 000 females (11.5%); colorectal and anus cancer with 13.1 deaths per 100 000 females (9.4%); and cervical cancer with 7.4 deaths per 100 000 females (6%). In OECD countries, trachea, bronchus and lung cancer are in first place among women and the most relevant difference with Brazil can be found in cervical cancer that only accounts for 2% of all women's cancer deaths. Among men, trachea, bronchus and lung cancer explains the largest number of deaths in both Brazil and the OECD, but the proportion is higher in the latter with 24.1% versus 13.8% in Brazil. Prostate and colorectal and anus cancer are the second and third causes of cancer deaths among Brazil's men, while in the OECD colorectal and anus cancer takes the second place (Figure 3.4).

Figure 3.4. Main causes of cancer mortality by sex in Brazil and OECD countries 2019 (or nearest year)



Note: Colorectal includes colon, rectum and anus cancer. Lung includes trachea, bronchus and lung cancer.

Source: OECD (2021^[1]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

As in OECD countries, age-standardised mortality rates in Brazil due to all cancers are higher among men than among women: 202.8 men's deaths per 100 000 males versus 135.8 women's deaths per 100 000 females. The gender difference in cancer mortality can be explained at least partly by higher prevalence of some risk factors among men.

3.2.2. Breast and cervical cancer have long-standing opportunistic screening programmes delivered mainly through PHC

The control of breast and cervical cancer in Brazil has decades of development, with National Cancer Institute (*Instituto Nacional de Câncer*, INCA) and the Ministry of Health as the main actors in terms of national planning, while PHC in municipalities have had the provision of care role. Importantly, cancer screening in Brazil is opportunistic, meaning that it does not happen within a population-based programme designed and managed at the central level to reach most of the population at risk, but as a result of a recommendation made by a health care provider during a routine medical consultation or by self-referral of individuals. A summary of the main historical developments of breast and cervical cancer screening is presented in Box 3.1.

Box 3.1. Almost 40 years of breast and cervical cancer screening in Brazil

The first actions towards breast and cervical cancer control can be traced back to 1984, when the Program of Integral Assistance to Women's Health (*Programa de Assistência Integral à Saúde da Mulher*), which proposed care beyond the traditional attention to the gravidic-puerperal cycle, included actions for the prevention of these two cancers. In 1986, an Oncology Programme was created and in 1991 was finally hosted in the INCA. In the 1990s and early 2000s, different pilot projects and programmes, mainly for cervical cancer were developed.

In 2005, the Ministry of Health launched the first National Oncology Care Policy (*Política Nacional de Atenção Oncológica*), with emphasis on cervical and breast cancer. The control of these cancers is now a priority on the country's health agenda and it is part of the Strategic Action Plan for Coping with Chronic Non-Communicable Diseases in Brazil, 2011-22 (*Plano de Ações Estratégicas para o Enfrentamento das Doenças Crônicas Não Transmissíveis no Brasil, 2011-22*).

In May 2013 occurred the latest update of the national cancer care policy, instituting the National Policy for the Prevention and Control of Cancer in the Health Care Network of People with Chronic Diseases in SUS (*Política Nacional para a Prevenção e Controle do Câncer, PNPCC, na Rede de Atenção à Saúde das Pessoas com Doenças Crônicas no âmbito do SUS*). That same year, the Cancer Information System (SISCAN) was created, a web platform version that integrates the Cervical Cancer Information Systems (SISCOLO) and Breast Cancer (SISMAMA).

More recently, in 2015 the guidelines for early detection of breast cancer, and in 2016 the national guidelines for cervical cancer screening and a manual for quality management of cytopathology laboratories were published.

Source: INCA (2020^[4]).

In Brazil's PHC, cervical cancer screening is performed through a cytopathological examination (Pap smear), which is offered to women aged 25 to 64 years and who have already had sexual activity. The recommended routine for screening in Brazil is to repeat the Pap smear every three years, after two consecutive normal examinations performed at an interval of one year. The repetition in one year after the first test aims to reduce the possibility of a false negative result in the first round of screening. The three-year periodicity is based on the WHO recommendation and the guidelines of most countries with an

organised screening programme (INCA, 2016^[5]). Table 3.1 summarises the characteristics of cervical cancer programmes in OECD countries.

Table 3.1. Target age and periodicity of cervical cancer screening in OECD countries, 2019

Every year	Every two years	Every three years	Every five years	Other
Czech Republic (15 and over), Greece (20 and over), Luxembourg (15 and over), Mexico (25-64), the Slovak Republic (23-64)	Australia (20-69), Japan (20 and over), Korea (30 and over), Latvia (25-70), Costa Rica (20-64)	Belgium (25-64), Chile (25-64), Denmark (23-50), France (25-65), Hungary (25-65), Iceland (23-65), Ireland (25-44), Israel (25-64), Italy (25-64), Lithuania (25-60), New Zealand (20-69), Slovenia: (20-64), Spain (30-39), Sweden (23-50), Switzerland (no age limit)	Denmark (51-64), Estonia (30-55), Finland (30-60), the Netherlands (30-60), Ireland (45-60), Norway (25-69), Spain (40-69), Turkey (30-65), England (25-64)	Canada: no periodicity (20-69). Germany: no periodicity (20 and over). USA: Every one, two or three years depending on the insurance coverage (21 and over).

Note: Target women's age in brackets.

Source: OECD (2021^[11]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

As for breast cancer, the Brazilian Ministry of Health recommends a biennial mammogram for women aged 50 to 69 years as screening strategy in PHC. In addition, the Ministry of Health recommends against teaching self-examination as a method of screening, and also against screening using ultrasound, thermography, tomosynthesis and magnetic resonance imaging. To assure an early diagnosis of breast cancer, the Ministry of Health endorses that SUS actors develop awareness strategies, suggests urgent referral to diagnostic breast cancer services of patients with specific red-flag signs, and that all the services are to be provided in the same centre (Ministério da Saúde and INCA, 2017^[6]). Table 3.2 summarises the target age and characteristics of breast cancer screening programmes in OECD countries.

Table 3.2. Target age in breast cancer screening programmes, 2018/19

Nationwide population-based		Population-based but not nationwide		Non-population-based
Wider age range (20 years+)	Narrower age range	Wider age range (20 years+)	Narrower age range	Wider age range (20 years+)
Australia (50-69), Belgium (50-69), Denmark (50-69), Finland (50-69), France (50-74), Germany (50-69), Hungary (45-65), Iceland (40-69), Israel (51-74), Korea (40+), Latvia (50-69), Lithuania (50-69), Luxembourg (50-69), the Netherlands (50-75), New Zealand (45-69), Norway (50-69), Poland (50-69), Portugal (45-69), Slovenia (50-69), Spain (50-69), Sweden (50-69)	England (53-69), Estonia (50-65), Ireland (50-64 but 50-69 by 2021), Northern Ireland (53-70), Wales (53-70)	Canada (50-69), Czech Republic (45+), Italy (50-69), Japan (40+), Mexico (50-69), Switzerland (50-70), Turkey (40-69)	Chile (50-64)	Greece (40+), the Slovak Republic (40-69) and United States (40 or 50+)

Note: Target women's age in brackets.

Source: OECD (2021^[11]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

The financing of these screening programmes is made through the Ministry of Health, which allocates resources to federated entities (states and municipalities), with these chiefly municipalities responsible for managing and providing the services. In 2020, a new financing model for PHC (the *Previne Brasil* Programme) was instituted combining financial resources from weighted capitation, pay-for-performance and resources for actions in specific contexts (see Chapter 4 of the report). Within the pay-for-performance component (15% of total funding), one of the seven initial target indicators rewards the teams that perform cervical cancer screening.

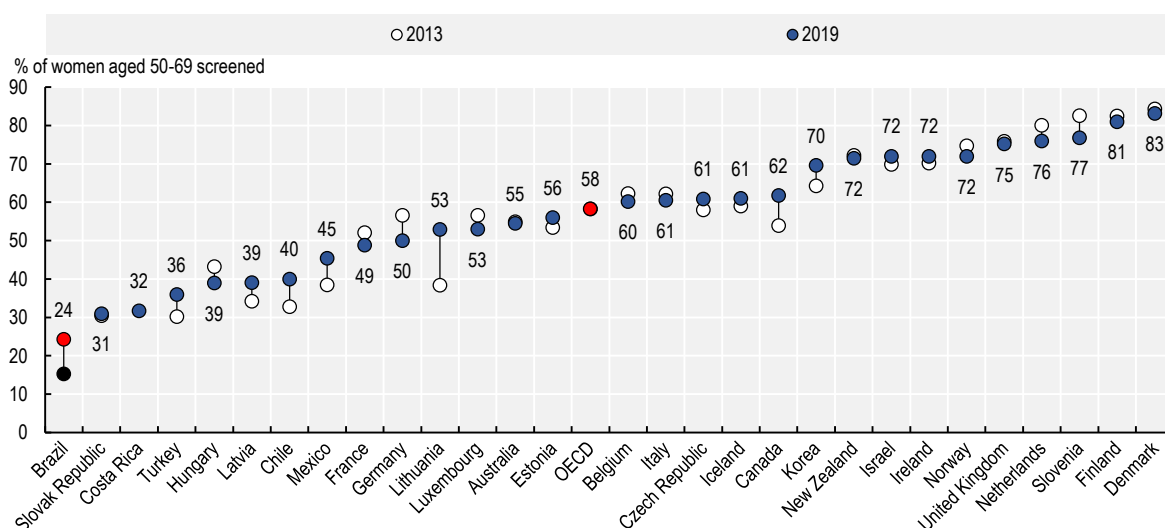
Regarding cancer information system, the PAINEL-Oncology is an instrument developed to monitor Law N°12.732, of 22 November 2012, which establishes a 60 days maximum waiting time for the start of

treatment for patients with proven malignant neoplasia. The information on PAINEL-oncology refers to the time to start the first oncological treatment calculated from the information of the dates of diagnosis and treatment recorded. It also presents the cases diagnosed through pathological examinations. Cervical and breast cancer diagnosis information has been available since 2013 since the implementation of the Cancer Information System (Sistema de Informação do Câncer, SISCAN). For the other cancers, diagnostic information arising from biopsy or anatomic-pathologic (except for cervical and breast) is available from May 2018, but from other sources it was possible to calculate the time of onset of the first treatment from 2013 as well. The data presented in the PAINEL-oncology refers exclusively to users who have a national health master card (DATASUS, 2021^[7]). In addition, a network of 17 Population-Based Cancer Registries, 16 located in capital cities and one of them in a non-capital city, provides crucial information about incidence, trends and survival (INCA, 2013^[8]).

3.2.3. Breast and cervical cancer screening needs to drastically increase coverage among target populations

Screening for breast cancer in Brazil has improved in recent years. Between 2014 and 2019, it increased from 15.2% to 24.2% among the target group of women between 50 to 69 years of age. However, breast cancer screening coverage in Brazil is very low when compared to OECD countries, standing below all OECD countries and well below the 58% of average coverage (Figure 3.5).

Figure 3.5. Breast cancer screening coverage in Brazil and OECD countries, 2013 and 2019

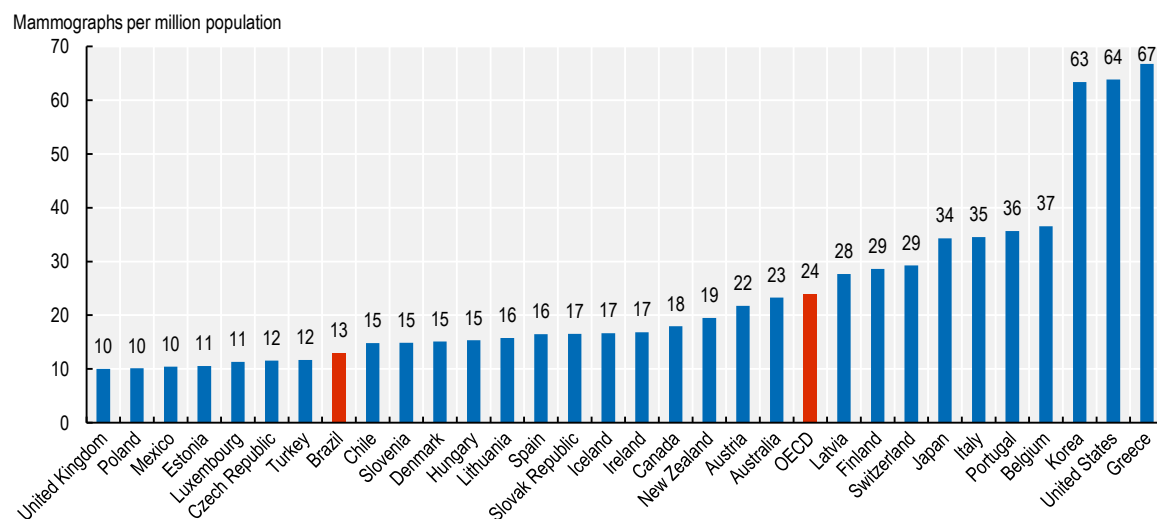


Note: Programme data is used for all countries. Brazil represent years 2014 and 2019.

Source: Ministry of Health for Brazil; OECD (2021^[11]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

One of the key enablers for conducting breast cancer screening is the availability of mammography machines. In 2020, Brazil had 13 mammography machines per million people, almost half than that of the OECD average of 24 and close to the lowest availability of 10 in Mexico, Poland and the United Kingdom (Figure 3.6). This finding shows that Brazil has space to put more investment into medical technologies aiming to improve equitable access for the population. At the same time, such expansion in access can be accompanied by the development of regulatory frameworks in the areas of registration, assessment and purchasing rules. The existence of updated clinical guidelines for breast cancer screening in Brazil is a step ahead in the necessary task of promoting rational use of diagnostic technologies aiming to reduce the use of unnecessary diagnostic tests and subsequent procedures and treatments.

Figure 3.6. Availability of mammography machines in Brazil and OECD countries, 2019 (or latest year available)



Note: Brazil represents year 2020.

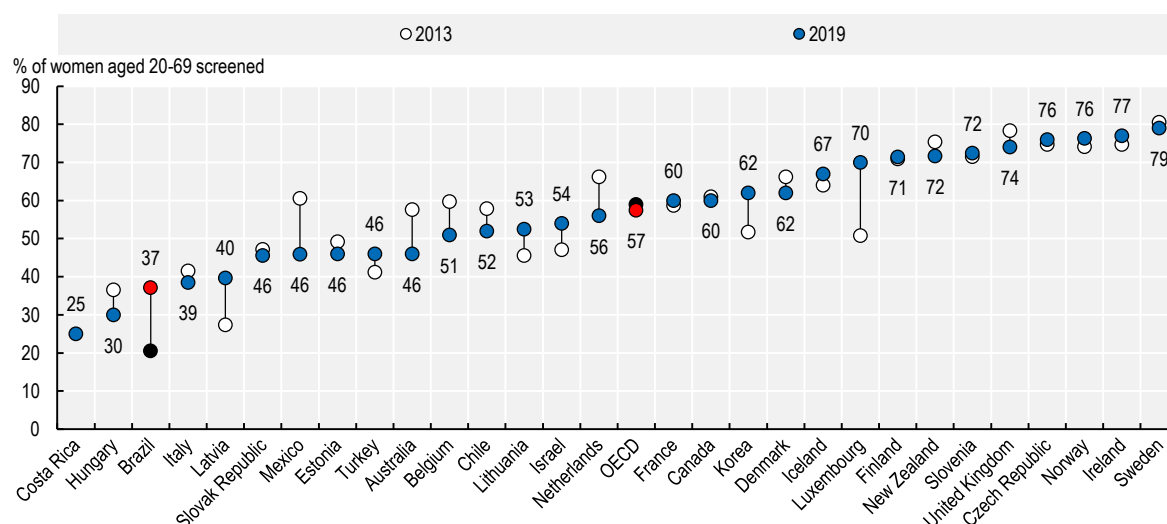
Source: Colégio Brasileiro de Radiologia e Diagnóstico por Imagem; OECD (2021^[1]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

A systematic review including 30 studies identified the main health system factors that led to a late diagnosis of breast cancer in Brazil (Da Costa Vieira, Formenton and Bertolini, 2017^[9]). Results highlighted problems related to the distribution of mammography machines, with a higher proportion of unused mammography machines in the North and Northeast of Brazil, and that most mammography occurred in the private sector. Another study analysing the National Health Survey 2013 found that 59.4% of women users of SUS and 83.9% of women with private health insurance declared having a medical request for a mammography. Having private health insurance, higher education level, and being white were positively associated with having this medical request (Silva et al., 2017^[10]). These findings highlight some of the problems with the health system organisation and with social inequities found in Brazil.

Cervical cancer screening in Brazil has substantially improved from 20.5% in 2014 to 37% in 2019. Yet, it also remains well below the OECD average coverage of 57% and below all OECD countries, except Costa Rica and Hungary (Figure 3.7).

A study that performed a trend analysis of the cervical cancer screening program's quality indicators in all Brazilian regions and states from 2006 to 2013, found that the estimated target population who underwent Pap testing was lower than that recommended by international guidelines in the North, Northeast and Central-West regions. The trends for this indicator remained stationary over the years in all regions of Brazil. Authors concluded that "the cervical cancer screening programme is still far from efficient" (Costa et al., 2018^[11]). Another study using data for 2008 found that rural areas were more likely to have never received a Pap smear screening, while North and Northeast regions showed the highest prevalence of never-screened women. Poorer, less educated, non-white skin colour and younger women were less screened as well, showing the patterns of social inequities (Martínez-Mesa et al., 2013^[12]).

Figure 3.7. Cervical cancer screening in Brazil and OECD, 2013 and 2019



Note: Programme data is used. Brazil represent years 2014 and 2019.

Source: Ministry of Health for Brazil; OECD (2021^[1]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

3.2.4. Brazilian society is quite active and key stakeholders are regularly involved in decision making around cancer control issues

Brazil has good examples of stakeholder involvement for cancer policies. Voices of local and regional stakeholders such as municipal and state administrations and health care providers are represented in institutional spaces such as the National Health Council, CONASS and CONASEMS. In addition, representatives of the civil society and users of SUS apply the so-called democratic social control by participating in the National Health Conference, the Councils of Rights and the Councils of Health. The last two have expression at the federal, state and municipal levels (Ministério da Saúde, 2013^[13]). Among the topics included, cancer regularly takes part of the agenda.

INCA develops several initiatives to engage civil society. The institution has a Consultative Council (Conselho Consultivo, CONSINCA) with representatives from technical-scientific entities related to cancer prevention and control; SUS service providers; SUS management councils; and SUS users. In recent years, INCA has organised conferences in different parts of the country with women's movements aiming to introduce the issue of cancer control into their working plans. Moreover, INCA has convened meetings with journalists with the objective of discussing the best ways to communicate information related to cancer control, including special meetings with radio managers of the North of Brazil, since in this region radios have a large audience in the population.

"All Together Against Cancer" (*Todos Juntos Contra o Câncer*) (TJCC, 2021^[14]) is a civil society movement that brings together representatives of different sectors, such as health managers, medical entities, hospitals, professional associations, researchers, media professionals, patient organisations, and others, committed to guaranteeing the patient's right to universal and equal access to health care. The movement develops several initiatives to assure the implementation of the National Policy for Cancer Prevention and Control, including the areas related to prevention and screening.

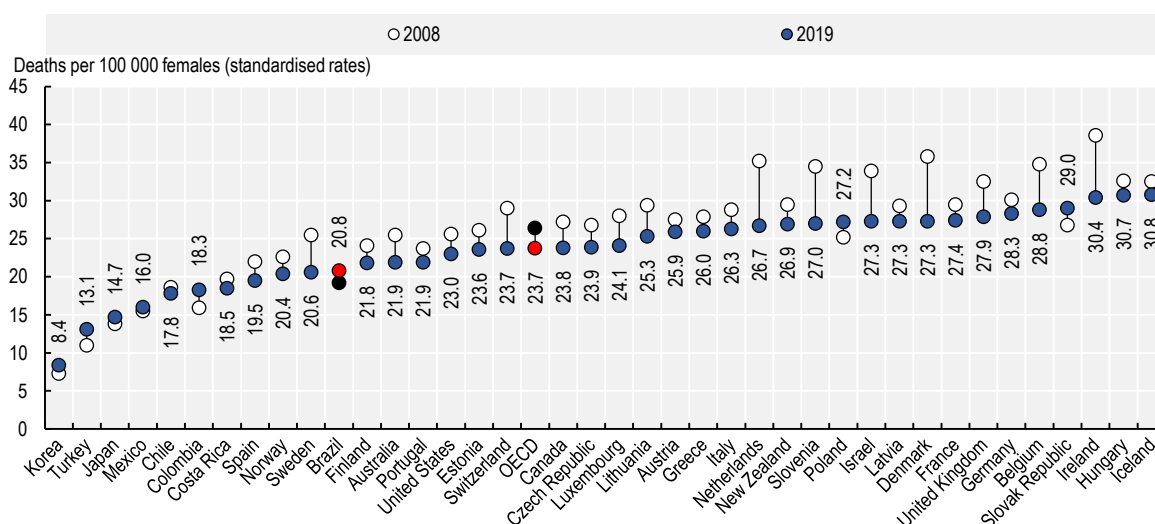
From the private sector, the "System S" carries out actions aimed at supporting promotion, prevention and treatment of cancer. The "System S" refers to nine institutions that are independently managed by business federations and confederations of the main sectors of the economy. Although they provide services of

public interest (e.g. education, social services, transport, etc.), these entities are not linked to any sphere of government, providing them a different space of action.

3.2.5. Despite prevention initiatives, breast and cervical cancer outcomes remain relatively poor with mortality increasing and survival decreasing in recent years

Between 2008 and 2019, breast cancer mortality increased from 19.2 to 20.8 deaths per 100 000 women in Brazil (+8.3%), which is the opposite of what happened as an average amongst OECD countries: a decrease from 26.4 to 23.7 (-10%). Only eight out of the 37 OECD countries experienced a growth in breast cancer mortality, with Turkey, Colombia and Korea having an increase of 15% or more in the period (see Figure 3.8). Despite Brazil's breast cancer mortality is lower than the OECD average, the increasing trend is worrisome.

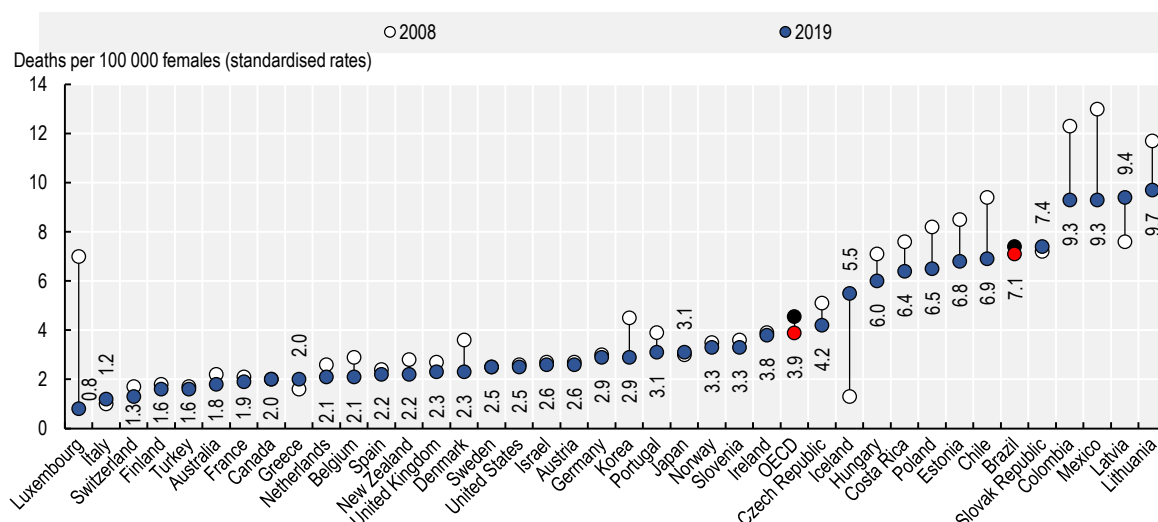
Figure 3.8. Breast cancer mortality in Brazil and OECD countries, 2008 and 2019



Source: OECD (2021^[1]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

The picture for cervical cancer mortality in Brazil seems more problematic. Between 2008 and 2019, mortality remained stable around 7.1 and 7.4 deaths per 100 000 women, compared to a decrease in OECD countries from 4.6 to 3.9 (-14.7%). Only seven out of 37 OECD countries experienced an increase, with Iceland, Greece, Italy and Latvia increasing by 15% or more (see Figure 3.9). In this case, Brazil's cervical cancer mortality is substantially higher than the OECD average.

Figure 3.9. Cervical cancer mortality in Brazil and OECD countries, 2008 and 2019



Source: OECD (2021^[1]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

These worrying trends are also observed in terms of 5-year net cancer survival. In all the selected cancer sites, Brazil has lower survival rates compared to OECD averages. Comparing the 2005-09 and 2010-14 periods, breast cancer survival dropped by more than two years in Brazil, while in the OECD it increased by one year. Colon cancer survival also decreased by more than two years, when in the OECD it increased by almost two years. Cervical cancer survival decreased by 11.3% and lung cancer survival declined by 11% in Brazil between 2000-04 and 2010-14, while in the OECD they both increased by 1.5% and 24.8%, respectively (Table 3.3).

Table 3.3. Trends in survival for breast, cervical, colon and lung cancer in Brazil and the OECD

Five-year net survival, population aged 15 and over

	2000-04	2005-09	2010-14
Brazil Breast	73.4	77.1	74.9
OECD Breast	81.0	83.3	84.3
Brazil Cervical	67.5	62.6	59.9
OECD Cervical	64.5	65.3	65.5
Brazil Colon	44.5	50.6	48.3
OECD Colon	56.1	59.4	61.2
Brazil Lung	9.1	8.5	8.1
OECD Lung	13.7	15.3	17.1

Note: Data for breast cancer considers only female population.

Source: CONCORD programme, London School of Hygiene and Tropical Medicine.

3.2.6. HPV vaccination has been very positive but it can be enhanced

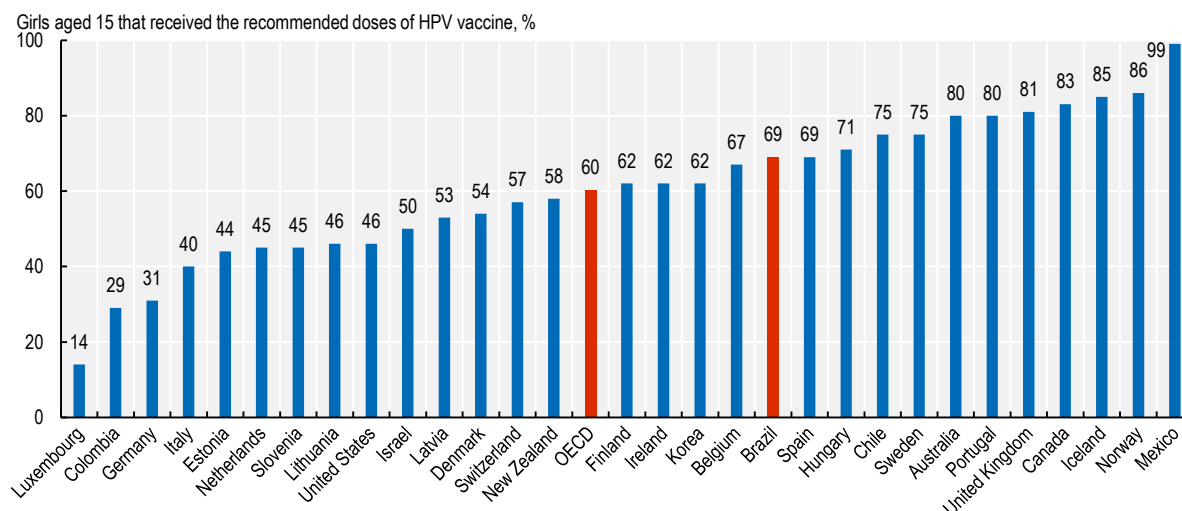
In 2014, the Ministry of Health, through the National Immunization Program, started the vaccination campaign for girls between 9 and 13 years against the Human Papilloma Virus (HPV). In 2017, girls aged 14 and boys between 11 and 14 years old were included as well. The vaccine is quadrivalent, which offers protection against HPV subtypes 6, 11, 16 and 18, preventing diseases such as precancerous lesions, cancer of the cervix, vulva and vagina, anus and genital warts. The vaccine is offered by SUS in

Basic Health Units and in vaccination campaigns in schools. The Practical Guide on HPV – Questions and Answers was launched by the National Immunisation Programme, in collaboration with INCA, seeking to clarify the main doubts on the subject (INCA, 2020^[15]).

In 2018, coverage of HPV vaccine in Brazil was 69%, above the OECD average of 60% (see Figure 3.10). However, it is below the target coverage of 80% that was defined by Brazilian authorities (INCA, 2020^[15]).

Figure 3.10. Human papilloma virus vaccine coverage in Brazil and OECD countries, 2018

Girls aged 15 that received the recommended doses of HPV vaccine, percentage



Source: Global Health Observatory: Explore a world of health data. (WHO, 2021^[16]), <https://www.who.int/data/gho>.

Although not yet available to be applied, ANVISA has already approved a new vaccine against HPV, which protects against nine types of virus. It is offered for girls from 9 to 14 years old and boys from 11 to 14 years old, for people with HIV – AIDS in the age group of 9 to 26 years old. The vaccination schedule is three doses (interval of 0, 2 and 6 months). Other age groups may have vaccines available at private services, if indicated by their doctors.

3.2.7. A national strategy for cancer prevention and screening in PHC would synergise efforts to improve cancer outcomes in Brazil

Brazil can develop a single national strategy addressing cancer prevention and screening in PHC. Scientific evidence and social participation can be two key guiding principles in order to move towards population-based screening programmes. This strategy can consider the renewal of existing screening programmes for cervical and breast cancer; the expansion to new cancer sites such as colon and lung; the introduction of systematic personalised screening invitation letters and reminders; the redesign of cancer information systems; and the use of information and communication technologies to improve public awareness and health literacy.

Brazil can move towards a population-based cervical cancer screening programme, while introducing HPV-DNA primary testing in PHC settings

In the area of cervical cancer screening, especially after the introduction of the HPV vaccine, Brazil could consider instituting primary testing for the DNA of oncogenic HPV types as novel method for screening.

This would ideally require moving towards a population-based screening programme, further analysing the current epidemiology in the country and conducting a complete health technology assessment to understand the different factors that will determine the efficacy and cost-effectiveness of implementing such an organised programme. It would be important to use reliable, validated HPV tests in qualified laboratories, accredited by authorised accreditation bodies (e.g. ANVISA) and in compliance with international standards. Training for health workers would be another crucial component, along with communication strategies for women in target population.

A recent systematic review focusing in low- and middle-income countries found that cytology-based screening was the least effective and most costly screening method, while self-collected HPV testing was cost-effective when it yielded population coverage gains over other screening methods. The cost of the HPV test and loss to follow-up were also important factors (Mezei et al., 2017^[17]). The European guidelines recommend the use of HPV primary testing within an organised, population-based programme for cervical cancer screening, avoiding co-testing (HPV and cytology primary testing) at any given age (von Karsa et al., 2015^[18]). The American Cancer Society 2020 guidelines also recommends HPV primary testing as the preferred screening method (Fontham et al., 2020^[19]).

Such a national strategy should include quality assurance in monitoring cervical cancer screening performance. The European guidelines define such a performance through achieving: (1) an invitation coverage of at least 95% of the target women; (2) an examination coverage of at least 70% (85% is desirable); and (3) a participation rate of at least 70% (85% is desirable) (von Karsa et al., 2015^[18]). Box 3.2 summarises the experience of Finland and Italy with implementing HPV testing.

Box 3.2. HPV primary screening test programme in Finland and Italy

In Finland, women of 30-60 years of age are invited to take part in cervical cancer screening every five years. Some municipalities also invite women aged 25 and/or 65 for screening. About 70% of those invited participate in the screening for cervical cancer. The conventional cervical screening test used in Finland has been the Pap test, while the HPV test is applicable to cervical cancer screening for women over the age of 35. Samples for HPV and Pap are taken from participants in the HPV screening. Following a positive HPV test, the Pap sample is also analysed, after which any further management required is decided. The screening HPV test must meet the requirements of the IARC and WHO. Before being implemented, the validity of new HPV tests must always be verified by comparing them with the most studied HC2 or PCR GP5+/6+ tests. A referral to colposcopy is given to about 1% of the participants. Additionally, about 5% are invited for a follow-up test before the next screening invitation. An annual review provides a comprehensive overview of the Finnish cervical cancer screening programme, presenting information about attendance, outcomes, time series and costs of screening. The history of cervical cancer screening in Finland and its impact on cancer burden is also discussed.

In 2013, based on a health technology assessment report, the Italian Ministry of Health included HPV test every five years as an option for screening programmes for women of over 30 years of age. Stand-alone HPV was defined as primary test, with cytology only to be used as triage test in HPV+ women. Women with negative findings in the cytology analysis are invited for repeat HPV testing after 12 months and referred to colposcopy if still positive. The National Prevention Plan 2014-18 put as an objective the full implementation of HPV-based screening by 2018.

Source: Finnish Cancer Registry (2020^[20]), Cervical cancer screening – Syöpärekisteri, <https://cancerregistry.fi/screening/cervical-cancer-screening>; Ronco (2020^[21]), Core elements of the new HPV-based cervical cancer screening programme in Italy, <https://www.hpvworld.com/articles/core-elements-of-the-new-hpv-based-cervical-cancer-screening-programme-in-italy>.

Self-sampling with HPV testing has been explored as a novel strategy to increase screening coverage and could be considered in Brazil, for instance, within pilot studies. A systematic review including 34 studies, mostly from high-income countries, found greater screening uptake among HPV self-sampling participants compared with standard of care (e.g. Pap smear, visual inspection with acetic acid, clinician-collected HPV testing) (RR: 2.13, 95% CI 1.89 to 2.40). Uptake was higher when HPV self-sampling kits were sent directly to women's homes (RR: 2.27, 95% CI 1.89 to 2.71) or offered door-to-door by a health worker (RR: 2.37, 95% CI 1.12 to 5.03). However, meta-analysis showed no statistically significant difference in linkage to clinical assessment/treatment between arms (RR: 1.12, 95% CI 0.80 to 1.57) (Yeh et al., 2019^[22]). The European guidelines state that the clinical accuracy of HPV primary testing on self-collected samples is sufficient to conduct pilot programmes (von Karsa et al., 2015^[18]), while the American Cancer Society did not include a recommendation about it because the Food and Drug Administration has not yet approved self-sampling (Fontham et al., 2020^[19]).

Brazil already has practical experiences with using HPV-DNA testing for screening. For instance, a study evaluated HPV-DNA screening in São Paulo, including 16 102 women, finding that high-risk HPV-DNA prevalence was 14.9%, whereas cytology abnormalities were found in 7.2% of the women. HPV DNA detected a significant number of patients with premalignant lesions missed by cytology and all 16 cervical intraepithelial neoplasia positive cases were HPV-DNA+. Authors concluded that HPV genotyping may reduce the burden of colposcopic referral and that HPV-DNA testing was shown to be feasible and advantageous over current cytologic screening in the public health system (Levi et al., 2019^[23]). Another study conducted in three Brazilian cities found that the most relevant barriers for a more comprehensive screening programme relates to excessive travel required to access clinics, inconvenient service hours, and lack of public education. In addition, cytology had many unsatisfactory results and low positivity, follow-up was incomplete at all steps and laboratories provided delayed cytology and biopsy results (3+ months). Finally, screening clinics did not arrange or track colposcopy referrals; colposcopy clinics received no information from referring clinics, and electronic medical records have not yet replaced paper recordkeeping, leading to errors and loss of medical records (Ribeiro et al., 2018^[24]). The identification of these and other barriers would be very important for policy design and an eventual implementation phase.

Breast cancer screening can evolve to a population-based organised programme and more tools can be provided in PHC

As in the vast majority of OECD countries (see Table 3.2), Brazil can develop a population-based breast cancer screening programme. In practice, this would require an effective call-recall structure, device a screening registry and safeguard robust quality assurance at all levels. The current process for population's registration in Brazil's PHC goes in the direction of individually identifying the eligible target population in each area served (e.g. municipalities), so it represents an opportunity for developing organised screening programmes.

The European guidelines on breast cancer screening and diagnosis describes seven characteristics of an organised screening programme (ECIBC, 2020^[25]): 1. a policy specifying target population; 2. screening method and interval; 3. an active invitation of the entire target population; 4. a team responsible for overseeing screening centres; 5. a decision structure and responsibility for health care management; 6. a quality assurance system utilising relevant data; and 7. monitoring of cancer occurrence in the target population.

Building capacity in PHC teams would be crucial. As stated by WHO guidelines (WHO, 2017^[26]), early diagnosis capacities requires guidelines or protocols and sensitise providers to cancer signs and symptoms to improve diagnostic accuracy; give sufficient time to evaluate patients and train practitioners in how to diagnose cancer including use of medical devices; promote a clear communication between the provider and the patient; and develop reliable referral mechanisms. In addition, mammography machines

distribution across the country should be also revised to achieve a balanced supply according to the population's need.

Brazil could build up from local experiences that have made progress towards an organised screening programme. For instance, a breast cancer organised screening programme in the community of the Andaraí in Rio de Janeiro targeting asymptomatic women aged between 50-69 years from SUS presented a mammographic coverage rate of 70%, suggesting that a population-based screening is feasible (Gioia et al., 2018^[27]).

Systematic personalised invitations and reminders from PHC teams could improve cancer screening coverage

As part of population-based organised screening programmes, Brazil could adopt a more systematic and personalised approach in inviting target populations, as has been implemented in many OECD countries. An increasing number of OECD countries send a personal invitation letter for cancer screening to each individual in the target group, issued through a registry in a systematic manner. In Brazil, however, there is no national initiative regarding such invitations, which in practice are only implemented in some local levels depending on the initiative of particular municipalities or PHC teams.

In the majority of OECD countries, the invitation letter includes information on the benefits and potential harms of cancer screening such as false-positive screening results, over-diagnosis and over-treatment, and asks for either signed or verbal informed consent for screening (IARC, 2017^[28]). This practice is in accordance with international recommendations as the WHO recommends organised population-based mammography screening programmes to provide information on both benefits and risks of mammography screening so that target women are able to make an informed decision before undergoing mammography (WHO, 2014^[29]). In many OECD countries, additional efforts are also made to invite people with positive screening results in the past for follow-up assessment, and in some countries such as Denmark, Finland, Germany, Ireland, Italy, the Netherlands, Spain, Sweden and the United Kingdom, appointment date is fixed for mammography in the letter to further facilitate access to cancer screening (IARC, 2017^[28]; ECIBC, 2020^[30]).

A systematic review identified the most effective interventions to increase participation in organised screening programmes. Interventions such as postal reminders, telephone reminders, PHC physician's signature on invitation letter, and giving scheduled appointment instead of open appointment were all effective to increase uptake of breast, cervical and colon cancer screening. Mailing a kit for self-sampling cervical specimens increased participation in non-responders as well (Camilloni et al., 2013^[31]). Another systematic review found that text messaging (short message service, SMS) appear to moderately increase screening rates for breast and cervical cancer and may have a small effect on colorectal cancer screening (Uy et al., 2017^[32]). Future studies would also need to examine the cost associated with each of the invitation options and its impact in order to identify the most appropriate cost-effective strategies that may be different across regions in Brazil.

Screening in Brazil's PHC could explore expanding to sites such as colon and lung cancer

As colorectal and lung cancer are among the most important cancers in the country (see Figure 3.4), Brazil could explore the possibility of creating national screening programmes for these cancer sites.

For colorectal cancer, which represents the third major cause of cancer mortality in Brazil for men and women, opportunistic screening is available. The recommendation in SUS is to prioritise early diagnosis actions and personalised approach for high-risk groups. Recognising Brazil's different epidemiological and health care network realities, further studies are still needed to support the analysis of the feasibility of introducing screening in these different contexts (INCA, 2019^[33]).

As reference, an increasing number of OECD countries have introduced population-based colorectal cancer screening programme for free to target population in recent years, and in many of these countries target population is the population of 50s and 60s years of age often with an upper age limit ranging between 64 and 74 (OECD, 2013^[34]). The European guidelines recommend “only organised screening programmes should be implemented, as opposed to case-finding or opportunistic screening as only organised programmes can be properly quality assured”. It also states that several issues should be considered: the legal framework, the availability and accuracy of epidemiological and demographic data, the availability of quality-assured services for diagnosis and treatment, promotional efforts, a working relationship with the local cancer registry, and follow-up for causes of death at individual level (Malila, Senore and Armaroli, 2012^[35]). The United States recommends that adults age 50 to 75 be screened for colorectal cancer, either with stool tests, flexible sigmoidoscopy, colonoscopy, and CT colonography (virtual colonoscopy) (USPSTF, 2021^[36]), varying depending on the patient’s history and health system availability (Bibbins-Domingo et al., 2016^[37]).

In relation to lung cancer, which represent the second cause of cancer deaths in Brazil (see Figure 3.4), general population screening is not currently recommended in the country and any examination should be discussed between the patient and the doctor (INCA, 2020^[38]). Lung cancer screening is not common in OECD countries, but it exists in Canada, Japan and the United States. The Canadian Task Force on Preventive Health Care recommends screening for lung cancer with three consecutive annual low-dose computed tomography (LDCT) scans in high-risk individuals adults aged 55-74 years who currently smoke or quit less than 15 years ago, with a smoking history of at least 30 pack-years (CTFPHC, 2016^[39]). In Japan, annual chest X-ray is recommended for people aged 40 and over, and sputum cytology is also recommended for smokers aged 50 and over who have smoked more than 600 cigarettes over their lifetime (OECD, 2019^[40]). In the United States, the USPSTF released an updated recommendation in March 2021. Lung cancer screening is recommended as an annual screening with LDCT in adults aged 50 to 80 years who have a 20 pack-year smoking history and currently smoke or have quit within the past 15 years. Screening should be discontinued once a person has not smoked for 15 years or develops a health problem that substantially limits life expectancy or the ability or willingness to have curative lung surgery (USPSTF, 2021^[36]).

Brazil could develop a thorough evidence-based analysis, including cost-effectiveness studies, to identify the potential harms and benefits of implementing colorectal and/or lung cancer screening in the country, along with appropriate definitions in areas such as eligibility criteria, screening intervals, existing local initiatives, monitoring schemes and quality improvement, infrastructure and equipment, and payment mechanism.

A redesign of cancer information systems can help manage screening programmes more effectively

Brazil has developed a monitoring system for cancer screening (SISCAN), cancer care (PAINEL-Oncologia) and population-level cancer epidemiology (Population-based cancer registry), which provides a good information infrastructure base. In addition, population-based surveys (Vigitel) regularly collect screening coverage and barriers to screening in view of seeking ways to increase screening coverage.

The main challenges for the cancer information infrastructure in Brazil, including screening information, relates to coverage, interconnectedness and feedback. Coverage aims to reach the full extent of the target population for each screening programme (including actions in the private sector). Interconnection challenges relate to linking existing information systems, for which an expansion of the national health card would be fundamental. Feedback to different actors in the health system, importantly to all PHC teams, is crucial to ensure that decisions and actions are informed by regular and updated information.

A comprehensive information system based on registries, capable of following each patient individually, is essential for efficient management of screening programmes and cancer care delivery. It can identify target

population who have and have not participated in the screening programmes, those who are monitored outside of the programme due to their previous diagnosis of cancer and/or genetic predisposition to specific cancer and those who do not consent to do screening. Using such a system, personalised invitations and reminders for cancer screening, which are important for increasing screening coverage, can be sent systematically to target population.

Integrated information systems generate data that need to be analysed periodically to assess the effectiveness of existing cancer screening protocols such as target group, screening frequency and/or methods and across population with different background. Brazil has started producing this data, which is also publicly available on line through DATASUS. Several OECD countries use information around cancer screening to improve quality of screening programmes as well. They provide feedback to individual providers and benchmarking is also possible. A systematic literature review supports these additional efforts as it found that provider assessment and feedback contribute to increased coverage for cervical, breast and colorectal cancer screening (Sabatino et al., 2012^[41]).

Better information and communication technologies could improve public awareness and health literacy

Alongside more personalised invitation to cancer screening, Brazil could develop further communication and information-sharing strategies to improve public awareness on cancer prevention and screening and health literacy of the population. Since 2002, the Ministry of Health of Brazil has been investing annually in actions to raise awareness of cancer prevention measures, for instance, through the October Pink Campaign. The campaign consists of lighting buildings and monuments conveying the message “Prevention is necessary”, while also promoting healthy lifestyle among women, encouraging HPV vaccination, overcome fear or stigma associated with cancer, among other topics.

Box 3.3. Sundhed.dk, the Danish e-health portal to support transparency and patient empowerment

Sundhed.dk, the Danish e-health portal launched in 2003, is the official website for the public Danish health care services and enables patients and health care professionals to find information and communicate. It is a public internet-based 24/7 portal that collects and distributes health care information among citizens and health care professionals. In a secure part of the portal the patient has access to:

- Personal health data on treatments and notes from hospital records, information about medication, laboratory test results, vaccination data and visits to the general practitioner (GP);
- Various e-services including making appointments with GP's, prescription renewals and electronic communication with the GP;
- Information on waiting times at all public hospitals and ratings of hospitals in terms of patient experienced quality;
- Patient networks and the sundhed.dk handbook for patients.

This portal brings the entire Danish health care sector together and provides an accessible setting for citizens and health care professionals to meet and efficiently exchange information. By serving both the citizens and the health professionals, the portals aim is to enable the two to achieve co-operation based on the same data. This empowers the citizen and gives the health professionals better tools to improve quality in care (OECD, 2013^[34]; European Observatory on Health Systems and Policies and Petersen, 2019^[42]).

Municipalities, with support from the State and the Federal Government, works to promote screening tests, through campaigns using printed media, social networks and local media. During determined periods in the year, some municipalities implement mobile units, which go to communities in the most distant parts of the country as a mean of increasing screening access, diagnosis and treatment. In addition, INCA has developed manuals or flyers that are sent to PHC teams every year for informing the public in the community and at work about healthy lifestyles to prevent cancer and the importance of early detection for cancer. INCA also develops videos and written material that is publicly disseminated in websites and social media.

As systematic sharing of information related to cancer is limited, particularly for cancer without screening programmes, Brazil could also better develop information-sharing strategies to help patients to seek care, including cancer diagnostic services, at the right place at the right time. In this context, an increasing number of OECD countries have developed information-sharing platforms to better involve and enable patients in taking care of their own health and navigating through health care systems for their conditions. For example, some OECD countries including Denmark (see Box 3.3), England and Estonia have developed a platform such as a website or e-Health account to share evidence-based information on health care for different diseases including cancer throughout patient pathway with a view to promoting health literacy of the population and to support them to seek health care including cancer screening, diagnosis and treatment appropriately in a timely manner.

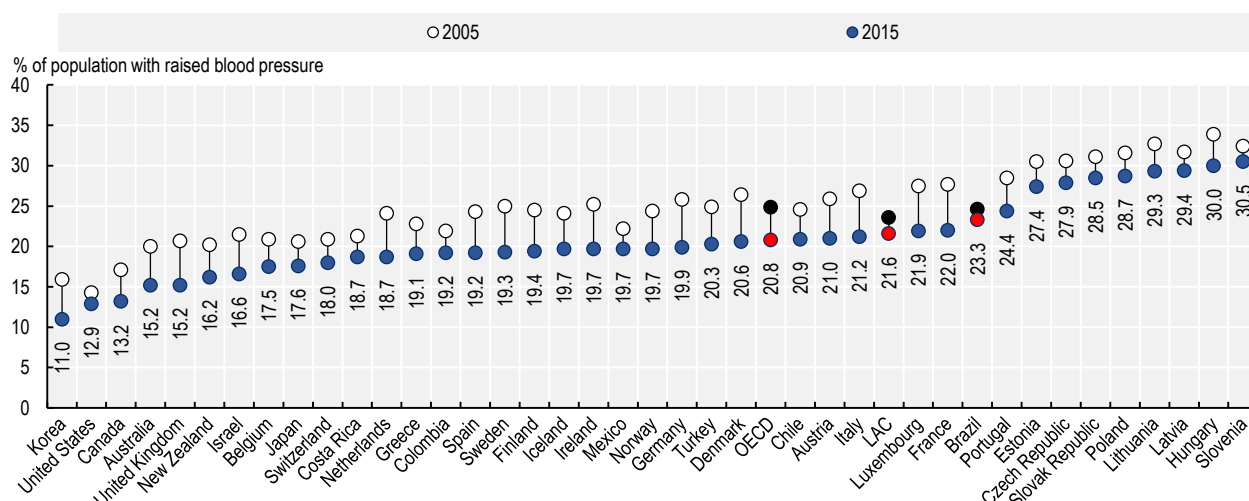
3.3. Screening for hypertension and diabetes can be further strengthened at the PHC level

3.3.1. Hypertension is one of the major health risk factors in Brazil

Hypertension or high blood pressure manifests by causing headaches, nosebleeds or difficulty breathing. If left untreated, hypertension can contribute to the development of more serious cardiovascular problems such as kidney disease, stroke and myocardial infarction. The absence of hypertension is a result of promotion efforts such as physical activity and healthy diets. When hypertension develops, it can be controlled with medication as well as with life style adjustments. This indicator is thus a proxy for both health promotion and medical services, usually delivered in PHC (WHO, 2019^[43]).

In 2015, the average prevalence of raised blood pressure in Brazil was 23.3%, higher than the OECD average of 20.8% and the LAC average of 21.6%. Between 2005 and 2015, all OECD countries reduced the prevalence of raised blood pressure with an average of -16.4% reduction. Brazil also reduced this prevalence but only by -5.3%, a decrease lower than all OECD countries (Figure 3.11). Changes in risk factors and improvements in detection and treatment of raised blood pressure have, at least partly, contributed to these general reductions, but other factors such as improvements in early childhood nutrition and year-round availability of fruits and vegetables, might explain it as well (Zhou et al., 2017^[44]).

Figure 3.11. Raised blood pressure among adults in Brazil and OECD countries, 2005 and 2015

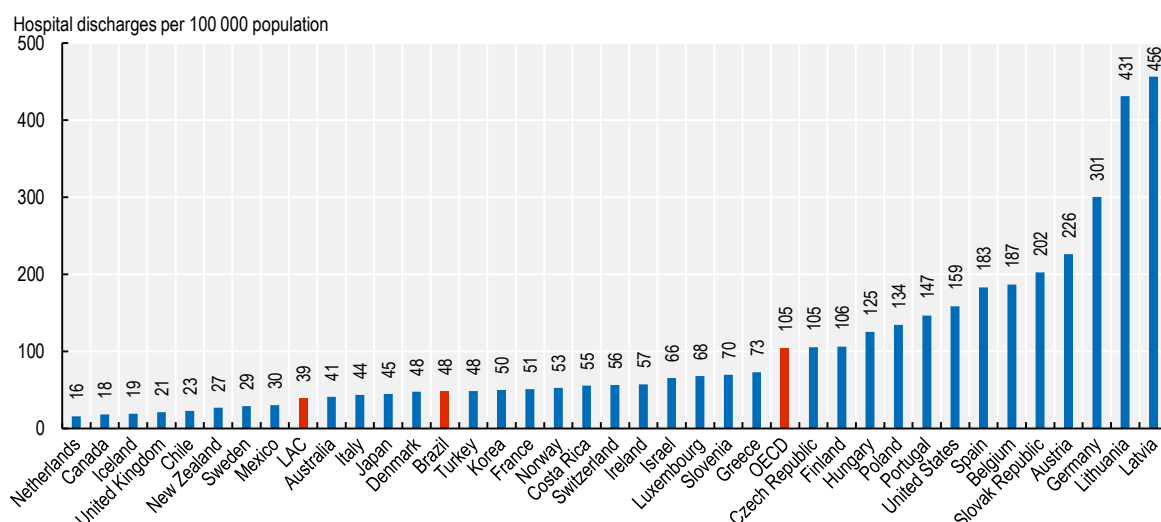


Note: Raised blood pressure (BP) defined as systolic BP ≥ 140 or diastolic BP ≥ 90 . Age-standardised estimate.

Source: Global Health Observatory: Explore a world of health data. (WHO, 2021^[16]), <https://www.who.int/data/gho>.

PHC is expected to keep people well by providing a consistent point of care over the longer term, treating the most common conditions, tailoring and co-ordinating care for those with multiple health care needs and supporting the patient in self-education and self-management. In this way, a good PHC performance can reduce the rates of hospital admissions of several diseases, including hypertension and diabetes. In Brazil, the rate hospital admission due to hypertension was of 48 per 100 000 people in 2019, which is less than half the OECD average of 105 but higher than the LAC average of 39 (Figure 3.12).

Figure 3.12. Hospital admission due to hypertension in Brazil and OECD countries, 2019 (or latest available)



Source: OECD (2021^[11]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>; Ministry of Health of Brazil.

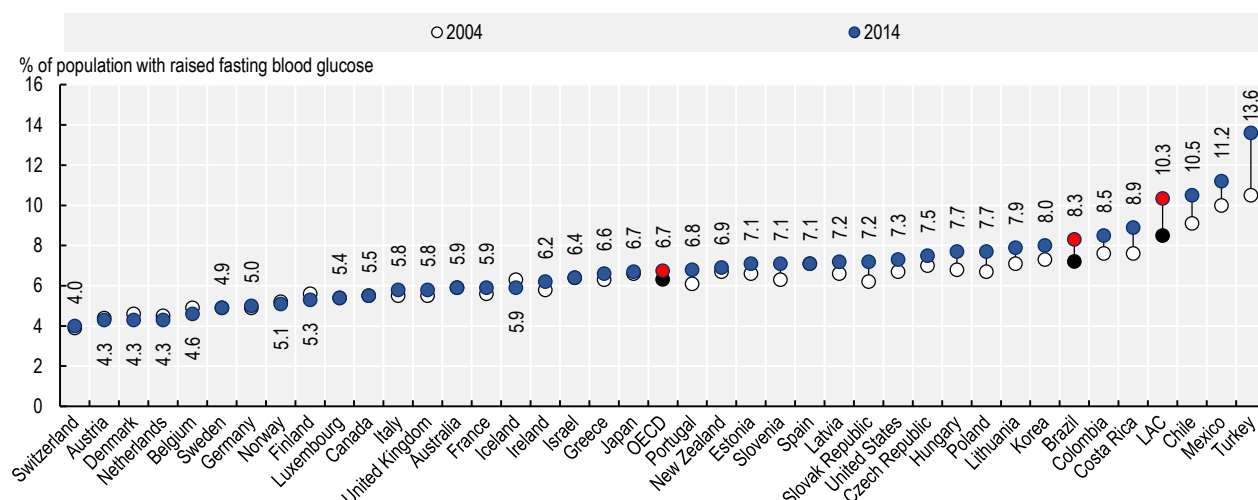
According to a recent study analysing high systolic blood pressure (HSBP) in Brazil, the prevalence of HSBP in 2017 was 18.9%, with an annual 0.4% increase rate, while age-standardised death rates attributable to HSBP decreased from 189.2 (95%UI 168.5-209.2) to 104.8 (95%UI 94.9-114.4) deaths per 100 000 from 1990 to 2017. However, the total number of deaths attributable to HSBP increased by 53.4% and HSBP raised from third to first position, as the leading risk factor for deaths during the period. Regarding total disability-adjusted life-years (DALYs), HSBP raised from fourth in 1990 to second cause in 2017 (Nascimento et al., 2020^[45]).

3.3.2. When compared to OECD countries, diabetes represents a higher burden of disease in Brazil

Raised levels of blood sugar can lead to the development of diabetes. Fasting blood glucose (FBG) contributes to diagnose and monitor diabetes, and can be under control because of effective treatment with glucose-lowering medication and as a result of health promotion activities. PHC has a central role in accomplishing these actions (WHO, 2019^[43]).

In Brazil, 8.4% of the population had raised FBG in 2014, above the 6.7% average in OECD countries. Importantly, between 2004 and 2014, Brazil recorded one of the largest increases in its population with raised FBG, increasing by 15.3% (Figure 3.13). Only Turkey, Costa Rica, the Slovak Republic and Chile had larger increases amongst OECD countries, evidencing a worrying trend in Brazil.

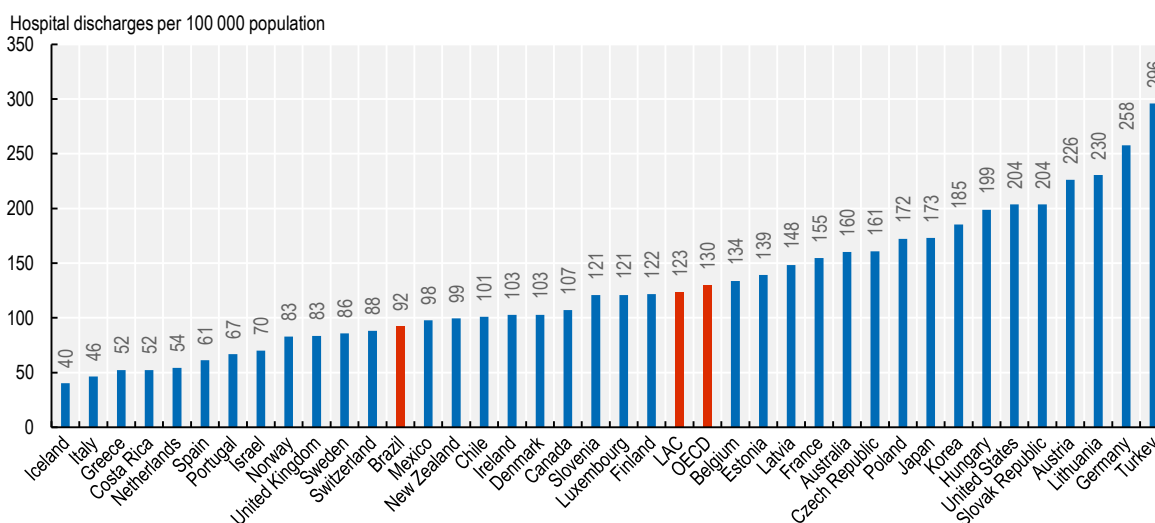
Figure 3.13. Raised fasting blood glucose among adults in Brazil and OECD, 2004 and 2014



Source: Global Health Observatory: Explore a world of health data. (WHO, 2021^[16]), <https://www.who.int/data/gho>.

As stated previously, a well performing PHC can control and reduce the number of hospital admissions due to diabetes. In Brazil, the rate of diabetes hospital admission in 2019 was of 92 admissions per 100 000 people, below the OECD average of 130 and the LAC average of 123 (Figure 3.14). The differences between Brazil and OECD countries in terms of access to hospital care and the profiles of chronic non-communicable disease burden remain as some of factors to be studied more in depth aiming to understand these differences (OECD/The World Bank, 2020^[46]).

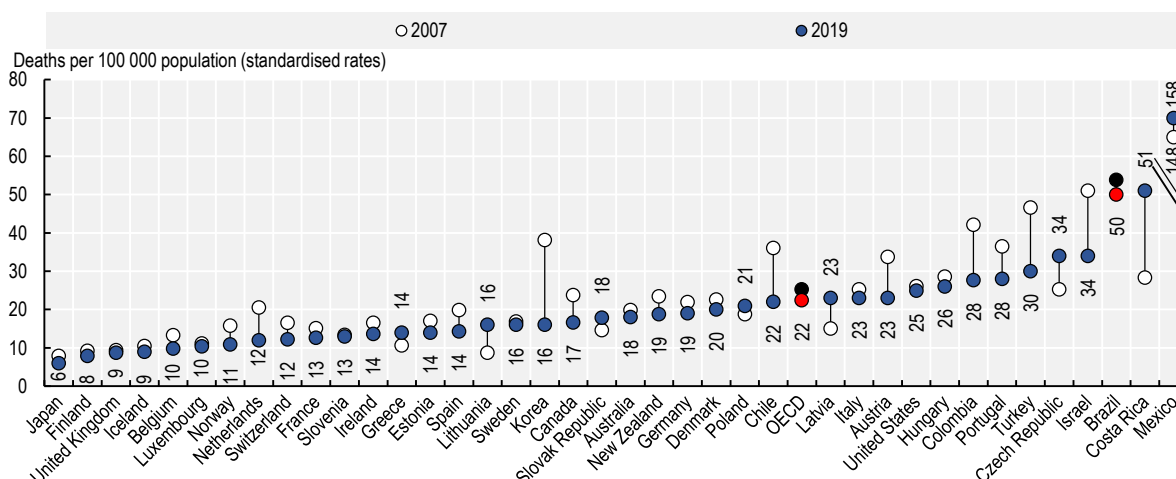
Figure 3.14. Diabetes hospital admission in adults in Brazil and OECD, 2019 (or nearest year)



Source OECD (2021^[1]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

Diabetes mortality in Brazil reached 50 deaths per 100 000 people in 2019, higher than the OECD average of 22 and only lower than Costa Rica and Mexico. Between 2007 and 2019, diabetes mortality was reduced by 9.8%, higher than the average reduction of 7.8% amongst OECD countries (Figure 3.15).

Figure 3.15. Diabetes mortality in Brazil and OECD countries, 2007 and 2019



Source: OECD (2021^[1]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>.

The impact of diabetes on the Brazilian health system was further studied by (Quarti Machado Rosa et al., 2018^[47]). They found that in 2014, 313 273 hospitalisations were due to diabetes in adults (4.6% of total adult hospitalisation), representing USD 264.9 million. The average cost of an adult hospitalisation due to diabetes was USD 845, 19% higher than hospitalisation without diabetes. Hospitalisations due to cardiovascular diseases related to diabetes accounted for the higher proportion of costs (47.9%), followed by microvascular complications (25.4%) and diabetes per se (18.1%). Another study analysing the burden of diabetes in Brazil, found that the overall estimated prevalence of diabetes in 2017 was 4.4% (95%UI

4.0-4.9%). While the crude prevalence of type 1 diabetes has remained relatively stable from 1990, type 2 prevalence has increased by 30% for males and 26% for females. In 2017, approximately 3.3% of all DALYs were due to diabetes and 5.9% to hyperglycaemia. Diabetes prevalence and mortality were highest in the Northeast region and growing fastest in the North, Northeast, and Central-West regions. Future projections suggested that the diabetes mortality burden will increase by 144% by 2040, more than twice the expected increase in crude disease burden overall (54%). By 2040, diabetes is projected to be Brazil's third leading cause of death and hyperglycaemia its third leading risk factor, in terms of deaths (Duncan et al., 2020^[48]).

3.3.3. Hypertension and diabetes screening in Brazil is embedded in a broader PHC check-up programme

Overall, basic health checks and risk screening for chronic diseases in Brazil relies on PHC family health teams, and predominantly on opportunistic screenings. Blood pressure and blood glucose are usually checked during PHC basic check-ups for chronic diseases. These check-ups also include other measures such as cholesterol, screen for cardiovascular disease based on age, family history and risk factors such as body mass index (BMI) that can help diagnose persons at-risk of chronic diseases, or diagnose chronic diseases in their earlier stages when they can be managed with fewer complications.

An official Ministry of Health screening guideline for chronic non-communicable diseases was published in 2010 and it is valid until today (Ministério da Saúde, 2010^[49]). For both hypertension and diabetes, screening target populations are quite wide. Screening for hypertension is recommended for adults (over 18 years of age) without the knowledge that they are hypertensive. It suggests screening every two years for people with blood pressure below 120/80 mmHg and annual screening if systolic blood pressure is between 120 and 139 mmHg or diastolic blood pressure between 80 and 90 mmHg. For diabetes, the guideline recommends screening in asymptomatic adults with sustained BP greater than 135/80 mmHg and does not apply to other screening criteria such as obesity, family history or age range. In addition, there are specific booklets (*Cadernos de atenção básica*) for the management of hypertension (Ministério da Saúde, 2013^[50]) and a recent protocol for diabetes (CONITEC, 2020^[51]), which also consider a screening component. For both diseases, the family health model proposes that trained health professionals work in an interdisciplinary manner, link with service users, and take on responsibility for ensuring the provision of comprehensive health care. For instance, for diabetes this includes from screening, classification, and diagnosis, along with an initial assessment, healthy lifestyle guidance and follow-up medical appointments. It also considers blood glucose control and treatment with or without medication, prevention and management of acute and chronic complications, care of feet and oral health care (Ministério da Saúde, 2013^[52]).

A study assessed hypertension care continuum in Brazil's PHC, finding that over one-third of the Brazilian adult population had measured hypertension or prior diagnosis. Nearly 90% of these had recent contact with the health system, but only 65% were aware of their condition. Only 62% of these regularly sought care for hypertension, but of these 92% received treatment. Hypertension control was 33% overall, but increased to 57% among those who passed through all stages of the care continuum, showing the importance of high-performing PHC teams (Macinko, Leventhal and Lima-Costa, 2018^[53]).

Regarding health information infrastructure, Brazil developed a specific registration and monitoring system for hypertensive and diabetic patients: HIPERDIA. It is managed by the Ministry of Health, through the Secretariat of Health Care Services, in conjunction with the the State and Municipal Health Secretariats. Municipalities that are members of the National Pharmaceutical Assistance Programme for Hypertension and Diabetes Mellitus send, through the HIPERDIA system, information on the registration and monitoring of patients with these diseases. All this information is processed and made available by DATASUS of the Ministry of Health (Ministério da Saúde, 2021^[56]).

Box 3.4. People with pre-existing chronic conditions risk more severe COVID-19 health outcomes

Across OECD countries, people with existing chronic conditions (including cancer, chronic kidney disease, chronic obstructive pulmonary disease, heart conditions, immunocompromised state, diabetes type 2, sickle cell disease and obesity) are the most hard-hit patient population in the COVID-19 pandemic. People with chronic conditions are facing a 'double threat': they are more vulnerable to complications and death from COVID-19, and they experience indirect health effects from disruptions in essential care:

- First, people living with certain chronic conditions are at higher risk of severe COVID-19 symptoms. Early in the pandemic, it was shown in China that older patients and those with chronic conditions were more likely to have a severe to critical COVID-19 condition, to show deterioration of their health condition and to die from COVID-19 (Zhang and et, 2020^[53]). Researches in the United States, Italy and the United Kingdom, confirm more severe COVID-19 outcomes and higher risk of hospitalisation among patients having underlying health conditions (OECD, 2021^[54]);
- Second, people living with chronic conditions, also face indirect impacts of the pandemic due to disruptions in continuity of care or foregoing care. The lock-down and fears of contamination in waiting rooms have led many patients to forego care, including those who require regular follow-up for chronic diseases, such as cancer (OECD, 2021^[54]). For example, in England, urgent referrals from PHC for people with suspected cancers decreased by nearly 80% in comparison with levels before the COVID-19 crisis. In France, the number of cancer diagnoses decreased by 35% to 50% in April 2020, as compared to April 2019. Preventive cancer screenings in the United States for breast cancer, colon cancer, and cervical cancer have also dropped between 86% and 94% in March 2020 compared to average volumes in 2017-19. In Australia, there were around 145°000 fewer screening mammograms conducted by BreastScreen Australia in January to June 2020 compared with January to June 2018.

Strong PHC has the capacity to mitigate these indirect effects during the pandemic, but also to reduce the pressure on the entire health system by providing comprehensive and preventive care.

Source: (OECD, 2021^[54]). Strengthening the frontline: How primary health care helps health systems adapt during the COVID-19 pandemic. OECD Policy Responses to Coronavirus (COVID-19).

In the context of the COVID-19 pandemic, and recognising that people with NCDs are at higher risk, the Ministry of Health issued specific guidelines for supporting this group of the population through PHC. The aim was to follow-up of this population with adequate regularity, aiming at clinical stability, reducing the chances of unfavourable outcomes during the pandemic period. Amongst other components, the strategy included the identification, registration and risk stratification of people with obesity, diabetes and/or systemic arterial hypertension through surveillance actions and timely diagnosis of the adherent population. In addition, the early identification and prioritisation of follow-up and monitoring of these individuals with influenza syndrome or with suspected or confirmed COVID-19. The response included multi-professional therapeutic assistance, the prevention of coronavirus transmission in individuals with obesity, diabetes and/or hypertension, and the implementation of intersectorial and community actions for promotion of health, considering the epidemiological situation of COVID-19 in each territory (Ministério da Saúde, 2020^[57]). These are all welcome strategies given that people with chronic non-communicable diseases are facing a 'double threat': they are more vulnerable to complications and death from COVID-19, and they experience indirect health effects from disruptions in essential care (Box 3.4).

3.3.4. Quality improvement and pay-for-performance schemes in Brazil's PHC have included hypertension and diabetes management

Checkup activities are part of the PHC Service Portfolio (*Carteira de Serviços da Atenção Primária à Saúde*), which receives financing from the Federal Government transferred to States and Municipalities, with these entities responsible for managing the provision of services. In 2011, the Ministry of Health created the Programme to Improve PHC Access and Quality (PMAQ), aiming at scaling up access and quality of care offered by means of financial incentives directed toward municipal health management, conditioned on formal agreements of commitments and indicator evaluation agreed between PHC teams, municipal health service managers, and the Ministry of Health (see Chapter 4). As part of PMAQ, the evaluation of FHTs included topics such as early detection of hypertension, laboratory exams for diabetes, and prescription refills for users on continued care/programs such as hypertension and diabetes without the need for marking medical appointments.

More recently, the new *Previne Brasil* Programme, implemented in 2020, includes a pay-for-performance component, which rewards performance linked to seven indicators (see Chapter 4). Two of them relates to hypertension and diabetes (Ministério da Saúde, 2020^[58]):

- Indicator 6: Percentage of people with hypertension having blood pressure checked in each semester.
- Indicator 7: Percentage of people with diabetes with a request for glycated haemoglobin.

Even though these indicators are not directly related to screening but to the follow-up of patients who already have the disease, it is important to highlight that patients who are diagnosed may benefit from this incentive. However, it would be important to monitor how this programme could influence screening activities for hypertension and diabetes in PHC.

3.3.5. Hypertension and diabetes secondary prevention in Brazil should be revised to improve the efficacy and efficiency of its screening activities

Brazil could develop population-based hypertension and type 2 diabetes screening targeting high-risk population groups

Several OECD countries have health check-ups for chronic conditions, usually targeted at particular groups of the population, undertaken periodically for example every five years, and sometimes provided by health care professionals other than doctors. In Estonia, health check-ups and guidance are provided by family nurses for people aged between 40 and 60 with hypertension or diabetes (Habicht et al., 2018^[59]), and in 2007, Korea introduced the National Screening Programme for Transitional Ages, targeting people at age 40 and 66 (Kim et al., 2012^[60]). In England, the NHS Health Check was introduced for people aged between 40 and 74 in 2009 and an invitation letter is sent every five years to those who do not already have diabetes, heart disease, or kidney disease or have not had a stroke, in order to screen them for the risk of developing chronic conditions including heart disease, stroke, kidney disease, type 2 diabetes, or dementia (available only for those above 65 and above). This check-up is often undertaken by a nurse or health care assistant (Gmeinder, Morgan and Mueller, 2017^[61]; NHS, 2018^[62]). Table 3.4 shows the recommendations for hypertension screening from selected evidence-based guidelines along with the latest recommendations in Brazil.

However, a recent review commissioned by WHO Europe found that the available evidence does not support the recommendation on systematic population-level screening for hypertension (Eriksen et al., 2021^[66]). Instead of recommending systematic screening, the preferred alternative would be targeting those in PHC who may be at a higher risk owing to age or the presence of a risk factor (i.e. case-finding). In Brazil, a study identifying the main risk factors for developing hypertension found that the prevalence increased with age, low educational level, increased body mass index and abdominal waist, and urinary

sodium excretion (Cipullo et al., 2010^[67]). This type of studies in the Brazilian population could help to define the target group in order to guide future hypertension screening recommendations.

Table 3.4. Selected recommendations for high blood pressure screening

	Population group	Screening interval	Persons at increased risk of high BP
Brazil (Ministério da Saúde, 2010 ^[49])	18 years and over, without known hypertension	• Every two years if BP below 120/80 mm Hg. • Every year if systolic BP 120-139 mm Hg or diastolic BP 80-90 mm Hg	None mentioned.
European Society of Cardiology and of Hypertension (Williams et al., 2018 ^[63])	18 years and over	• Every 1-5 years or when opportunity arise if BP below 120/80 mm Hg • Every 3 years if BP 120-129/80-84 mm Hg • Every year if BP 130-139/85-89 mmHg	In older patients (>50 years), more frequent screening should be considered for each BP category.
United States (USPSTF, 2016 ^[64])	18 years and over, without known hypertension	• Annual for adults aged ≥ 40 years and persons at increased risk for high BP. • Adults 18-39 years with BP < 130/85 mm Hg who do not have other risk factors, every 3-5 years.	Have high-normal blood pressure (130-139/85-89 mm Hg); those who are overweight or obese, and African Americans.
WHO HEART guidelines (WHO, 2018 ^[65])	18 years and over, during routine visits to PHC	• Every 1 to 5 years if normal • No mention of other interval	Have had a prior heart attack or stroke; have diabetes; have chronic kidney disease; are obese; use tobacco; have a family history of heart attack or stroke.

Note: BP: Blood pressure.

Table 3.5. Selected recommendations for type 2 diabetes screening

	Population and screening interval	Persons at increased risk of DM2
Brazil (CONITEC, 2020 ^[51])	• All adults over 45 years of age. • Adult overweight (BMI >25 kg/m²) and with one risk factor or moderate CVD risk. • If normal, screen every 3 years. • If pre-diabetes, screen every year.	Sedentary lifestyle; first-degree relative with DM; women with previous pregnancy with a fetus weighing ≥ 4 kg or with a diagnosis of gestational DM; hypertension or antihypertensive use; HDL cholesterol ≤ 35 mg/dL and/or triglycerides ≥ 250 mg/dL; women with polycystic ovarian syndrome; other clinical conditions associated with insulin resistance (e.g. obesity III, acanthosis nigricans); history of CVD.
Canada (Diabetes Canada, 2018 ^[69])	• Screen every 3 years in individuals ≥ 40 years of age • Screen every 3 years in individuals at high risk according to a risk calculator • Screen earlier and/or more frequently (every 6 to 12 months) in people with additional risk factors for diabetes	First-degree relative with DM2; Member of high-risk population (e.g. African, Arab, Asian, Hispanic, Indigenous or South Asian descent, low socio-economic status); History of prediabetes; History of gestational diabetes; History of delivery of a macrosomic infant; presence of end organ damage associated with diabetes; and presence of vascular risk factors (e.g. smoking), associated diseases (e.g. HIV), and use of drugs associated with diabetes (e.g. statins).
United States (USPSTF, 2015 ^[70])	• Adults aged 40 to 70 years who are overweight or obese. • Every 3 years for adults with normal blood glucose levels.	Family history of diabetes; history of gestational diabetes or polycystic ovarian syndrome; members of certain racial/ethnic groups (e.g. African Americans, American Indians or Alaskan Natives, Asian Americans, Hispanics or Latinos, or Native Hawaiians or Pacific Islanders) may be at increased risk for DM2 at a younger age or at a lower BMI.
WHO HEART guidelines (WHO, 2018 ^[65])	• Adults who are 40+ years old and who are overweight (BMI >25) or obese (BMI >30) • No screening interval mentioned.	Have had a prior heart attack or stroke; have diabetes; have chronic kidney disease; are obese; use tobacco; have a family history of heart attack or stroke.

Note: DM2: diabetes mellitus type 2. BMI: body mass index. CVD: cardiovascular disease.

Similarly as with hypertension, diabetes screening in OECD countries is normally embedded in more general health check-ups. In Australia, for example, PHC physicians can provide health assessment for people who are at risk of developing a chronic disease. This assessment is provided to people aged between 45 and 49 once if they have at least one risk factor (lifestyle habits or a family history) for developing a chronic disease such as type 2 diabetes or heart disease. The assessment is also provided to people aged 75 and over with an interval of 12 months or longer (Department of Health, 2014^[68]). In view of reducing the prevalence of lifestyle-related diseases including cancer, cardiovascular diseases and diabetes, Japan introduced the specific health check-up (Tokutei kenshin) to the population aged between 40 and 74 in 2008. All insurers in the Japanese health system are obliged to provide a specific health check-up to people in this age group every year as they are considered to have higher risks of developing lifestyle-related diseases. Insurers need to provide a nationwide standard set of health check-up items (OECD, 2019^[40]). Table 3.5 shows the recommendations for diabetes mellitus type 2 screening from selected evidence-based guidelines along with the latest recommendation in Brazil.

According to the most recent scientific evidence, too much screening (e.g. by having a too wide target population) probably will not improve population health outcomes and can be an inefficient use of resources. Therefore, Brazil can aim for well-targeted hypertension and type 2 diabetes screening population-based programmes, based on its national epidemiology, health system characteristics and cost-effectiveness analysis, as an effective way to identify and manage chronic disease.

Strengthen disease management pathways with a people-centred perspective

Hypertension and diabetes screening should be just the first step in a clear disease management pathway in Brazil. Chronic disease management pathways (also called integrated care pathways), which can be embedded on clinical guidelines, should be available for all high prevalence chronic diseases, to give guidance to health care providers and patients over expectations of the care that should be delivered and received. The two existing clinical guidelines in Brazil (Ministério da Saúde, 2013^[50]; CONITEC, 2020^[51]) represent a very positive step towards developing best practice guidelines with explicit pathways of care. For the moment these pathways are focused on information for health professionals in PHC, but could be expanded to integrate other care providers (for example specialists, or patient-support groups), and be produced in a patient-friendly format.

Pathways of chronic disease care, produced in both patient-facing and clinician-facing formats, should clearly establish the professional responsibilities of health professionals at different stages of disease. These pathways can be used to set quality expectations for chronic diseases such as diabetes and cardiovascular disease, and standardise quality of care across Brazil. The pathways can also be used to clarify expected roles for different care providers. In the area of diabetes, clinical pathways can help by: (1) assisting to systemically evaluate the patient's clinical presentation and risk factors; (2) choosing the most appropriate behavioural and medication interventions; and (3) providing a timeline for patient follow-up and monitoring. In short, clinical pathways will assist in the co-ordination of care and follow-up (Wong, 2017^[71]).

Figure 3.16. NHS England RightCare Pathway: Diabetes, summary table

The National Opportunity	5 million with non-diabetic hyperglycaemia Most receive no intervention	940, 000 undiagnosed Type 2 diabetes	>50% of diagnosed receive no structured education within 12 months of diagnosis	60% of Type 1 and 40% of Type 2 are not completing care processes	Few areas have high quality Type 1 services embedded	30% of hospitals don't have multi-disciplinary foot teams	National variation in spend and safety issues on non-elective admissions
Service component	<u>Risk Detection</u>	<u>Diagnosis and Initial Assessment</u>	<u>Structured Education Programmes</u>	<u>Annual Personalised Care Planning</u>	<u>Type 1 Specialist Service</u>	<u>Service Referral and key relationships</u>	<u>Identification/ Management of admissions by Inpatient diabetes team</u>
Interventions	Cross Cutting: 1. Shared responsibility and accountability 2. Participation in NATIONAL DIABETES AUDIT 3. Consistent support for patient activation, individual behaviour change, self-management, shared decision making 4. Integrated multi-disciplinary teams						
	NHS Diabetes Prevention Programme	Protocol for diagnostic uncertainty	Education programmes (including personalised advice on nutrition and physical activity)	9 recommended care processes and treatment targets	Type 1 Intensive specialist service	1. Triage to specialist services 2. RCA for major amputations	Inpatient diabetes team, shared records, advice line
Target outcomes	Decreased incidence of Type 2 diabetes	Improved detection	Better diabetes management and reduced complications	Reduced variation in completion of care processes	Reduced risk of Microvascular complications	Year on year reduction on major amputations	Reduction in errors in hospitals, reducing LOS
The evidence	Intensive behaviour change can on average, reduce incidence of Type 2 diabetes by an average of 26%	Diabetes prevalence model for local authorities and CCGs	Improved health outcomes and reduction in the onset of diabetic complications in both Type 1 and Type 2 diabetes	Control of BP, HbA1c and cholesterol reduces risk of macro and micro vascular complications	Type 1 services deliver year on year improvements in blood glucose control	MDFT and supporting pathway reduces risk of complications	Young Type 1 and older Type 2 diabetes patients have higher rates of non-elective admissions

Source: NHS England (2018^[72]), NHS RightCare, Diabetes pathway, <https://www.england.nhs.uk/rightcare/products/pathways/diabetes-pathway>

England has developed more complex pathways, which include expectations at different stages of the disease, the roles for different care providers, key interventions and target outcomes (NHS England, 2018^[72]). England's 'NHS RightCare Pathway: Diabetes' includes, for example, an expectation that care planning and an annual review take place for patients with both Type 1 and Type 2 diabetes, and the Pathway includes links to supportive documentation to help with care planning. In England, the expectation is that a lot of diabetes care is provided by multidisciplinary teams in community care settings, and the Pathway includes details of the services that the team would usually provide (patient education, pregnancy advice, foot protection team) (see Figure 3.16).

Alongside establishing clear chronic disease management pathways, there is a need to ensure that other levers within the system are effectively aligned with the pathway. This includes line up payment and reimbursement incentives, as well as ensuring that health professionals have the tools and capacities they need to undertake the responsibilities expected of them.

Create more capacity in PHC for screening and management of hypertension and diabetes through expanding the role of health workers

To strengthen prevention capacity and impact, Brazil should look to increase capacity in PHC. One way that several OECD countries have found to increase capacity relates to task shifting or expanding the role of other health workers (Box 3.5). Exploring whether there are ways for other health workers – for instance nurses, nutritionists or pharmacists – to play a role in delivering some of these key prevention activities is certainly a possibility for Brazil.

Box 3.5. Advanced nurse practitioners and community pharmacists in France

As in other OECD countries, France has extended the role of nurses and pharmacists, which is seen as a key policy lever to improve access in underserved areas where the number of PHC physicians is decreasing. The new decree establishing the profession of Advanced Nurse Practitioner (Infirmière en Pratique Avancée) was issued in June 2018. The Advanced Nurse Practitioner will work within a PHC team to manage patients having chronic conditions and take the lead in prevention and co-ordination.

In parallel, the role of community pharmacists is gradually increasing. Community pharmacists are allowed to perform three rapid diagnostic orientation tests: the capillary blood glucose test for diabetes screening; the oropharyngeal tests for influenza; and the group A streptococcal tonsillitis test. The objective is to determine if antibiotic treatment is necessary and if a visit to the doctor is required for a prescription. The community pharmacist can also participate in punctual screening programmes for chronic obstructive pulmonary disease (COPD). Lastly, pharmacists in France are now allowed to perform flu vaccination.

Source: OECD (2020^[78]), Realising the potential of PHC, <https://dx.doi.org/10.1787/a92adee4-en>.

For example, when it comes to diabetes, nurses could have the capacity and training to conduct an initial assessment of the patient, including ensuring the capacity to do blood sugar testing in all practices and working together with doctors and other team members in follow-up activities. In general terms, nurse-provided care has shown equivalent or better quality of care across a large range of clinical outcome measures for chronic conditions, including diabetes (Maier, Aiken and Busse, 2017^[73]). A systematic review found that nurse-led care was at least as effective as physician-led care in providing secondary prevention to patients with chronic conditions, where 84% of outcome parameters showed no statistically significant differences, and for the remaining 16%, secondary prevention outcomes improved in the nurse-group, primarily among patients with diabetes. In this patient group, nurse-led care was superior in preventing the onset of heart disease and it lowered cardiovascular risk (Martínez-González et al., 2015^[74]).

Community pharmacists can also be considered for task shifting policies, and in most OECD countries its role has evolved substantially in recent years. Pharmacists can play a key part in giving advice to patients and supporting them to navigate their health needs and manage their care. Recent systematic reviews have shown that pharmacist involvement in care may improve health outcomes for chronic conditions such as diabetes and hypertension (Pousinho et al., 2016^[75]), while it can turn to be cost-effective for health systems (Wang, Yeo and Ko, 2016^[76]). In Brazil, pharmacists do not seem to have a large presence in PHC teams, even stating that pharmacists' isolation in PHC prevails (Barberato, Scherer and Lacourt, 2019^[77]). Therefore, such a policy would require a thorough analysis and planning to define the best way to invest on further introducing community pharmacists in Brazil's PHC teams.

The health information system needs to be developed further to support targeted population-based screening for better monitoring and evaluation

Screening and other health care activities for hypertension and diabetes have the potential of generating a wealth of valuable health information, which could be used both for better managing population health, designing and targeting more effective public health interventions, and for research. However, HIPERDIA, Brazil's health information system for hypertension and diabetes in PHC, does not collect information about screening and it is not capable of capturing the pathway of care for each patient either (e.g. referrals, waiting times, consultations with specialists, laboratory tests, medications, etc.). It is also hard for PHC teams to review and act on this information as they do not receive regular feedback, and patients cannot

have access to his personal health data. It is also difficult for the system to link with other databases such as hospitalisations, death registries, etc.

Many OECD countries have nationally standardised digital medical records and they are able to use up-to-date data for monitoring and evaluation of the health system by automatically extracting data from electronic clinical records. For example, 13 countries including Canada, the Czech Republic, Denmark, Finland, Israel, Korea, New Zealand, Norway, Singapore, Sweden and the United Kingdom (England, Scotland and Wales) regularly link data from different national datasets such as hospital and mental hospital in-patient data, cancer registry data and mortality data. Using these linked data, these countries are able to actively monitor health care quality and health system performance.

In order to monitor and further develop secondary prevention strategies, Brazil needs to develop a health information system, which allows linking different data sources. Experiences of other OECD countries in developing a strong health information system within a data governance framework that protects patients' health information privacy, which is laid out in OECD (2019^[79]), would be useful for Brazil in following such paths.

A stronger health information system could also facilitate streamlining the efforts to invite target population for hypertension and diabetes screening. Currently, these screenings are largely opportunistic and there is no information about who is receiving them. Moreover, it is not possible for municipalities or PHC teams to identify whom to send the invitations. The current process for population's registration in Brazil's PHC represents an opportunity for developing organised screening programmes. Brazil could send personalised invitation letters, without duplication, only to those who are in the target population, and not currently undergoing treatment, and to those who have not followed up on worrying results from a previous check-up. These targeted personalised approaches are considered more effective in recruiting people in need of health check-ups (Bunten et al., 2020^[80]).

In the future, Brazil could consider bundled payments as a mechanism to improve management of chronic conditions in PHC

Brazil has recently innovated on its payment scheme in PHC with the *Previne Brasil* programme. For the future, another alternative to explore are bundled payments, in particular, for chronic conditions. These consist of one payment per patient with a chronic illness to cover the cost of all health care services provided by the full range of providers during a specific time period. It has been introduced in OECD countries to incentivise co-ordination of care between providers, or provide a broader set of care (e.g. education, regular checks, occasional specific checks). Bundled payments can encourage collaboration within and across care settings, contribute to greater standardisation of care for example by requiring adherence to quality criteria, and can strengthen data availability by requiring the collection of monitoring indicators or integration of data systems across care settings, and control overall costs (OECD, 2016^[81]).

Although the design and characteristics of bundled payments differ between OECD countries, the models developed in Australia and Canada could be of particular interest to Brazil and others. In these countries, the bundled payment accounts for patient complexity, which is an important prerequisite to encourage the participation of PHC providers (OECD, 2020^[78]).

3.4. Conclusion

In Brazil, all-cancer mortality has not been reduced in the past decade, while breast and cervical cancer have reduced its odds for a longer survival after being diagnosed. This finding goes in the opposite direction from the outcomes obtained in OECD countries, calling for an urgent action to reverse these trends. Although breast and cervical cancer screening have increased in recent years, its coverage remain well

below the recommended standards, limiting its potential to contribute to better cancer outcomes. In order to tackle the burden of cancer, Brazil has strengthened cancer care governance and delivery, however, more still needs to be done, with an emphasis in promotion and prevention of cancer. There is scope to expand the coverage of screening – notably by developing population-based programmes, introducing more systematic personalised invitation for cancer screening, and developing better communication and education strategies by using new technologies. Brazil could also look to expand efforts to new cancer sites such as lung and colorectal, as they represent a high burden of disease, deserving further analysis of the potential benefits of investing in national screening programmes. Meanwhile, developing a more comprehensive monitoring system for cancer screening, with information at the individual level of each patient, would help Brazil in both the administration of the cancer screening programmes (e.g. assuring effective target population coverage) and be a rich source of data to periodically assess the effectiveness of existing cancer screening protocols.

On the other hand, hypertension and diabetes also represent a burden for Brazil's population health and health system. For example, the prevalence of both raised blood pressure and of raised fasting blood glucose is higher than the average of OECD countries. For diabetes, mortality has not been reduced in past years in Brazil, on the contrary to what has been observed in OECD countries. Therefore, Brazil could strengthen its PHC check-up activities by moving towards population-based hypertension and type 2 diabetes screening programmes, targeting well-defined high-risk population groups. This goes in hand with further developing clear disease management pathways, where PHC teams and each individual patient can take appropriate actions to manage its disease. The current clinical guidelines and the registration of patients in PHC are good starting points to continue in this direction. There is also a clear scope for PHC nurses to play a more active role in patient education, screening promotion and disease management, while community pharmacists might be further introduced in targeted actions to manage these chronic conditions. Health information systems needs to be further developed in order to provide more and better information to both PHC teams and patients, which would allow improving clinical management decisions in PHC and will empower patients to take a leading role in its own prevention and control actions.

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4 Quality and outcomes in primary health care

Looking to expand access to high quality PHC, Brazil has taken important steps to improve the distribution of doctors, develop new forms of service organisations, introduce new financing models, and implement a range of quality initiatives, well aligned with the experiences of OECD countries. However, the Brazilian PHC system is still characterised by a relatively low population coverage, large disparities in care quality, and a weak referral system. PHC has also traditionally provided few low-complexity procedures, and is not comprehensive enough to meet evolving patients' needs. Tackling these challenges requires strengthening the gatekeeping system while expanding the range of services provided by family health teams. Efforts are also needed to ensure implementation of quality initiatives throughout the country. Greater guidance and support from the federal government will be necessary to help municipalities with low capacity.

4.1. Introduction

Brazil recognised that a strong PHC is the foundation of a health system that is effective, efficient and responsive to patients' needs. Over the past decade, Brazil has implemented a wide range of reforms to develop the system accordingly and improve both care quality and the range of services provided by family health teams (FHTs). These include the development of standards of care, guidelines, health care services portfolio, the establishment of pay-for-performance programmes and the production and publication of data underpinning PHC. High level analyses however suggest significant scope for improving the quality and outcomes of PHC service delivery. While the burden of risk factors for health and of chronic conditions is growing in Brazil, compliance with preventive care norms, and the detection and management of chronic conditions give some cause for concern. In addition, PHC has traditionally been characterised by a low supply of health care services, offering few low complexity procedures and surgeries.

This chapter assesses Brazil PHC system with regards to the quality and outcomes of care provided. It starts with an overview of recent reforms to expand PHC and to foster quality improvement. The chapter then points to its performance in examining some indicators of PHC quality across OECD countries. The chapter concludes with some key suggestions to secure higher quality and outcomes of PHC services.

4.2. Recent reforms in PHC in Brazil

Following the implementation of the Family Health Strategy in 1994 (described in Chapter 2), several important reforms were introduced from 2011 to 2019 to continue to expand PHC in Brazil. The reforms intend to restructure PHC through new regulations to improve the distribution of doctors, to develop new forms of service organisation and new financing models for PHC.

4.2.1. Considerable efforts have been made to recruit and retain professionals in more remote areas

Several initiatives have been taken in Brazil over the past decade to recruit and retain professionals in more vulnerable areas. The *Inland Health Care Programme (Programa de Interiorização de Trabalho em Saúde)* implemented from 2001 to 2004 first intended to improve the distribution and training of doctors and nurses in order to ensure their provision in poor and remote parts of the country. However, only 469 doctors joined the programme. Another federal strategy was the Valuing Primary health Programme (*Programa de Valorização da Atenção Básica – PROVAB*) introduced in 2011 to encourage health workers to spend a year working in PHC in areas with a shortage of these professionals (ordinance No. 2, 087, of 1 September 2011). The programme provided for the payment of a federal scholarship of BRL 10 000 a month. The programme also entails a mandatory 12-months theoretical-practical postgraduate course and the completion activity in the PHC units. To guarantee the quality of the service provided, medical doctors were evaluated, on a quarterly basis, by managers and institutions, in addition to conducting self-assessments. Well-evaluated participants received an additional 10% score on their medical residency exam. While the programme helped to attract more doctors in remote areas, the number was not enough to meet the needs of the target municipalities. Medical professionals showed a limited interest for the programme. In 2013, the programme hired 3 800 doctors, who were allocated in 1 300 different municipalities (Pereira et al., 2016^[1]).

The Programme *Mais Médicos para o Brasil* (More Doctors Programme [MDP]) was then instituted as a federal government initiatives in 2013 to address shortfalls in the supply of doctors in PHC and diminish regional inequalities in health. The MDP was a supply side intervention with three prongs that complement each other, with a particular emphasis on strengthening PHC provision in underserved communities (Stein and Ferri, 2017^[2]; WHO, 2018^[3]):

- (1) Transfer of funds to municipalities to strengthen PHC infrastructure;
- (2) Improving access to and quality of medical school training; and
- (3) Recruitment of Brazilian and foreign physicians to work within FHS teams.

Priority municipalities for participation in the programme were ones with population living in extreme poverty, population living in indigenous reserves, municipalities with very low human development index, or situated in areas of extreme poverty and with rural communities in the Northeast and North Regions (Pereira et al., 2016^[1]).

The Ministry of Health paid the monthly salary of doctors enrolled under the programme. The agreement stated that Brazil paid about BRL 10 000 (EUR 2 126; USD 2 321) a month for each professional. Food, lodging and transportation costs were covered by the municipality. Doctors had a three-year contract, renewable once (Biernath, 2020^[4]). It is not mandatory for the MDP physicians to take the national competency exam that is required for other physicians to practise medicine in Brazil. Instead, the MDP physicians are required to complete a mandatory three-week training course provided by the Brazilian health authorities before they could be deployed in municipalities. In addition, all MDP physicians are required to attend brief training sessions organised by Brazilian authorities in regular intervals. Online-learning courses are also available to the MDP physicians.

Overall, the MDP led to positive improvements. The programme has allowed to make considerable investment in the infrastructure of PHC units in all participating municipalities with the provision of new equipments and refurbishments (Pereira et al., 2016^[1]). More recent evidence confirms that the MDP successfully channelled additional resources for strengthening the PHC infrastructure, with the MDP resources leading to the construction of 3 496 new basic health units and refurbishment of 3 417 units by 2015 (Santos et al., 2017^[5]). According to Biernath (2020), more than 26 000 PHC facilities were built or renovated throughout Brazil.

In addition, the MDP have contributed to improvements in access to medical training. Data obtained from the Higher Education Census suggest that the MDP achieved its medium-term objective of expanding the number of new vacancies by 2017. An estimated 92% of the new medical undergraduate vacancies created from 2010 and 2018 took place after the MDP was launched in 2013 (Figueiredo et al., 2021^[6]). It allowed to create 11 500 new places for medical training and 12 400 places for medical residency, opened to both Brazilian and foreign doctors (WHO, 2018^[3]).

Lastly, evidence shows that the MDP has improved incentives for health professionals to work in underserved areas, and recruiting medical professionals from outside the country. As Brazilians were slow to accept posts in underserved communities (only 938 doctors applied for 15 460 job openings in the first months of the programme) (Biernath, 2020^[4]), co-operation agreements between Brazil and Cuba facilitated by the Pan American Health Organization (PAHO) and the World Health Organization (WHO) have ensured an adequate provision of doctor. More than 20 000 Cuban doctors have been mobilised to the Brazilian health system (representing two-third of the professionals who joined the programme) (WHO, 2018^[3]). In total, 18 240 job positions were created during the MDP. Among these, 8 332 were filled by Cubans, 4 525 by Brazilians who had graduated locally, 2 824 by Brazilians who had graduated abroad, and 451 by doctors of other nationalities (Biernath, 2020^[4]).

Hone and colleagues (2020) suggested that the MDP was associated with a 12% increase in the number of PHC physicians by its fifth year of implementation, with more pronounced effects among municipalities that had lower levels of physician availability at the start of the programme (Hone et al., 2020^[7]).

Between 2013 and 2014, the number of municipalities with fewer than 0.1 doctors per thousand inhabitants decreased by 75%, from 374 to 95 (Pereira et al., 2016^[1]). More generally, the number of municipalities with a shortage of doctors fell from 1 200 in 2013 to 777 in 2015 (Biernath, 2020^[4]).

In addition, the improved supply of medical doctors in remote regions in Brazil had prevented a total of 521 000 hospital admissions in 2015, generating large savings for the Brazilian health system (Biernath, 2020^[4]). The MDP led to reductions in adult hospitalisations primarily due to infectious gastroenteritis, bacterial pneumonias, asthma, kidney and urinary infections, and pelvic inflammatory disease (Maffioli et al., 2019^[8]), though it fell short of reducing hospitalisations due to other preventable conditions like hypertension (Özçelik et al., 2020^[9]). Amenable mortality was also significantly reduced following the MDP, with greater benefits in municipalities prioritised for doctor allocation and where doctor density was low before implementation (Hone et al., 2020^[7]). The positive impact of the MDP includes improved patient satisfaction with public health care services and better doctor-patient relationship (Biernath, 2020^[4]) (Chapter 5 of this report provides additional details on the MDP).

The More Doctor Programme is still in place today but it will be gradually replaced by a new federal programme called *Programa Médicos pelo Brasil* (see Chapter 5).

4.2.2. New configurations of care have been established to promote greater co-ordination and integration of care

The family health support centres (*Núcleos de Apoio à Saúde da Família*, NASF) were originally created in 2008 to expand the capacity of the Brazilian health care system to meet the increasing needs of the population, and move from vertical organisations of health care to horizontal modes of organisation.

The 2017 PNAB further encouraged the development of NASF to expand the scope of PHC for users and provide higher resolution of PHC through greater care co-ordination and longitudinally. NASFs were renamed Extended Family Health and PHC Units (NASF-AB). The organisation is structured on a territorial basis, depending on local needs. NASFs-AB should operate in an integrated manner to provide clinical, medical and pedagogical support for the PHC professionals (Barros da Silva, Carlos da Silva and de Araujo Oliveira, 2020^[10]). NASFs- AB are multi-disciplinary teams, expected to function in an integrated manner. The overarching objective is to support FHTs and PHC teams for specific populations (Street Offices, river team) through supporting them in the management of health problems, as well as broadening the range of services offered in the community (Ministerio da Saude, 2021^[11]).

Three types of NASFs-AB are recognised:

- NASFs-AB 1 includes 5 to 9 FHTs and/or PHC teams for specific populations (street consultancy team, riverside team, and river). Minimum 200 hours per week (minimum 20h and maximum 80h weekly workload)
- NASFs-AB 2 includes 3 to 4 FHTs and/or PHC teams for specific populations (street consultancy team, riverside team, and river). Minimum 120 hours per week (minimum 20h and maximum 40h weekly workload)
- NASFs-AB 3 includes 1 to 2 FHTs and/or PHC teams for specific populations (street consultancy team, riverside team, and river). Minimum 80 hours per week (minimum 20h and maximum 40h weekly workload)

The NASF-AB composition is defined based on the health needs of the territory and the supported family or PHC teams. It can be composed of social workers, physical educators, pharmacists, physiotherapists, speech therapists, nutritionists, psychologists, gynaecologists, obstetricians, psychiatric doctors, geriatric doctors, among others.

Activities carry-out by NASFs-AB include clinical care in the PHC facilities, home visits, multi-sectoral actions, health prevention, health education, case discussions with the teams and joint construction of Unique Therapeutic Projects (Barros da Silva, Carlos da Silva and de Araujo Oliveira, 2020^[10]) (Sales et al., 2020^[12]). Multi-disciplinary teams working in the NASF-AB have opportunity to share problems,

exchange knowledge and practices, and define together possible interventions through the Community of Practices (Comunidad de Practicas).

NASFs are guided by the following priorities:

- It is a team formed by different professions and specialties;
- It is recognised as a specialised support for PHC, but it is not outpatient specialties or hospital service;
- It should be available to support scheduled and emergency health situations;
- It performs actions in collaboration with the FHT or the PHC team to increase the supply of health services offered in PHC settings.

Existing evaluations of NASFs-AB show overall positive results (Barros da Silva, Carlos da Silva and de Araujo Oliveira, 2020^[10]; Sales et al., 2020^[12]). The actions developed by the NASFs were positively perceived by the multi-disciplinary teams. Nurses and physicians acknowledged that the multi-professional formation allowed for a greater provision of services, and with greater co-ordination and communication between the team members (Sales et al., 2020^[12]). A NASF team in the city of Recife (Pernambuco, Brazil) has been found to improve the quality of referral, and also to generate greater care co-ordination and continuity (Barros da Silva, Carlos da Silva and de Araujo Oliveira, 2020^[10]).

The technical Note of 27 January 2020 presents the new configuration of the Expanded Center for Family Health and Basic Health Care (NASF-AB). Given Brazil's vast territory and the heterogeneity of local environments, this technical note gives the flexibility to municipal managers to organise and define the team composition, the work-week hours, and team arrangements.

Another interesting development in Brazil is the establishment of the health care networks *Rede de Atenção à Saúde* (RAS) (CONASS, 2020^[13]). Established in 2011, RAS are large health care networks aimed at reducing health fragmentation. FHTs and PHC teams have a key role to play here, in being identified as the hub to co-ordinate care between and across the health sector.

RAS are similar to territorial communities of health professionals aimed at improving patient-centred care. They are organised by *Regiões de Saúde* (Health Regions) which are agglomerations of municipalities, with the objective of integrating organisation, planning and provision of health care services at regional level (from PHC, mental health, public health, emergency care to specialised care and hospital care). RAS are agreed by Municipalities and States by the *Comissões Intergestores*, according to population health needs, regional and municipal capacities (in terms of operational structure including support, logistical and governance systems), and the health care model.

There are thematic *Rede de Atenção à Saúde* including:

- Maternal and Child Health Care Network (Rede Cegonha)
- Urgency and Emergency Care Network (Rede de Atenção as Urgencias)
- Health care network for people with chronic diseases (Rede de Atenção à Saúde das Pessoas com Condições Crônicas) – such as cancer, overweight or obesity;
- Health care network for people with disabilities (Rede de Cuidados à Pessoa com Deficiência)
- Mental health care network

CONASS has been actively working on the development of RAS over the past years, notably with the development of *Planificação da Atenção Primária à Saúde* (PAPS) and *Modelo de Atenção às Condições Crônicas* (MACC) (CONASS, 2020^[13]; CONASS, 2018^[14]). PAPS and MACC provide technical and managerial guidance to support health managers and workers in states and municipalities to establish and organise RAS. PAPS consists of a set of workshops, tutorials and short-term trainings for health professionals and the managerial teams of the states and municipalities. The development of MACC in Brazil is based on the well-known Kaiser Permanente Pyramid Population Management, the Wagner

Chronic Care Model, and the Dahlgren-Whitehead “rainbow model” of social health factors (CONASS, 2018^[14]).

The use of PAPS and MACC was piloted across several municipalities, for instance in Curitiba (Parana), Santo Antonio do Monte (Minas Gerais) and Taua (Ceará) (see Box 4.1). Evaluations of pilot programmes have been positive in terms of better management of chronic conditions through greater integration of primary and specialised health care (CONASS, 2018^[14]; CONASS, 2020^[13]).

Box 4.1. Establishment of RAS in Taua (Ceará)

A new configuration of PHC was developed in the municipality of Tauá (Ceará) with the establishment of a RAS in 2014. The project was led by CONASS in partnership with the Secretary of Health of the State of Ceará, Municipal Secretary of Health of Tauá, School of Public Health of Ceará and Fiocruz Ceará.

The project began through the use of the PAPS that proposes new way of organising health services in this municipality. The development of RAS were also supported by several workshops with the participation of all health professionals, and the use of the MACC to identify at risk population including children under 2 years old, pregnant women, hypertensive and diabetes patients.

Risk stratification was used to better understand the health and risk profiles of the community, to undertake proactive management of patients’ needs and define intervention in primary and specialised care before the occurrence of any adverse outcomes.

Teams’ actions are based on indicators defined by the management, such as:

- preventive actions for congenital syphilis (early detection of infection for pregnant women, timely notification and treatment of pregnant women and partners, appropriate referral to the maternity)
- updating immunisation for elderly people;
- better monitoring of chronic conditions in line with recommended clinical practice guidelines, and care co-ordination with secondary care for high-risk and very high-risk patients.

Evidence of the impact of this RAS in Ceará is positive, with reduced complication from chronic conditions; reduced mortality rates from cardiovascular disease, improved quality of health care service and greater supply of PHC services.

Sources: CONASS (2020^[13]), Estudos sobre a planificação de atenção à Saúde no Brasil – 2008 a 2019: uma revisão de escopo.

4.2.3. Brazil has embarked on an important reform to finance PHC

Federal financing of PHC has recently changed with the Ordinance 2 979 (12 November 2019) which defines the *Previne Brasil*. Up until 2019, the federal transfers to the municipalities was based on a mechanism for regular and automatic transfer of federal funds. Every municipality received per capita funds per year to cover the cost of basic health care activities and incentives for implementing programmes recommended by the Brazilian Ministry of Health. Federal transfer consisted essentially of:

- Two capitation components -“PAB Fixo” and “PAB Variável”. The PAB Fixo “fixed component” was multiplying a municipality-specific value with the number of residents in the catchment area of municipalities. In 2019, this fixed value varied among municipalities between BRL 23 and 28 depending on some few socio-economic factors such as GDP per capita and the poverty ratio. The variable capitation component depended on the accreditation of FHT, implementation of some

strategies and priority programmes such as the Family Health Strategy or the Community Health Agents Programme;

- A voluntary pay-for-performance bonus based on the National Programme for Improving PHC Access and Quality (PMAQ) (described in the next section);

In 2020, a new PHC funding model – *Previne Brasil* – has come into effect. This new funding model substantially modifies the way the federation allocates resources for PHC to the municipalities (Ministério da Saúde, 2019^[15]). The objective of the Previne Brasil is to accelerate the expansion of the FHTs (to reach 50 000 FHT by 2022), and to strengthen quality of PHC. The new financing model includes:

- A weighted capitation calculated according to the population registered with the FHT or the PHC team, the socio-economic vulnerability and demographic profile of the registered population, and some geographical factors.
- A Pay-for-performance (P4P) bonuses. The programme is compulsory and focussed on seven priority areas, including prenatal care, women's health, child immunisation and care for chronic conditions (hypertension and diabetes mellitus) (describes in the next section). The P4P programme considers the results of the indicators achieved by the accredited teams and registered in the national register of health facilities.
- Incentives for strategic actions and priority areas, which target 16 programmes (Table 4.1). Actions cover specific characteristics according to the needs of each municipality.

Table 4.1. Incentives for strategic actions in Brazil, 2020

Areas of actions	Actions
Health promotion	Health on the Spot Program, Health at School programme, Academia da Saúde Programme
Dental care	Oral Health Team; Mobile Dental Unit, Dental Specialties Centre, Regional Dental Prosthesis Laboratory
Information and Communication Technology	Computerization of PHC facilities
Medical education	Incentive to municipalities with medical and multi-professional residency
Supporting vulnerable population	Prison PHC Team, Actions for adolescents in situations of deprivation for individual liberty, Ribeirinha FHT, Basic Fluvial Health Unit, Street Office team
Other areas	Community health agents, Microscopist

Source: Ministério da Saúde (2021^[16]), Previne Brasil, <https://aps.saude.gov.br/gestor/financiamento/incentivosacoes/>

The new Brazilian approach to adjust capitation payments in PHC is in line with OECD policies in this area (it is already implemented in Chile, Israel, Portugal, New Zealand, the United Kingdom, or the Netherlands for example). The new capitation formula intends to allocate resources based on needs. The single capitation amount will be applicable to the population registered with family and PHC teams, adjusted for socio-economic, demographic and geographical factors. Low socio-economic status will be identified if people receive benefits from the *Programa Bolsa Família* or *Benefício de Prestação Continuada* or have small pensions. The demographic factor identifies children below the age of 6 and people above the age of 64 as those with higher needs. Finally, the municipalities are clustered into five different groups based on their urbanisation using an existing typology from the Brazilian Institute of Geography and Statistics (IBGE).

When it comes to the weighting of patients, the reference patient without low socio-economic status in the age group of 6-64 years registered in the most urban type of municipality (type 1) has a base value of 1. This base value will increase by 30% if the patient has either low socio-economic status or falls outside

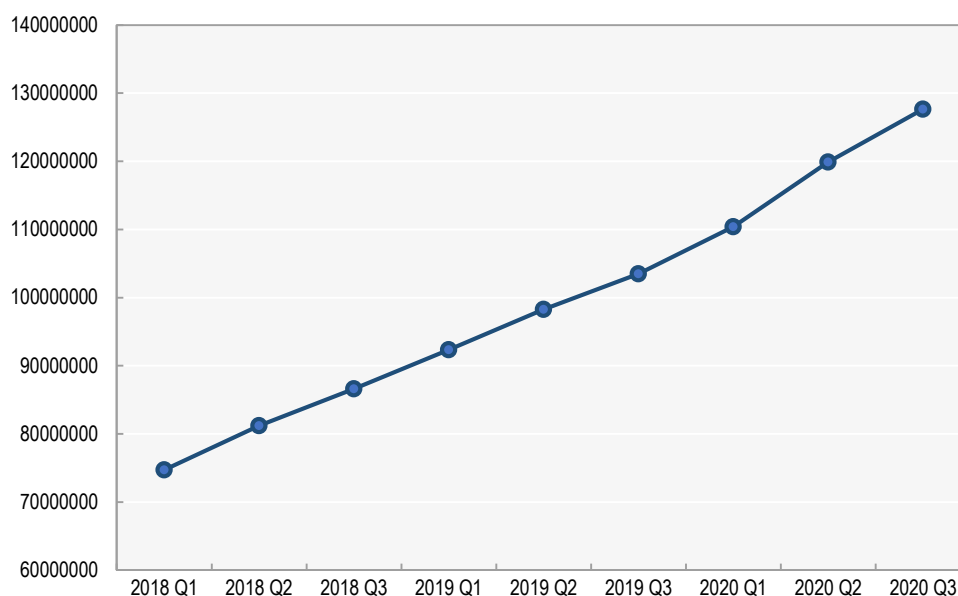
the above-mentioned age group. If a patient is registered in a municipality with medium type of urbanisation (types 2 and 3) the amount received will be multiplied by a factor of 1.45. For the two most rural types of municipalities (types 4 and 5) the amount will be multiplied by a factor of 2. Consequently, the new capitation adjustment is better taking into account differences in need (associated with age) than the previous method. Adjusting the capitated payments according to risk is an important step to make sure municipalities with higher population needs due to higher age and less favourable socio-economic situation receive the resources they need to provide care for everyone. This can limit the risk of increasing inequalities in access to PHC services.

In 2020, the value received by the municipalities from the federation ranged from BRL 51.35 for a patient in middle age (6-64 years) without low socio-economic living in an urban municipality to BRL 133.52 for a patient with either low economic status or an age outside the age bracket 6-64 living in the most rural type of municipality. These calculated amounts can be adjusted downwards if health workers in the FHT lack the required level of qualification.

There are also differences in the registration targets for each FHT based on the degree of urbanisation. In the most urban municipalities, FHTs are expected to register 4 000 people. In the two medium urbanised types of municipalities, this number stands at 2 750. In the most rural municipalities, expectations are that 2 000 people register with FHTs. Variations per FHTs are acceptable within defined limits. The new funding mechanism provides an incentive for FHT to register people in order to include the 50 million people who are not registered yet. Better registration will induce a more adequate identification of people linked to each FHT to encourage more appropriate longitudinal and co-ordinated care.

Already the number of registered people significantly increased since 2018 (Figure 4.1). The number of people registered with a FHT increased by 70% between 2018 and 2020, with around 61% of the population covered by a FHT (reaching 127 million of people registered by November 2020).

Figure 4.1. The number of people registered with a FHT has increased since 2018



Source: Ministério da Saúde (2021^[17]), Secretary of Primary Health Care, Individual Records of Family Health Teams.

In 2019, the federal government announced that municipalities will receive BRL 401 million as a financial support for increasing the number of registered people (this will amount to BRL 8 900 for each municipality)

(Ministerio da Saude, 2019^[18]). Although several researchers advocate the difficulties for some municipalities to be able to register people with FHTs and the fear of losing federal transfers (Harzheim et al., 2020^[19]), it is important to highlight that a more systematic registration system in Brazil will help build a profile of the health needs of the registered population. Registering people with FHTs is a prerequisite for better knowledge and longitudinal monitoring of people. In addition, the threat of receiving discounted capitated amounts if health workers lack the required qualification should encourage municipalities to ensure that health workers are sufficiently qualified.

In 2020, 52% of the budget costs for the provision of PHC was allocated based on weighted capitation, 24% as salaries for community health workers (a subcomponent of the strategic actions), 15% of incentive payments for strategic actions and priority areas and 9% as performance bonuses. In the context of the COVID-19, the federal government decided to maintain the amount of funds received in 2019 for each municipality, and to provide punctual financial assistance to family and PHC teams (Box 4.2 and Table 4.2).

In 2020, the budget allocated to PHC from the federal government was BRL 20.9 billion, compared to BRL 17.5 billion in 2019 (CONASEMS, 2020^[20]). In total, the federal government increased its PHC budget by around BRL 3.4 billion between 2019 and 2020.

Box 4.2. Supports provided to family and PHC teams during the COVID-19 pandemic

The Brazilian Federal Government provided punctual financial assistance to family and PHC teams through a series of block grants during the COVID-19 pandemic. These exceptional payments were given on a general basis to ensure a swift and effective PHC to manage COVID-19 cases, and to maintain routine care for non-COVID-19 patients. Municipalities with vulnerable population groups (indigenous groups, or incarcerated individuals) were specifically identified to receive additional funding. Supplementary grants were also given to mental health institutions, with amounts ranging from BRL 28 305 to BRL 400 000, depending on the size and type of psychosocial unit.

Beyond financial support, the Ministry of Health has published multiple guidelines to support family and PHC teams to managing the surge in health care demand while maintaining continuity of care for all patients.

FHTs and PHC facilities are expected to:

- Help diagnose and manage mild COVID-19 patients through testing, and providing care and appropriate exams when available;
- Follow-up people with COVID-19 to help them safely and effectively manage their care at home;
- Facilitate quick and effective referrals to hospitals for deteriorating cases.

As in other OECD countries, special COVID-19 care centres (see Table 4.2) have been established with specific health professionals and equipment to manage patients experiencing mild COVID-19 symptoms. Specific safety standards and protocols were established for high-risk patients. For example, safety protocols for delivery rooms were implemented to ensure staff is asymptomatic. In dental care, a set of ten recommendations were given to municipalities to adapt the delivery of care, including incentivising telemedicine while ensuring the continuity for dental emergencies. Informative campaigns have been developed to provide patient education on COVID-19.

To ensure care continuity, an online tool called “Consultório Virtual de Saúde da Família” has been made available to PHC facilities and FHTs. PHC workers can receive the necessary equipment, training and technical support on a 24 hours basis. PHC workers can also receive psychological support through a telephone helpline.

Table 4.2. Examples of policies introduced in Brazil to maintain care continuity for all patients during the pandemic

	Policy measures
Tele Health	Helpline to assist PHC workers to manage COVID-19 cases
Online teleconsultation platform and TeleSUS	Teleconsultation service to guarantee the continuity of PHC. Around 2 300 professionals were certified to use the system.
COVID-19 Assistance centres	Development of 3 395 centres throughout Brazil. They received special monthly federal funding, ranging from BRL 60 000 to BRL 100 000 according to municipal population size.
Community Reference Centers	Community centres responsible for screening suspected COVID-19 cases, providing in-person consultations when necessary, monitoring people at home, testing high-risk patient groups, and disseminating the TeleSUS service. The focus groups for these centres are underprivileged communities and favelas. Around 130 centres were established and BRL 8.2 million spent each month.
COVID-19 contact tracing	Effort aimed at integrating health surveillance and PHC to better identify and follow COVID-19 cases and contacts at local levels
Other use of digital health to maintain access to care for patients with chronic conditions	Three punctual projects implemented in collaboration with the Federal University of Minas Gerais aimed at developing digital platforms to guarantee continuity of care and to provide other digital instruments to reduce referrals to specialist care. The three projects combined amounted to more than BRL 9 million.

Several OECD countries have also expanded PHC to reduce pressure on health systems and minimise complications and direct death from the COVID-19 crisis (OECD, 2021^[22]):

- Use of COVID-19 community care facilities to manage patients experiencing mild COVID-19 symptoms, provide patient education on COVID-19 and work in close co-operation with other PHC centres, such as in Iceland, Slovenia and the United Kingdom;
- Expanding home-based programmes to improve access to care for all patients during the crisis and to alleviate pressure on hospitals in Canada, Spain or the United States;
- Leveraging telemedicine to support access to essential services and information. Removal of institutional barriers were required during the first wave of the COVID-19 pandemic, such as expanding insurance benefits to include telemedicine (Estonia), expanding provider payments (Canada and Australia), designing new telemedicine services (Canada and Belgium), and lifting requirements for face-to-face encounters prior to being seen virtually (France).
- Rearranging tasks and responsibilities in PHC, notably to allow community pharmacists to extend prescriptions and prescribe chronic disease medications, such as in France, Portugal or the United States.

Source: Brazilian Ministry of Health; OECD (2021^[22]), Strengthening the frontline: How primary health care helps health systems adapt during the COVID-19 pandemic, <https://dx.doi.org/10.1787/9a5ae6da-en>.

4.2.4. The Agency for the Development of PHC is being established

The Agency for the development of PHC (ADAPS) is an autonomous social agency created by the Law 13 958 of December 2019. It is a privately incorporated non-profit organisation. The Agency will promote

the execution of policies for the development of PHC at the national level, including the new *Programa Médicos pelo Brasil* (see Chapter 5).

The advisory Board will consist of 12 representatives: six from the Brazilian Ministry of Health, one from CONASS, one from CONASEMS, one from the Brazilian medical association, one from the federal board of medicine, one from the Brazilian National Federation of Physicians and one from the National Council of Health (CNS).

The ADAPS will be responsible for PHC service provision, professional training and qualification, research and extension, incorporation of health care management technology, monitoring and assessment of health care activities, and execution of the new *Programa Médicos pelo Brasil*. It is expected that the new ADAPS will provide support for actions within the municipalities to execute the PNAB. This is a welcome development to provide support to vulnerable municipalities with low capacities and greater health care needs.

4.3. Brazil has put in place a number of initiatives to support improvements in quality of care

Several initiatives have been introduced within SUS to improve the quality and the performance of PHC in Brazil. These include the development of standards of care and guidelines, of health care services portfolio, of pay-for-performance programmes and the production and publication of data underpinning PHC.

4.3.1. Few institutions are responsible for improving quality of PHC

As mentioned in Chapter 2, the Política Nacional de Atenção Básica (PNAB) describes the aspects of the governance, the roles of different actors and other aspects of policy settings and governance for PHC.

The SUS National Audit Department (DENASUS) performs audits of Municipal Health Secretariats to verify the compliance with the PNAB. Based on auditing activities, DENASUS is also responsible for monitoring compliance with the recommendations from the Ministry of Health. The Department proposes and promotes methods to assist the execution and implementation of policies. It can provide technical and methodological support to the states, the federal district, and the municipalities as part of the SUS National Auditing System. However, DENASUS is a central entity of SUS National Auditing System, and does not work as an inspectorate for health to provide independent verification that standards of care are being met.

4.3.2. Standards of care and guidelines help structure the FHT responses to most health problems

In Brazil, practice clinical guidelines are developed by governmental and non-governmental organisations at municipal, state or federal government levels. The Ministry of Health and some professional and scientific societies such as the Brazilian Medical Association, the Brazilian Society of Diabetes or the Brazilian Society of Cardiology are involved in clinical practice guidelines (CPGs).

There are also the PHC Notebooks¹ (*Cadernos de Atenção Básica em Atenção Primária*) which are clinical practice guidelines specifically designed for FHTs to improve the delivery of high quality PHC (Ministerio de Saude, 2021^[23]). They are produced by CONASS to provide guidance on for example how to better meet the health needs of people, ensuring the diagnosis of morbidity, initiating and monitoring the treatment and control of diseases or risk factors. These guidelines are easily available to FHTs.

Many of the PHC Notebooks address specific diseases or risk factors such as care for the smokers (*Cadernos de Atenção Básica* 40), obesity (*Cadernos de Atenção Básica* 38), arterial hypertension

(Cadernos de Atenção Básica 37), diabetes mellitus (Cadernos de Atenção Básica 36) and mental health (Cadernos de Atenção Básica 34).

There is one PHC notebook to address chronic diseases (Cadernos de Atenção Básica 35) published in 2014. The notebook highlights the current importance of chronic conditions, the complexity of such diseases and of interrelated risk factors. It addresses concepts common to several chronic diseases and presents guidelines for the organisation of PHC, for example on health promotion and disease prevention common to tobacco consumption and cardiovascular diseases, or common to tobacco consumption and arterial hypertension. The PHC book presents some risk stratification models, and some case studies to guide health professionals. It also includes strategies for changing habits, promoting healthy eating and practicing physical activity, approaches to building and monitoring care plans and support for self-care (Ministerio de Saude, 2014^[24]).

While such guidelines are useful, the PHC notebooks are still too narrowly defined and does not meet patients' expectations on integrated health care. There are too few clinical practice guidelines addressing integration of care for high prevalence chronic diseases (see also Chapter 3). In addition, information about their effective use is lacking. Implementation is the responsibility of the municipalities, but there is no mechanism at federal level to monitor compliance, and no systematic incentives to stimulate guideline uptake. Recent evidence show that compliance to clinical practice guidelines are unsatisfactory given inadequate follow-up observation of diabetes control and blood pressure control for people with hypertension (Leite Simão et al., 2017^[25]; da Silva Rêgo and Radovanovic, 2018^[26]). Another study shows that the degree of compliance with preventive care norms could be improved in Brazil (Guanais et al., 2019^[27]). Less than 50% of men in the target population reported a blood pressure check in the last year and a measurement of cholesterol levels in the last five years.

In addition, while high-quality clinical practice guidelines are critical for improving health care management, some studies have noticed the low quality of CPGs in Brazil. Available evidence shows that around 77% of CPGs were deemed “low quality” (de Godoi Rezende Costa Molino et al., 2016^[22]). Factors undermining the quality of CPG included a lack of multi-disciplinary team for the development group, no consideration of patients' preferences, a lack of rigor of development and conflict of interest among other (de Godoi Rezende Costa Molino et al., 2016^[22]).

4.3.3. The PHC Portfolio defines the protocols and services offered in PHC

The SAPS established the PHC Portfolio (*Carteira de Serviços da Atenção Primária à Saúde*, CaSAPS). The portfolio defines and presents the procedures and services that should be offered in PHC to reduce the heterogeneity of existing practices between municipalities, but also between PHC units in the same municipality. It helps FHTs to organise themselves in the service routine, and seek appropriate competences and skills to offer the actions and services define in the protocol through several actions such as continuous personal development strategies, monitoring, and evaluation (da Cunha et al., 2020^[28]).

The CASAPS is one of the most important tools to support municipalities and FHTs to provide high quality care. The portfolio has already been used in Rio de Janeiro, Curitiba and Florianópolis among other places (da Cunha et al., 2020^[28]). Rio de Janeiro developed his own portfolio of PHC services to address the issue of having a high concentration of patients at the secondary and tertiary levels of care seeking low complexity procedures, such as ear washing or nail extraction (Salazar, Campos and Luiza, 2017^[29]). This is a source of wasted resources for the health care system, overburdening emergencies and hospitalisation services, and reducing the quality of care for PHC conditions (Salazar, Campos and Luiza, 2017^[29]).

The portfolio contains 210 actions and services organised in five groups: “health promotion for adults and elderly” (44 items), “Health care for Children and adolescent's (27 items), “PHC procedures” (17 items), and “Oral health care” (21 items). Each item is associated with recommendations, materials for consultation, and lists of inputs for implementation.

The elaboration of the portfolio is based on a review and evaluation of services implemented in Brazilian cities, and on consultations with several bodies including the Ministry of Health, CONASS, CONASEMS, the Professional Associations of the Brazilian Society of Family and Community Medicine (SBMFC) or the Brazilian Association of Family and Community Nursing (ABEFACO) (da Cunha et al., 2020^[28]).

In the municipality of Rio de Janeiro, the use of the Services Portfolio was associated with a better performance compared to other municipalities not using similar portfolio (da Cunha et al., 2020^[28]).

While the development of such PHC portfolios contributes to strengthen the attributes of a strong and comprehensive PHC sector, an important challenge relates to the need to have appropriate training for PHC providers to be able to deliver high quality actions and services. FHTs need to have the right and sufficient equipment and capacity to execute the PHC procedures and services.

4.3.4. National Programme for Improving PHC Access and Quality had a positive impact on quality

The federal government introduced a pay-for-performance programme (P4P) for FHTs in 2011 called the National Programme for Improving Access and Quality of PHC (PMAQ-BA in Portuguese). The programme defined standards of care in accordance to the PNAB to evaluate the performance of FHTs towards a wide range of structure, process and outcomes indicators.

Three cycles have taken place (in 2011/2013, 2013/2015 and 2015/2019). Although the programme was voluntary, almost 100% of all FHTs, oral health teams, NASFs and dental specialty centres were registered in the third round of the PMAQ (compared to 71% in round 1 and 91% in round 2).

The indicators selected were infrastructure, management for PHC development, worker appreciation, access and quality of health care and user satisfaction. In the 3rd round, more than 650 indicators were included around structural quality of care (for example availability of drugs and equipment), process of care (for example antenatal care and treatment completion rates), quality of care (for example patient satisfaction), utilisation of health care (patient volume), and management processes (proportion of appointments that are scheduled) (Kovacs et al., 2021^[30]).

Certification processes consisted of self- and external assessment of the teams, involving facility assessments, examination of health indicators and interviews with health care professionals, municipal managers and service users. External evaluations were conducted by independent academic institutions by collecting indicators, and by conducting interviews (Macinko, Harris and Rocha, 2017^[31]). A mean score was then calculated, and grouped according to the overall human development index for their municipalities to create several strata of similar socio-economic environments.

Each team received the bonus according to its ranking compared to a mean score for their stratum:

- Those at or below the mean score for their stratum received 20% of the bonus;
- Those above the mean received 60% of the bonus;
- Those significantly above the mean receive 100% of the bonus.

From 2011 to August 2018, the Ministry of Health has invested BRL 9.85 billion (USD 2.52 billion). The bonus ranges from BRL 1 700 (USD 434.55) to BRL 11 000 (USD 2 811.79) per team and month based on their performance (de Medeiros et al., 2020^[32]).

One important consideration was the fact that each's team score was made publicly available on the Ministry of Health's website. It enhanced transparency and allowed citizens and health managers to see the overall performance of each team and the associated assessments. Each team was compared to municipal, states, and national averages, which helped nurture a quality improvement culture in PHC services.

Available data shows that PMAQ has led to an increase in federal investment for infrastructure, and a large increase in performance incentives (Macinko, Harris and Rocha, 2017^[31]). In addition, the PMAQ has been associated with an increase in the provision of care to pregnant women and children under 2 years, with more significant improvement among the worst performing PHC providers (de Medeiros et al., 2020^[32]). The positive impact of the PMAQ is also confirmed by a recent evaluation showing positive effects on the quality and access of PHC (Soares and Ramos, 2020^[33]). On average, municipalities that adhered to the PMAQ had rates of hospitalisation for chronic conditions 9% inferior to those that did not adhere *ceteris paribus*. The effect of the programme was more pronounced in the Northeast regions of Brazil (rates of hospitalisation for chronic conditions 14% inferior for municipalities that adhered to the programme), and less pronounced for Southeast region (rates of hospitalisation for chronic conditions 5% lower for municipalities that adhered to the programme). Kovacs and al (2021) also showed that existing income inequalities in the delivery of quality PHC were eliminating during the three period of PMAQ implementation, due mainly to the design feature that adjusted financial payments for socio-economic inequalities (poorer municipalities received higher rewards than richer municipalities) (Kovacs et al., 2021^[30]).

However, several failures of the programme were noted, including the very large group of indicators used (with almost 900 questions) and a too large focus on structure indicators, with little emphasis on outcomes indicators. According to the Ministry of health, the high number of indicators resulted in monitoring problems and lack of adherence by municipal managers. It is also reported that external evaluation as part of PMAQ was highly criticised, and thus not carried-out by health professionals. Lastly, the payment made by the Ministry of Health were directed towards the municipality and not the FHT. Municipalities were free to provide bonus payments to the team but little is known on the actual practice (Macinko, Harris and Rocha, 2017^[31]).

Despite these failures, the PMAQ programme had resulted in improvement of quality of PHC, notably because it was possible during the eight years of implementation to monitor improvements in infrastructure, medical supplies, medications and health outcomes.

4.3.5. The *Previne Brasil* established a new pay-for-performance scheme

With the reform of the PHC financing model, the federal government introduced a new P4P programme in 2020 as part of the *Previne Brasil* programme.

The new P4P programme is compulsory and focussed on fewer priority areas than the previous PMAQ programme. The initial number of indicators used in this programme was set at seven in 2020 and will rise to 21 by 2022.

In 2020, priority is given to prenatal care, women's health, child immunisation and two chronic conditions (hypertension and diabetes mellitus) (Ministério da Saúde, 2019^[15]). The selected indicators are:

- Share of pregnant women with six or more prenatal consultations in the first 20 weeks of pregnancy
- Share of pregnant women with tests for syphilis and HIV carried out
- Share of pregnant women with dental examination
- Pap smear test carried out to detect cervical cancer
- Coverage of inactive polio and pentavalent vaccines
- Share of patients diagnosed with hypertension which have their blood pressure measured every six months
- Share of patients with diagnosed diabetes with controlled haemoglobin level.

For each of the seven indicators, weights of 1 or 2 are assigned to calculate the Final Synthetic Indicator. The weights reflect the clinical and epidemiological relevance of the indicator, as well as the level of difficulty in achieving the targets. This translates the effort of managers and teams to carry out actions,

programs and strategies. Table 4.3 presents the overall targets, the targets for 2020 and the weights defined for each indicator.

Table 4.3. Targets and weights assigned for each performance indicator, 2020

	Domains and indicators	Overall targets	Targets for 2020	Weights
Women health	Share of pregnant women with 6 or more prenatal consultations in the first 20 weeks of pregnancy	>=80%	60%	1
	Share of pregnant women with tests for syphilis and HIV carried out	>=95%	60%	1
	Share of pregnant women with dental examination	>=90%	60%	2
	Pap smear test carried out to detect cervical cancer	>=80%	40%	1
Care for children	Coverage of inactive polio and pentavalent vaccines	>=95%	95%	2
Care for chronic diseases	Share of patients diagnosed with hypertension which have their blood pressure measured every 6 months	>=90%	50%	2
	Share of patients with diagnosed diabetes with controlled haemoglobin level.	>=90%	50%	1

Source: Ministério da Saúde (2019^[34]), NOTA TÉCNICA N° 5/2020-DESF/SAPS/MS.

The performance score for each indicator will range from zero to ten, considering the targets for 2020 and the weighting scale. If the result for a given indicator is 30%, while the target is 60% for 2020, the final score for this indicator will be 5.0 (Ministério da Saúde, 2019^[34]).

Further indicators to be included in the years 2022 will be discussed after evaluation and will cover the following areas: care for new-borns; care for babies (under 1-year-old); HIV, Tuberculosis, dental care, hepatitis, mental health, breast cancer, multi-professional action, and global measures including patient experience. Patient experience measurement will rely on population-based surveys conducted by the Brazilian Institute of Geography and Statistics (IBGE), which use PHC Assessment Tool (PCATool), the Patient-Doctor Relationship Questionnaire (PDRQ-9) and the Net Promoter Score (NPS) (Ministério da Saúde, 2019^[15]).

While all these standards of care are based on scientific evidence, and selected on epidemiological and clinical relevance, there are all process indicators to assess the quality of care delivered.

The P4P programme and related transferred resources correspond to 9% (BRL 1.8 billion) of the budget linked to the *Previne Brasil* Programme. Performance indicators are collected every four months, and bonuses are paid monthly to municipalities based on the performance of their respective teams.

The amount of the performance payment vary according to the classification of team (Ministério da Saúde, 2020^[35]). The following performance payment, referring to 100% of the Final Synthetic Indicator, are equivalent to:

- BRL 3 225.00 for the FHT;
- BRL 2 418.75 for the PHC Team Modality II 30h; and
- BRL 1 612.50 for the PHC Team Modality I 20h.

Results are publicly available on the Health Information System for PHC (SIAB) in the form of reports and dashboards, and on the e-Gestor AB web platform so that all municipalities and states can follow and adopt quality improvement strategies (e-Gestor AB, 2021^[36]).

4.3.6. The Brazilian Government has invested a lot of efforts in the production and publication of data underpinning PHC

Several initiatives have been implemented to monitor the resources and activities in the PHC sector in Brazil. Within the Ministry of Health, the Department of Health Information (DATASUS) holds several databases including the national register of health facilities (CNES), the National Program of Immunization (SI-PNI), the System for prenatal care (SIPRENATAL) and the PHC Information System (SISAB, previously called SIAB) to mention a few.

The SISAB was established by Ordinance GM/MS N° 1 412 of July 2013 for the financing purpose and for implementing the PNAB. Community health workers collect on a monthly basis health and social data, monitoring of risks groups, priority health problems, child's follow-up, and on certain diseases and risk factors for health including alcoholism, disability, diabetes, epilepsy, hypertension, malaria or tuberculosis (Lopes, Monteiro and Santos, 2020^[37]). The SISAB also integrates the information related to social programmes including Health at School Programme (*Saúde na Escola*) or the Health Academy Programme (*Academia da Saúde*).

While the SISAB gathers an impressive amount of indicators underpinning PHC, most indicators are inputs and process indicators including for example the number of FHTs, the number of people registered with a FHT, the number of community health agents, or the number of pregnant women receiving first prenatal care services (SISAB, 2021^[38]).

It is fair to note that Brazil has developed some actionable indicators to support quality improvement in PHC, notably through the PMAQ strategy and the newly *Previne Brasil* programme. In 2020, quality indicators designed as part of the P4P programme are process indicators focussing on maternal health, cervical screening programme, immunisation coverage, and management of hypertension and diabetes. Still, they are good measures of the quality of care delivered by each FHT.

Input and process indicators are made publicly available, broken down by Federative Unit and Region, by urban or rural areas and by age group. The data can be presented in various formats such as tables, graphs, maps and technical documents in order to facilitate the interpretation and analysis of the data. This is an important quality initiative in Brazil as open publication promotes transparency within the PHC sector and expand the possibilities for improving care quality. However, the data is not published at more local, granular level. To date, patients are not allowed to monitor activity and quality of PHC at FHT level. Municipal health managers and FHTs have however access to their performance results through the e-Gestor AB portal (e-Gestor AB, 2021^[36]).

At the same time, national reports based on the DATASUS databases are publicly available to assess and monitor population health and the health care system. The *Indicadores de Saúde no Brasil: conceitos e aplicações* for example reports data on health expectancy, diseases-specific mortality rates, diseases-specific incidence rates, prevalence of risk factors for health and some health care resources and activities.² The 2019 National Health Survey, which was jointly published by the IBGE and the SAPS, focuses in particular on access and use of PHC services.³ The Survey provides data on access in PHC, it provides some prevalence rates for risk factors for health and some quality indicators around maternal health or cancer screening programmes. Importantly, the PHC module of the survey consisted of 26 items distributed in 10 components based on the PHC Assessment Tool (PCAT) developed by Starfield and Shi (Pinto and Silva, 2021^[39]). The use of PCAT allowed to assess the attributes and the characteristics of a strong PHC: first contact access, longitudinal, co-ordination, comprehensiveness, family orientation, and community orientation (the next section of this chapter presents the results of the 2019 National Health Survey). Incorporating a standardised and internationally validated instruments for the evaluation of PHC services is a major step in Brazil to improve the provision of high quality patient centred care. It demonstrates the high political will to strengthen PHC services in Brazil.

It is also important to note that the e-SUS PHC System was developed to be used in multiple computerisation and care settings, ranging from PHC units without computers to those with computers and the internet (Sousa et al., 2019^[40]). The e-SUS PHC System is made of two softwares: simplified data collection and individual health records. The simplified data collection has been specifically developed for non-computerised environments. It is a software for entering a set of files that include the work process of FHTs. Today, only around 40% of FHTs do have adopted the individual health records (see also Chapter 6).

While there is lot of data underpinning PHC in Brazil, some studies have noted the poor reliability and quality of the collected information for planning and public decision making (Lopes, Monteiro and Santos, 2020^[37]). As municipalities can be penalised with cancellation of their funding resources, the work process is rather motivated by production targets. It is reported that community health workers, responsible for the data collection, have not properly completed the forms because they do not have the knowledge and training to perform the task. So far, these data have not been used to estimate diseases prevalence, characterise the epidemiological profile of the population, assess health status and even more important be linked with other information system. (Lopes, Monteiro and Santos, 2020^[37]). This is a challenge also reported by the OECD Brazil Health System review (OECD, 2021^[41]).

4.4. Outcomes associated with PHC

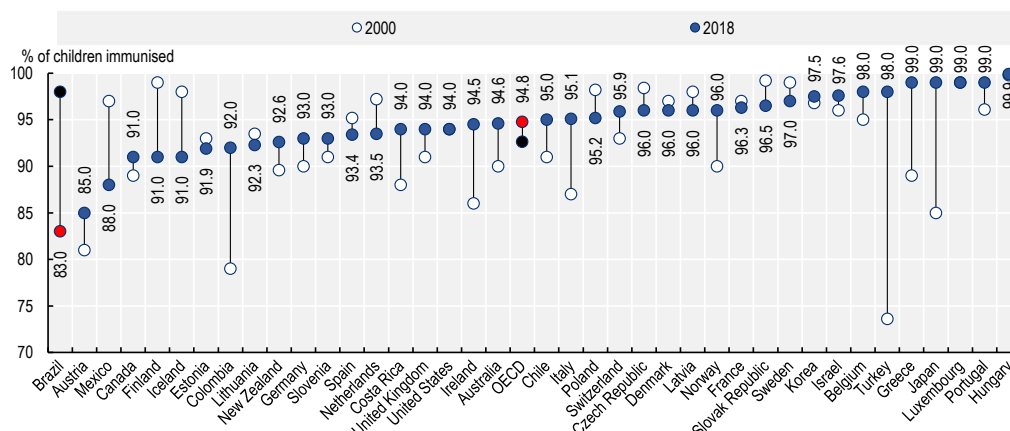
Despite progress, several indicators pertaining to quality of PHC place Brazil among poor performers. PHC in Brazil has traditionally been characterised by providing few low complexity procedures and surgeries. In addition, international figures demonstrate that Brazil lags behind other countries regarding to preventive care and management of chronic conditions. Large regional differences in health care quality also suggest that some remote and vulnerable areas require more attention, more support than others to improve quality of PHC.

4.4.1. Brazil lags behind other countries with regards to preventive care

There is strong evidence that vaccines provide safe and effective protection against diseases such as diphtheria, tetanus, pertussis, measles, and hepatitis B. The proportion of children protected from these diseases can be considered as an indicator of quality in PHC. According to the WHO Global health Observatory, 84% of Brazilian children aged around 1 were vaccinated against measles, below the average in OECD countries (95%). Additionally, 83% of Brazilian children aged around 1 were vaccinated against diphtheria, tetanus, and pertussis, compared with OECD average of 95% (Figure 4.2). In a similar vein, Brazil achieved 83% coverage for vaccination against hepatitis B for children aged around 1, compared with the OECD average of 91%.

There are also significant inequalities in the rates of vaccination against yellow fever in Brazilian states where vaccination has been recommended, ranging, in 2018, from 21.8% in Sergipe to 100% in the Federal District and Roraima (the national average for all states where vaccination is recommended is 64.1%).

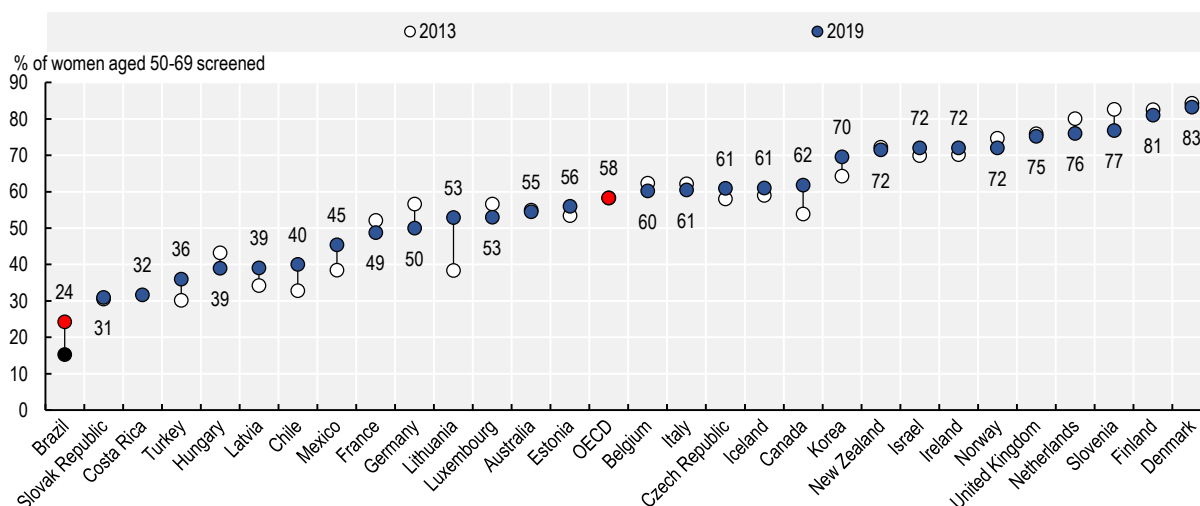
Figure 4.2. Vaccination rates for diphtheria, tetanus toxoid and pertussis (DTP3), children aged around 1, 2000 and 2018 (or latest year available)



Source: OECD (2021^[42]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>; Brazilian Ministry of Health.

International figures also show that breast and cervical cancer screening needs to improve. Although screening for breast cancer in Brazil has improved in recent years, breast cancer screening coverage in Brazil (at 24%) is very low when compared to OECD countries. Brazil stands below all OECD countries and well below the 58% of average coverage (Figure 4.3). In a similar vein, cervical cancer screening in Brazil stands at 37.1% in 2019, well below the OECD average coverage of 58% (see Chapter 3 on Screening in PHC).

Figure 4.3. Breast cancer screening coverage in Brazil and OECD countries, 2013 and 2019 (or latest year available)



Note: Programme data is used for all countries. Brazil represent years 2014 and 2019.

Source: OECD (2021^[42]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>; Brazilian Ministry of Health.

Inappropriate preventive care has also been highlighted by previous studies. The use of patient-reported experience measures for the prevention, detection and management of chronic diseases has shown that only around one people over five reported that their PHC physicians sent a reminder for check-up among

SUS users (Guanais et al., 2019^[27]). In addition only 22% of SUS users indicated that they had received some recommendation on healthy lifestyles from their family doctor, and less than 40% of public users had a preventive visit in the past two years.

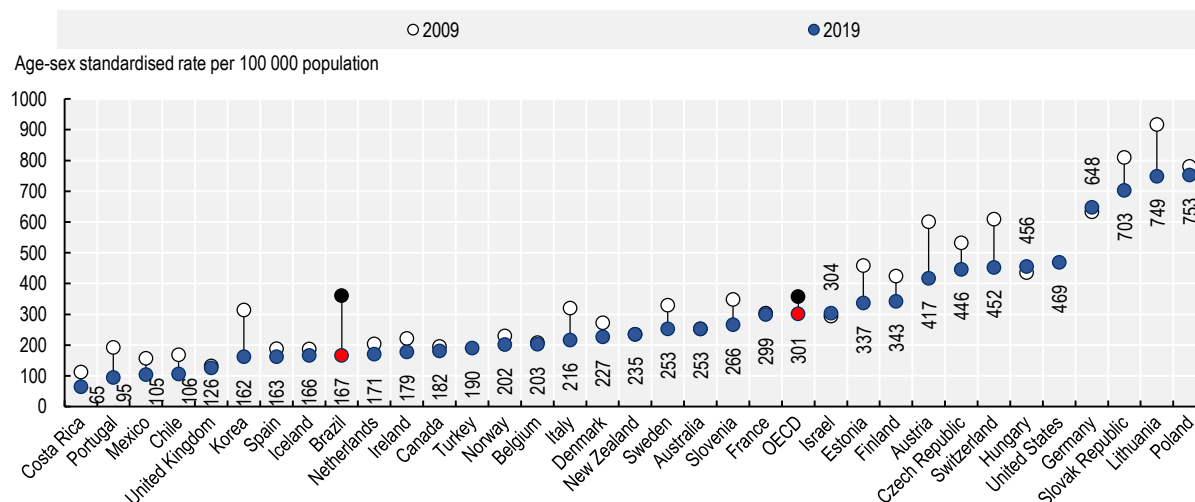
More preventive care is warranted in Brazil at a time when the burden of disease is shifting towards chronic diseases.

4.4.2. Indicators around the management of chronic conditions give some cause for concern

Hospitalisations for chronic conditions, also known as avoidable hospitalisations, are used as an indirect indicator of the overall effectiveness of PHC. There is growing evidence that proactive management of some chronic conditions may prevent or reduce the need for acute hospital admission (OECD, 2020^[43]). Ambulatory care sensitive conditions, such as asthma, chronic obstructive pulmonary disease (COPD), congestive heart failure, and diabetes, are conditions for which accessible and effective PHC can generally reduce the risk of complications and prevent the need for hospitalisation. The evidence base for effective treatment for these conditions is well established and much of it can be delivered at a PHC level. Therefore, a high performing PHC system should be able to avoid to a large extent any acute deterioration of the health status of people living with chronic conditions and prevent their admission to hospital.

In Brazil, hospitalisations for chronic conditions are lower than in OECD countries. The rate of diabetes hospital admissions in 2019 was 92 admissions per 100 000 people, below the OECD average of 131 (see Chapter 3). In addition, hospital admission rates for congestive heart failure (CHF) and hypertension are lower in Brazil than the OECD average, but still higher than other LAC countries including Costa Rica, Mexico, or Chile (Figure 4.4).

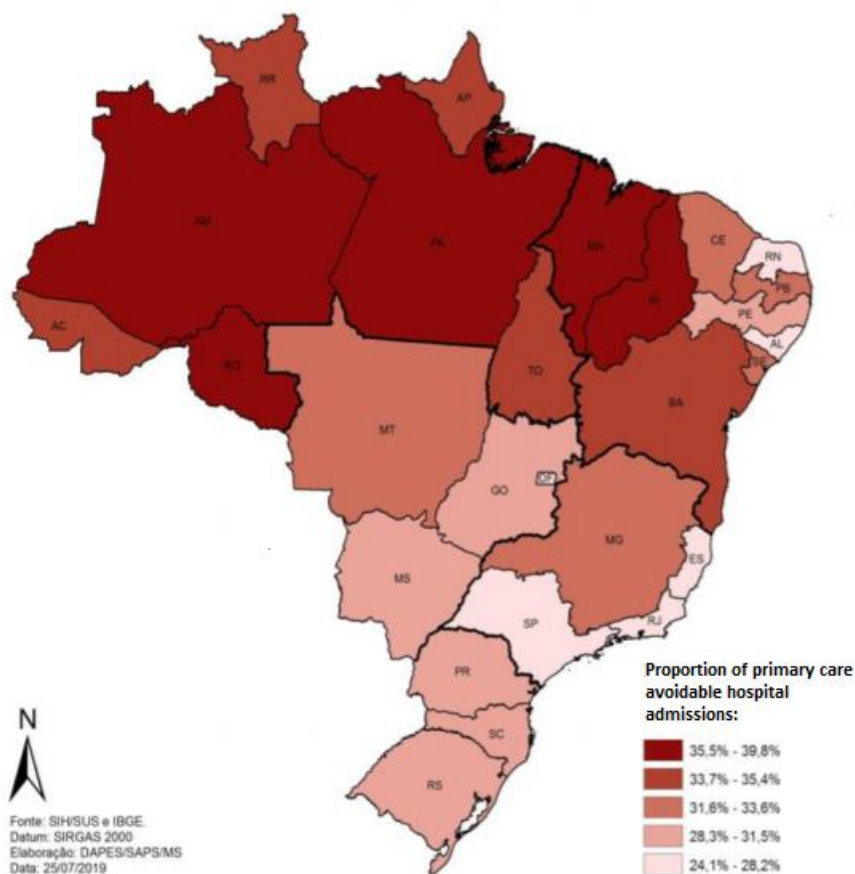
Figure 4.4. Congestive heart failure (CHF) and hypertension hospital admissions in adults, 2009 and 2019 (or latest year available)



Source: OECD (2021^[42]), Health Statistics Database, <https://www.oecd.org/health/health-data.htm>; Brazilian Ministry of Health.

In Brazil, lower rates of avoidable hospital admissions can be the result of a lower prevalence of chronic conditions than across OECD countries. At the same time, whilst the performance is good on average, there are significant differences across regions. There is a North-South gradient where North and Northeast regions show higher hospitalisation rates for chronic conditions (Figure 4.5).

Figure 4.5. Share of hospital admissions due to conditions that are treatable in PHC, population aged 60 or more by region, 2019

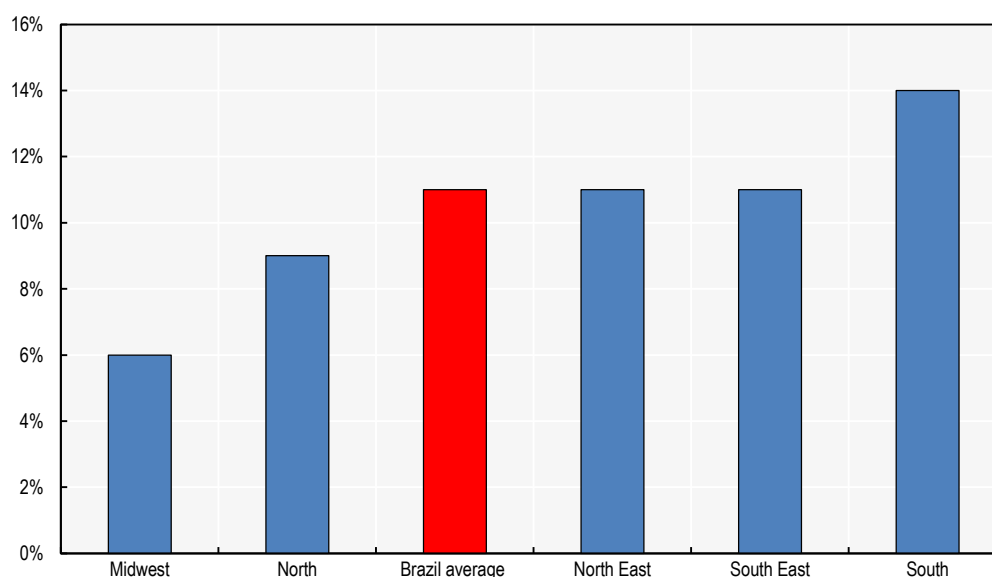


Source: Ministério da Saúde (2020^[44]), Plano Nacional de Saúde 2020-23.

The 2019 Health Interview Survey also corroborates the North-South gradient. Accordingly, diabetic patients have a higher likelihood of complications (as measured in hospitalisations) in the Northeast than in the Southeast (IGBE, 2020^[45]).

High hospitalisation rates for chronic conditions show that recommended care is not always provided for people with these conditions. This is confirmed by performance indicators reported in the SIAB system. Two indicators relate to whether people with diabetes and hypertension are receiving recommended care or effective management for their conditions. In 2020, only 4% of hypertensive patients had their blood pressure measured every semester, and only 11% of diabetics patients had controlled haemoglobin level (SISAB, 2021^[38]). Although such very low performance scores certainly reflect difficulties in recording and monitoring PHC activities by municipalities, it gives some cause for concern. Again, the South and Southeast regions report higher performance score than the North, Northeast and Central-West region regarding to the management of diabetes patients (Figure 4.6).

Figure 4.6. Share of diabetes patients with controlled haemoglobin level, 2020 (Q3)



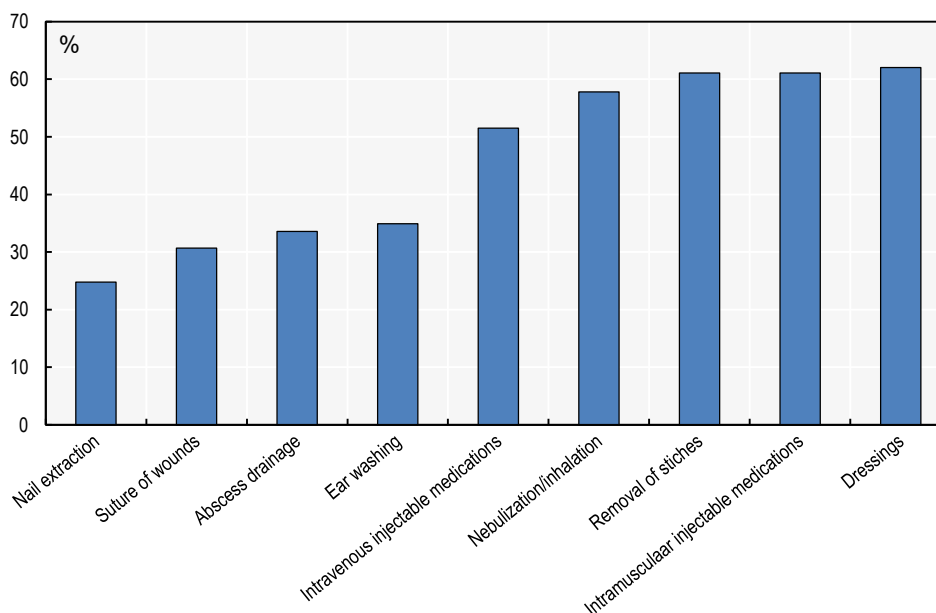
Source: Ministério da Saúde (2021^[46]), Secretary of Primary Health Care, Performance indicators by Regions, <https://sisab.saude.gov.br/paginas/acesoRestrito/relatorio/federal/indicadores/indicadorPainel.xhtml>

Such inequalities in favour of wealthier regions can be explained by a greater supply of PHC services, health workers, equipment and compliance with clinical practice guidelines or standard treatment guidelines. The use of therapeutic guidelines protocol for hypertension, diabetes, tuberculosis, leprosy and mental health service is found to be consistently lower in small geographical areas (Salazar, Campos and Luiza, 2017^[47]). Kovacs et al. (2021) also confirmed that FHTs located in poorer areas performed significantly worse than those in richer areas. A PMAQ performance score across 20 income groups has shown that a higher monthly household income of BRL 1 000 was associated with a 1.59 percentage point higher PMAQ score (Macinko, Harris and Rocha, 2017^[31]).

4.4.3. Simple procedures and small surgeries are not always provided by family health teams

Simple procedures and surgeries should be widely provided in PHC settings. This is both a matter of striving for better safety and care quality, and an economic necessity. However, available data from the PMAQ programme show that only 62.8% of FHTs performed low complexity procedures or small surgeries in 2012. Worst, only one-third of FHTs performed suture of wounds and abscess drainage. The highest national share for low complexity procedures is around 60% for removal of stitches, dressings, or intramuscular injectable medications (Figure 4.7). Still, this is too few if Brazil wants to increase the responsibility of PHC.

Figure 4.7. PHC has historically offered a low supply of procedures



Source: Adapted from Salazar, Campos and Luiza (2017^[47]), The Rio de Janeiro Municipality's Services Portfolio and Health Actions in Primary Care in Brazil, <https://doi.org/10.1590/1413-81232017223.33442016>.

The low supply of simple procedures and surgeries is a source of care fragmentation and inappropriate referrals to more specialised care. High turnover of medical doctor, inadequate training to practise as PHC physicians, and the lack of equipment or resources are leading factors contributing to the difficulty of performing such procedures for FHTs (Salazar, Campos and Luiza, 2017^[47]).

4.4.4. There is scope to improve patient experiences with public PHC services

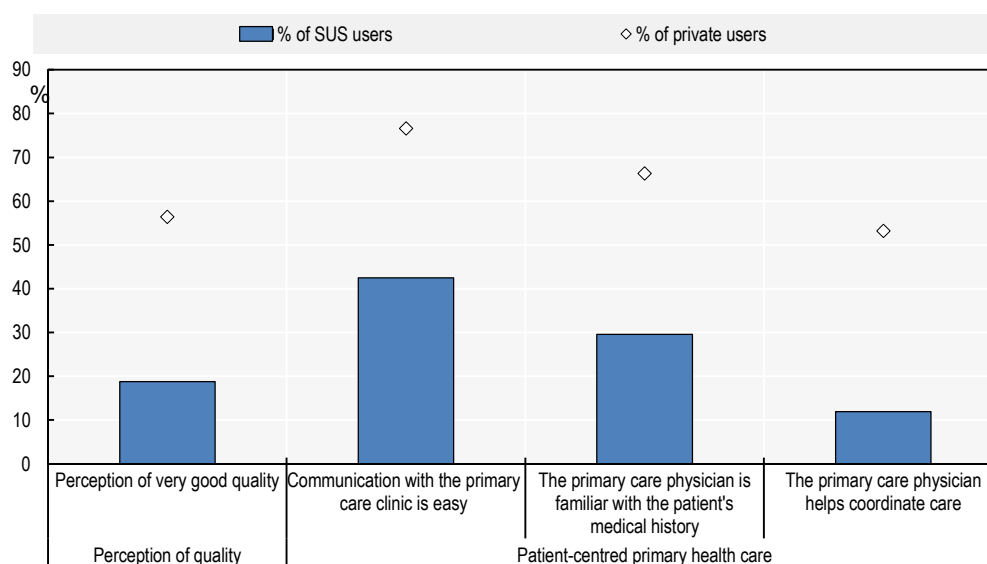
Recent data from the 2019 National Health survey show that patient experience with PHC in Brazil needs to further improve. PHC obtains an overall score of 5.9 on a scale of 0 to 10, which is below the benchmark value of 6.6 recommended by the PCAT methodology (IBGE, 2019^[48]). Some differences appear when considering registered households and non-registered households (Table 4.4). The general score was 6.0 for registered people versus 5.5 for non-registered people. The score was also more favourable among household that received at least one visit from community health workers or other members of the FHT, and for people having chronic conditions. People living in the South region also reported a higher score (6.3) than people living in the North (5.5).

Less recent data looked at people-centeredness of PHC in Brazil. Good provider-patient communication and patient involvement in care and treatment decisions are key dimensions of patient centeredness, alongside the dimension of co-production which entails patients self-management and care co-ordination (OECD, Forthcoming^[49]). Results from the PMAQ 2017 cycle showed that around 90% of patients reported that PHC doctors provided easy-to-understand explanation, and involved patients in decisions about care and treatment. An adaptation of the Commonwealth Funds International Health Policy Survey to the LAC context has shown a large variation in patient's experience between public and private users. SUS users were consistently less likely to report a positive experience with PHC compared to private users (Figure 4.8).

Table 4.4. Overall mean score of PHC (value from 0 to 10) depending on individual characteristics

Individual characteristics	Yes	No
Arterial hypertension	6.2	5.7
Diabetes	6.3	5.8
Heart disease	6.4	5.8
Depression	6.1	5.8
Is the household registered at the family health facility	6.0	5.5
Visit from community health workers	6.1	5.7
Overall score per Regions		
North	5.5	
Northeast	5.8	
Southeast	5.8	
South	6.3	
Midwest	5.8	

Source: IBGE (2019^[48]), Pesquisa Nacional de Saude in Portuguese: Atenção primária à saúde e informações antropométricas, <https://biblioteca.ibge.gov.br/visualizacao/livros/liv101758.pdf>

Figure 4.8. SUS users consistently report lower patient experiences with PHC than private users, 2013

Note: Prevalence are adjusted for sex, age, level of education and self-reported health status.

Source: Authors, adapted from Guanais et al. (2019^[27]), From the patient's perspective: experiences with primary health care in Latin America and the Caribbean.

4.5. Securing a greater quality dividend from PHC in Brazil

4.5.1. Strengthening the gatekeeping system to better recognise the central role of FHTs

While the overarching objective of the Family Health Strategy was to strengthen access to care, 20 years after its implementation only about 65% of the Brazilian population are currently registered with a FHT. The number of registered people is increasing in recent years following the *Previne Brasil* Programme

(Figure 4.1). Still, this is too low to address the new Brazilian's health care needs characterised by rising burden of chronic conditions and risk factors for health.

In addition, many Brazilians have direct access to outpatient specialties and hospitals for minor conditions that could be treated more effectively in primary care settings. The latest Health Information Survey for example found that around 50% of the diabetic population identified basic health units as their last contact to the health system, 11% used public speciality units in hospitals, 6% public urgent care units and 30% private practices or clinics (IBGE, 2020^[50]). The later example confirms that a significant proportion of the population uses public speciality units or emergency departments for provision of chronic services, while these types of conditions are best treated in PHC settings. These give strong arguments to strengthen the gatekeeping system in Brazil with a systematic registration system with FHTs, which better control and orient the patient's into specialist care.

Gatekeeping system with a named PHC doctor or a FHT who provide overall care co-ordination can drive better care for patients, particularly for those having chronic diseases. It refers to the arrangement where a patient is only seen by a hospital specialist if authorised by the PHC physician who is responsible for overseeing and co-ordinating the health needs and providing care. In a recent systematic review of the literature, Sripa, Hayhoe, Garg et al. confirmed that gatekeeping was associated with better quality of care and appropriate referral for further hospital visits and investigation (Sripa et al., 2019^[51]). Overall gatekeeping resulted in fewer hospitalisations and use of specialist care, but also led to lower patients satisfaction because it limits patient choice (Sripa et al., 2019^[51]). In addition, a systematic registration system allow to build a profile of the health needs of the registered population, which improve care continuity through better patients' relationship to PHC providers (Starfield, Shi and Macinko, 2005^[52]; Kringos et al., 2010^[53]). Having a PHC physician as a regular source of care consistently show benefits for a variety of health and health related outcomes (Starfield, Shi and Macinko, 2005^[52]). It allows for the provision of continuous and comprehensive care, focused on prevention and management of long-term conditions.

Box 4.3. Introducing a compulsory registration system in Norway

While before 2000, Norwegian citizens were able to consult GPs without restrictions, the Regular General Practitioner reform introduced a compulsory registration system. Pilot of the reform was undertaken in four municipalities in 1993 prior to national implementation.

With the reform, the general practitioner became responsible for providing and co-ordinating prevention, investigation and treatment of health care needs. He is also responsible for decisions on the need for referral to secondary care, and link with social security and social services when appropriate. The reform also specified that GPs should engage in public health activities, emergency care, out-of-hours care, and supervise students and doctors in training.

National implementation was a success, with close to 100% of Norwegians now registered with a GP. The scheme has been regarded as one of the most successful public services in Norway, with high satisfaction among both patients and GPs (Svedahl et al., 2019^[54])

Source: OECD (2014^[55]), *OECD Reviews of Health Care Quality: Norway 2014: Raising Standards*, <https://dx.doi.org/10.1787/9789264208469-en>.

Some OECD countries have introduced a compulsory registration system to encourage the right care at the right place (Box 4.3). In Norway, the Regular General Practitioner reform introduced in 2000 required all citizens to register with a named general practitioner (GP) of their choice. General practitioners became the usual source of care for people, and are now responsible for co-ordinating individual's prevention,

investigation, and treatment of health care needs. This approach is also taken in Chile, Portugal, and Italy: in these countries, the population are required to register with a PHC physician who control access to secondary health care (OECD, 2020^[43]). Strengthening the gatekeeping system by requiring all individuals to register with a FHTs is a reform that Brazil wishes to consider in the future to strengthen the role and accelerate the expansion of the FHTs. This would help build accurate patient registers, and better recognise the central co-ordinating role of FHTs.

4.5.2. Expanding the range of health care services provided by FHTs towards disease prevention, treatment and management of long term conditions

In recognition of the challenges set out in Section 4.4, there is an increasing need to support and encourage FHTs to deliver a modern PHC service. A key function of a modern PHC is the provision of continuous and comprehensive care, focused on health promotion, prevention, treatment, follow-up of diagnoses and management of chronic and long-term conditions.

Previous studies have shown that PHC, when well equipped, can deal with a large range of health problems, including health promotion, preventive care, treatment and follow-up of diagnoses and simple technical procedures (Starfield, Shi and Macinko, 2005^[52]; Kringos et al., 2010^[53]). A comprehensive scope of PHC services is consistently associated with lower hospitalisation for chronic conditions, improved health outcomes and reduce health inequalities (OECD, 2020^[43]; Starfield, Shi and Macinko, 2005^[52]; Kringos et al., 2010^[53]).

Underpinning greater comprehensiveness of PHC services entails delivering actions defined in the PHC Portfolio, and ensuring that all municipalities are able to properly deliver these actions and health care services. In Rio de Janeiro, the Health Services Portfolio has been found to be an important guiding tool to standardise and improve the range of services offered by FHTs (Salazar, Campos and Luiza, 2017^[47]). For all defined procedures, Rio de Janeiro reports better performance than national average or other urban centres. An important policy lever has been the use of management agreements to foster compliance with the Health Services Portfolio. One indicator called “rate of implementation of items of the services portfolio” is monitored quarterly as part of the agreement, which enables to highlight problems in the practice and take corrective actions. Such a good practice could be consolidated nationwide. Another option for consideration, would be to monitor compliance rates with the CaSAPs and link it to financial incentives attributed in the Strategic Actions of the *Previne Brasil* Programme.

In a similar vein, although a number of clinical guidelines are produced in Brazil, there are no requirements and few incentives for FHTs to comply with the standard level of care. There is a risk, then, that these guidelines are not adequately adopted by FHTs. Experiences from OECD countries can be useful for Brazil in its efforts to improve compliance to clinical guidelines. Some OECD countries provide grants to municipalities to encourage compliance with clinical practice guidelines. In Sweden, the central government provides grants to regional governments to encourage guideline implementation (OECD, 2013^[56]). New guidelines on dementia, for example, were accompanied by grants to be disbursed to local government, which were then free to use the additional funds as they saw fit. This approach maintains local responsibility for effective implementation, whilst drawing in national resources and support (OECD, 2013^[56]). Arguably, the Ministry of Health could use the *Previne Brasil* Programme to monitor and encourage more systematic compliance with clinical guidelines.

At the same time, there is a need to support continuing medical education (CME). As earlier mentioned, a number of bodies such as the CONASS, CONASEM, the Brazilian Medical Association or the Brazilian Society of Family Medicine organise several training, conferences and on-line learning. However, a more formal CME framework which makes clear the expectation upon each FHT members, and supports them to meet it will be required to provide comprehensiveness of PHC services, and in line with the best available evidence.

4.5.3. Quality monitoring and benchmarking within PHC needs to improve

A fundamental element within a strengthened PHC sector is a data infrastructure capable of monitoring and benchmarking PHC activities and outcomes in a consistent way at national, municipal and provider level (OECD, 2020^[43]).

Brazil collects several indicators on access to PHC, risk factors and quality for maternal health, child immunisation, breast cancer screening and management of hypertension and diabetes a part of the *Previne Brasil* Programme. While it is an important initiative toward quality monitoring, five quality indicators are process indicators, and only 2 indicators are intermediate outcomes measures. Better measurement of patient outcomes and experiences with PHC needs to be a priority in Brazil.

It will be important for Brazil to collect a richer set of quality indicators on a wider array of preventive activities and management of chronic conditions such as alcohol consumption, obesity, cancer screening or mental health. The experience from Portugal can be helpful to inform Brazil on the development of candidate indicators. Portugal has an extensive quality information infrastructure for PHC, which are collected as part of the contracting arrangements with PHC bodies (OECD, 2015^[57]). The indicators provide information to senior management to evaluate performance and achievement, benchmarked against other institutions (OECD, 2015^[57]). Indicators are collected across several domains – women's health and family planning; maternal health; child and youth health; hypertension; diabetes; cancer screening; mental health or patient experience (Table 4.5).

Table 4.5. Examples of indicators collected in Portugal

Domains	Indicators
Hypertension	Proportion of clients with hypertension, with at least one record of BMI in the last 12 months
	Proportion of users with hypertension with blood pressure recording in each semester
Diabetes	Proportion of users with diabetes, with at least one foot examination of recorded in the last year
	Proportion of clients with diabetes, with record management regimen (three items) in the last year
	Proportion of clients with diabetes, with consultation of nursing surveillance diabetes in the last year
	Proportion of clients with diabetes, with at least two HgbA1c tests in the past year
	Proportion of clients with diabetes, with the last record HgbA1c lower or equal to 8.0%
Mental health	Proportion of users with diabetes, with at least one or at least referencing a record of performing examination of the retina, in the last year
	Proportion of users aged over 18 years and a diagnosis of depression who were prescribed antidepressant therapy
Screening and prevention	Proportion of users aged over 65 years who were not anxiolytics or sedatives or hypnotics prescribed in the period
	Share of women with mammography in recent two years
	Proportion of users aged years with screening for colon and rectal cancer performed
	Proportion of users aged 14 and over, with quantification of smoking habits in the last three years
	Proportion of users aged over 14 years and with smoking, whom smoking-related consultation was held in the last year
Pharmaceuticals	Proportion of users aged 14 and over, with quantification of alcohol consumption, registered in the last three years
	Proportion of users aged over 75 years, with chronic lower than five prescription drugs
	Proportion of packaging billed drugs that are generic
Patient experience	Proportion of users aged over 65 years with no prescription trimetazidine in the last year
	Proportion of users satisfied or very satisfied
	Number complaints per 1 000 medical consultations conducted by doctors or nurses

Source: OECD (2015^[57]), *OECD Reviews of Health Care Quality: Portugal 2015: Raising Standards*, <https://dx.doi.org/10.1787/9789264225985-en>.

Another best practice example across the OECD is from the National Programme for Quality Indicators in Community Healthcare (QICH). The programme is used to monitor and improve the quality of preventive, diagnostic, and therapeutic PHC services in Israel. The indicators in QICH cover nine clinical areas: health

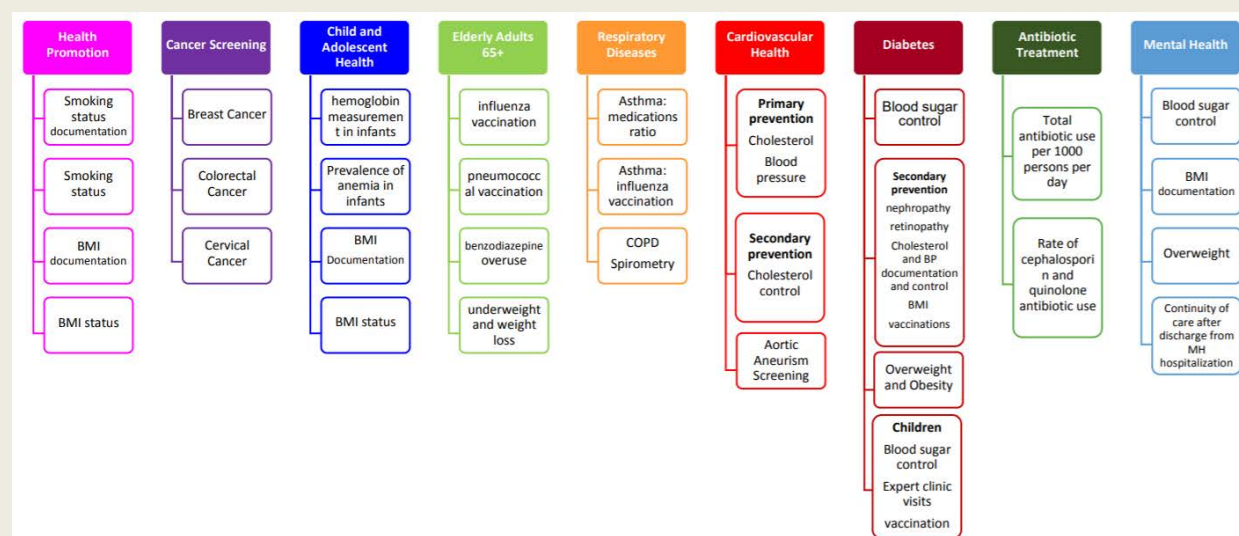
promotion, child and adolescent health, cancer screening (breast, colorectal and cervical cancer), elderly health, respiratory diseases, cardiovascular health, diabetes, antibiotic treatment and mental health. QICH incorporates a focus on primary prevention and management of chronic conditions, as demonstrated by the inclusion of indicators relating to risk factors in the general population, and the focus on COPD and asthma, diabetes, cardiovascular diseases and mental health (Box 4.4). Data quality for QICH is ensured through the use of standard indicator definitions by all health funds, and a systematic data quality audit cycle to ensure validity and comparability. Past evaluations have shown that the quality in community health care has improved over time for almost all categories of care – primary prevention, treatment and effectiveness of care (QICH, 2017^[58]). In the area of diabetes care for example, a recent evaluation has found an improvement in most quality indicators (from 53% to 75% for the composite quality score). Declines were noted in rates of blindness, diabetes-related end-stage kidney disease, lower limbs amputations and diabetes-related mortality (Calderon-Margalit et al., 2018^[59]). Quality improvements were also observed for elderly health (Podell et al., 2018^[60]). The extensive use of electronic health records by health plans are one key successful factor of the QICH.

Regular feedback to providers on quality and outcomes as well as open publication for patients with performance data at municipal level and FHT level could also drive aspiration for continuous improvement. On this front, Brazil will need to improve transparency at granular level, and ensure that patients can monitor activity and quality of PHC at FHT level.

Box 4.4. The National Programme for Quality Indicators in Community Healthcare in Israel

The indicators in QICH cover nine clinical areas: health promotion, child and adolescent health, cancer screening (breast, colorectal and cervical cancer), elderly health, respiratory diseases, cardiovascular health, diabetes, antibiotic treatment and mental health. In total 20-eight indicators are collected (Figure 4.9). These indicators are based on national and international guidelines reflecting the current scientific evidence, international practices, relevance for the Israeli Health care system, and the feasibility of production.

Figure 4.9. The National Programme for Quality Indicators in Community Healthcare



An internal audit is conducted within each health plan, the QICH programme's directorate also performs a data audit, and an external audit is carried-out by a certified external audit. Throughout the auditing

process, methodologies, control processes, documentation, and lessons learned were examined, nurturing a continuous improvement in indicator reporting.

Table 4.6. Performance results at national level for selected indicators, 2017-19

Domains	Indicators	2017	2018	2019
Health promotion	Documentation of BMI components (ages 20-64)	90.33%	89.51%	89.16%
Cancer Screening	Under-screening for cervical cancer (women aged 35-54 who were not screened in the last 5 years)	35.71%	34.64%	32.65%
Child health	Documentation of BMI components in children (age 14-18 years)	74.94%	76.71%	77.54%
Elderly health	Use of long-acting of benzodiazepines in older adults (aged 65 years or older)	2.29%	2.17%	1.88%
Respiratory diseases	Asthma medication ratio (AMR) greater than or equal to 0.5 (ages 5-45 years)	69.71%	70.67%	71.94%
Cardiovascular health	Rate of LDL cholesterol level less than or equal to 160 mg/dL in those at low-risk for heart disease (ages 35-74 years)	90.18%	90.35%	90.49%
Diabetes	Uncontrolled diabetes: HbA1c greater than 9% in individuals with diabetes mellitus (ages 18 years or older)	9.98%	9.60%	9.01%
Antibiotic	Documentation of molecular HCV test among those with a positive serology test	83.34%	85.55%	87.59%
Mental Health	Adequate control of HbA1c in individuals with SMI and diabetes mellitus (ages 18-84 years)	71.15%	71.83%	72.07%

Source: QICH (2017^[58]), National Program for Quality Indicators in Community Health Care in Israel, <https://en.israelhealthindicators.org/publications>.

To support greater patient-centred care, it is also important to collect patient-reported experience measures with PHC (OECD, 2021^[61]). This approach is taken internationally. In England for example, the GP Patient Survey assesses patient's experience of health care services provided by GP practices within the National Health Service (NHS) England (OECD, 2020^[43]). It assesses experience of access, making appointments, the quality of care received from health care professionals, patient's health and experience of their GP practice. Around 2 million patients registered with a GP practice are surveyed twice a year. The Care Quality Commission uses the results from this survey in their regulation, monitoring, and inspection of GP practices in England. The GP Patient Survey offers a description of each GP practice and its performance based on the latest survey. The analysis tool also provides comparison of performance between GP practices.

In 2017, the OECD launched the Patient-Reported Indicators Surveys (PaRIS) to address the need to understand the outcomes and experiences of people with chronic diseases (Box 4.5) (OECD, 2021^[61]). PaRIS is the OECD's Patient-Reported Indicator Surveys initiative where countries work together on developing, standardising and implementing a new generation of indicators that measure the outcomes and experiences of health care that matter most to people. The International Survey of People Living with Chronic Conditions will be the first of its kind to assess the outcomes and experiences of patients managed in PHC across countries. The PaRIS survey aims to fill a critical gap in PHC, by asking about aspects like access to health care, waiting times, as well as quality of life, pain, physical functioning and psychological well-being. The initiative will help open a dialogue with service providers about how to further improve the performance of health services and health systems to become more people-centred. Brazil might want to join the OECD PaRIS survey to understand how the outcomes and experiences of care in Brazil compare with those in OECD countries.

Box 4.5. The OECD PaRIS Survey

In 2017, the OECD launched the Patient-Reported Indicators Surveys (PaRIS) to address the need to understand the outcomes and experiences of people with chronic diseases. PaRIS offers an opportunity for gathering the evidence necessary to transform health care systems into patient-centred systems based on the needs of the people they serve. The initiative includes:

- the collection of validated, standardised, internationally comparable patient-reported indicators in three areas: hip and knee replacements, breast cancer care and mental health care
- the collection of a new set of internationally comparable measures which focus on patients with one or more chronic conditions, who are living in the community, and who are largely treated in PHC or other ambulatory care settings.

Source: OECD (2021^[61]), Patient-Reported Indicator Surveys (PaRIS), <https://www.oecd.org/health/paris/>.

The inclusion of the PCAT Tools in the 2019 National Health Survey is a positive step in Brazil to better understand users' experience with PHC services. This initiative should be maintained over the longer term.

4.5.4. Accreditation system for FHTs could help standardise and improve care quality

To standardise quality across the country, it is warranted to establish accreditation programme for PHC. Accreditation is a control mechanism to assure the quality of health care services, which often feeds into quality improvement (OECD, 2017^[63]). It uses explicit standards derived from the best available evidence to assess performance of health care organisation through survey, assessments, or audits. Accreditation is particularly needed when performance data are lacking.

Several OECD countries have established accreditation system in PHC, including Australia, Canada, England, New-Zealand or the United States (Tabrizi and Gharibi, 2019^[64]). In these countries, community oriented care, safe-care, high quality care, care continuity and human resource management had the highest priority among accreditation programmes (Tabrizi and Gharibi, 2019^[64]).

England's approach to health service accreditation is at the forefront of OECD efforts, and is a model for other health care systems to emulate (Box 4.6). It is unusually comprehensive as it accredits all providers of primary and social care. The accreditation is carry-out by the Care Quality Commission an independent statutory body responsible for registering care providers; monitor, inspect, and rate services, and take action to protect population who use services (Care Quality Commission, 2021^[65]). In Australia, the Australian Commission on Safety and Quality in Health Care, in collaboration with the Royal Australian College of General Practitioners, developed the voluntary National General Practice Accreditation Scheme. The Scheme aims to encourage safe and quality general practice in Australia. It involves independent third party review that practices meet the requirements set by the Royal Australian College of General Practitioners, and offer financial incentives to support activities that encourage continuing improvement in quality of care.

A starting point to improving quality in PHC will be to build accreditation system for all FHTs to carry out independent assessment of performance and identify areas that may require improvements. An accreditation system in Brazil could help standardise PHC in municipalities to achieve more consistent quality across the country.

Establishing a national inspectorate for health similar to the Care Quality Commission in England or the Haute Autorité de Santé in France could help to provide independent verification that standards are being met, identify good practice and support weaker centres to improve their standards. The new Agency for

the Development of PHC could also take this role, and act as an inspectorate for PHC. The Agency could for example be responsible for quality monitoring and improvement, and have a prescriptive role in producing overviews of current practice, current performance, and setting standards on performance and performance reporting. It could also develop tools such as evaluation framework, deep dive teams to visit and support municipalities with special needs, and levelling out resources when needed. This Agency could also provide a platform or forum for contact and exchange between municipalities, evaluating innovative practices and consolidating quality initiatives across the country when successful.

Box 4.6. Accreditation system in England is comprehensive

In England, the Care Quality Commission (CQC), an independent statutory body established in 2009, is responsible for the inspection for hospitals, adult social care, general practice, mental health care services, ambulances and community-based services. All providers of regulated activities, including NHS and independent providers, have to register with CQC and follow a set of fundamental standards of safety and quality below which care should never fall. The CQC assesses if providers are meeting these fundamental standards through monitoring and inspection.

The findings of such assessments are shared with the public, and citizens are encouraged to share their experience or report concerns to the CQC.

England's health care system is one of the few in the OECD to have a comprehensive accreditation programme for PHC. By March 2020, 7 237 GP services have been inspected and rated, while only 1 461 sites have not. Detailed individual practice inspection reports are publicly available on a website (Figure 4.10). The core services being inspected are urgent and emergency services, medical care, surgery, critical care, maternity and gynaecology, services for children and young people, end of life care, outpatients' services and diagnostic imaging. After inspection, the GP services must respond to areas of concern that was identified, develop an action plan to address them and make improvements. Detailed individual practice inspection reports are publicly available on a website.

Figure 4.10. Example of a CQC inspection in a GP surgery

CQC inspection area ratings	
(Latest report published on 8 March 2021)	
Safe	Inadequate 
Effective	Requires improvement 
Caring	Good 
Responsive	Requires improvement 
Well-led	Inadequate 

Source: Care Quality Commission (2021^[66]), CQC inspection ratings for GP services, <https://www.cqc.org.uk/search/services/doctors-gps?sort=default&distance=15&mode=html>.

4.5.5. Designing smarter payment for PHC

The *Previne Brasil* scheme is an important scheme to drive quality improvement in Brazil. In the years to come, it will be important to maintain and even expand the scheme. Compared to other OECD countries, the incentivised priority areas in Brazil have a stronger focus on child and maternal health. In OECD countries, the management of chronic conditions is frequently put at the centre of P4P schemes (OECD, 2020^[43]). The *Previne Brasil* programme plans to expand the number of indicators by 2022 to include other indicators around mental health, breast cancer, multi-professional action, and patient experience (Ministério da Saude, 2020^[67]). This will be an important step given the rising burden of chronic conditions and the need to address some risk factors for health including obesity. The experience from Portugal or England could inform the development of candidate indicators to be incorporated to the *Previne Brasil* programme.

A second key difference between the Brazilian P4P compared to most OECD P4P schemes is the fact that bonus payments are not made to the individual physician or the structure that he/she works for but to the municipalities that employs him/her, with no obligation for municipalities to pass on their bonuses to staff. This is not common in OECD countries in PHC. Generally, research suggests that P4P schemes with bonus payments aimed directly at professionals are more effective (Eijkenaar et al., 2013^[68]).

Table 4.7. New forms of payment for PHC across the OECD

Add-on payments	Bundled payments	Population-based payments
Australia, Austria, Canada, Denmark, France, Germany, Iceland, Israel, Italy, Mexico, Sweden	France, Italy, the Netherlands, Canada, Australia, Belgium	United States (Accountable Care Organisations), Germany (Gesundes Kinzigtal GmbH)

Source: OECD (2020^[43]), Realising the Potential of Primary Health Care, <https://doi.org/10.1787/a92adee4-en>.

Lastly, there are successful experiences across the OECD where innovative forms of payment for PHC have been introduced to encourage greater prevention, management of chronic disease and incentivise providers to effectively work together. Beyond P4P programme, there are three innovative forms of payments across OECD countries (OECD, 2020^[43]) (see also Table 4.7):

- **Add-on-payments:** Additional payments to remunerate specific activities in order to target some dimensions of the care provisions such as establishment of care plans, collaborative care meetings or the provision of patient education. In 2018, 11 OECD countries used add-on payments to pay for co-ordination or prevention. In Denmark, general practitioners get pay-for-co-ordination, notably when they have more responsibility for treatment of chronically ill patients (e.g. diabetes).
- **Bundled payments:** The objective is to bundle separate health care activities into a single tariff for chronic conditions. The single tariff should cover the cost of all health care services provided by the full range of providers during a specific time period. In this type of arrangement, a group of health care providers is collectively responsible for the delivery of a wide range of activities for a pre-defined population group, typically involving at least PHC and secondary care. Evidence shows that bundled payments lead to better collaboration within and across care settings and can contribute to a greater standardisation of care and to the development of sophisticated IT systems. In 2018, 5 OECD countries used bundle-payments. The Netherlands have for example established bundled payments to care groups for patients diagnosed with Diabetes type 2, COPD and cardiovascular diseases.
- **Population-based payments:** This payment scheme is an extension of bundled payment, being not restricted to particular episodes of care or conditions. It consists of one payment made to groups of health providers as well as management companies to cover most health care services for a

defined group of the population. Rather than paying providers in “silos”, the money follows the patients across providers, covers most health care services and has a more comprehensive view of population well-being. In 2018, only two OECD countries used population-based payments, including for example the “Gesundes Kinzigtal” Initiative in Germany.

4.5.6. Expand the development of *Rede de Atenção à Saúde* to promote integrated care

Given the context of an epidemiological transition towards longer lives and increasing prevalence of chronic conditions, an important challenge in Brazil is to shift the focus of health care services towards addressing longer episodes of health care needs. To this end, greater integration between PHC, emergency care, specialist care and social care services is needed. Interactions between various health care providers about patient cases, and transitions from one service to another, should be timely, safe, and seamless for the patient and its family (OECD, Forthcoming^[69]). PHC needs to play a key role in achieving this goal.

The impact of integrated care include better access, improved satisfaction for patients and health professionals, more appropriate care, enhanced preventive care, reduced avoidable hospital admissions, prolonged independent living and delayed admission to institutional care, improved health status and quality of life (Nolte and Pitchforth, 2014^[70]; Curry and Ham, 2010^[71]).

In Brazil, the *Rede de Atenção à Saúde* (RAS) are ambitious health care network aimed at reducing health fragmentation within the Brazilian health system. As earlier mentioned, the implementation of RAS is supported by Planificação da Atenção à Saúde (PAS) and Chronic Conditions Care Model (MACC) developed by CONASS (CONASS, 2020^[13]; CONASS, 2018^[14]). RAS is included in the scope of the Regiões de Saúde, with technical, logistics and managerial support to encourage effective integration.

However, implementation of RAS in Brazil is challenged by bureaucratic governance model and weak institutional leadership, workforce issues such as inadequate training and qualification toward inter-professional collaboration and co-ordination, a weak information system and payment policies that do support integrated care (Ministério da Saude, 2014^[72]). These have resulted in low and uneven diffusion of RAS initiative across the country. Brazil will need to work on these four fronts to promote integrated care and ensure effective implementation of RAS (Box 4.7).

Brazil could learn from other OECD countries, which have embarked on different pathways to operationalise the integration of care, such as the Spain Basque Country with Integrated Care Organisations, the United Kingdom with PHC Networks and the United States with Accountable Care Organisations.

Box 4.7. Promoting integrated care and ensuring effective implementation of RAS in Brazil

Stronger leadership and governance model

The importance of leadership and management for the successful implementation of integrated care as well as integrated care service delivery as already been demonstrated in recent studies (Borgermans and Devroey, 2017^[73]). While there is no one size fit all models, consolidated strategies seem often to be developed based on consultations with main stakeholders so that they can be implemented effectively by holding them collectively accountable and committed to achieving the common goals of promoting integrated care delivery.

Better workforce planning and modern educational training

To ensure that PHC workers work together with other health care professionals, there is a need to increase effort to develop appropriate training towards multi-disciplinary teams, case management, and

developing new roles for care co-ordination. In Germany, a national training programme focussing on case management was made available for nurses to strengthen their role in delivering care co-ordination. The Basque country in Spain created new roles such as liaison nurses and referral internists to co-ordinate and organise care. In Australia also, care co-ordinators for cancer patients or patients with chronic long-term conditions have been established to liaise health professionals, interface with the patient's health care team, help arrange health screenings, resolve barriers to accessing the health networks, and source follow-up health education.

Stronger data and information systems

Strong data and information systems are critical to support integrated care. With growing number of patients with complex needs receiving care by various services over long episodes of care, such systems can help improve communication and co-ordination between patients and their providers, increase the accuracy of diagnoses and clinical decision-making, help monitor and deliver appropriate services remotely, and provide patients more access to their health history. In Brazil data linkage remains insufficient, undermining information sharing in the context of integrated care (see (OECD, 2021^[41]).

Smarter payment models

Payment models need to incentivise providers from different health sector to effectively work together. Fragmented governance in Brazil may undermine resource allocation across health care, hinder integrated care delivery. At the same time, there are some good providers' payment mechanisms that encourage greater co-ordination and integration of care – such as add-on payments, bundled payments or population-based payment. Add-on payments reward more co-ordinated, safer and effective care, and are widespread across OECD countries in PHC (Table 4.7). Population based payment for example pool into a single payment all services delivered to patients including prevention and treatment across hospitals, primary and long-term care. In the United States, ACO receive a virtual budget for a wide range of services and they are incentivised by shared savings contracts by payers to co-ordinate care across different levels efficiently.

Source: OECD (2020^[43]), Realising the Potential of Primary Health Care; OECD (Forthcoming^[69]) Strengthening Performance of Integrated Care Delivery across OECD countries.

4.6. Conclusion

Looking to expand access to high quality PHC, Brazil has been taking important steps to improve the distribution of doctors, develop new forms of service organisations and introduce new financing models for PHC. The development of guidelines and health care services portfolio; and the production and publication of data underpinning PHC are also important steps towards ensuring high quality care across the country. All these developments are well aligned with the experiences of OECD countries, and have led to large improvements in access to PHC and population health outcomes.

However, the Brazilian PHC system is still characterised by a relatively low population coverage. A significant proportion of the population also have direct access to outpatient specialties and hospitals for minor conditions, perhaps because PHC has traditionally provided few low complexity procedures and surgeries. Available evidence also points to the need for enhancing the contribution of PHC to prevention and management of chronic conditions, notably in North and Northeast regions which present consistently poorer health care.

These shortcomings give strong arguments to strengthen the gatekeeping system in Brazil with a systematic registration system with FHTs, which will control and orient the patient's into specialist care. This is a prerequisite to accelerate the expansion of FHTs, helping to build a profile of the health needs of the registered population, and better recognise the central co-ordinating role of FHTs. Actions are also needed to support FHTs to deliver a more modern PHC service provision, focused on prevention, treatment, follow-up of diagnoses and management of chronic conditions. The federal government will need to play a more prescriptive role to strengthen comprehensive PHC evenly across the country. These would include ensuring that all municipalities are well equipped and trained to properly deliver actions and services defined in the CaSAPs. There are also opportunities to monitor and encourage compliance with clinical guidelines and CaSAPs using the *Previne Brasil* Programme – either through the Strategic Actions or the P4P component. Collecting a richer set of quality indicators on a wider array of preventive activities and management of chronic conditions (such as alcohol consumption, obesity, cancer screening or mental health, and patient experience) would also support greater patient-centred PHC. An accreditation system in Brazil could help standardise PHC quality – notably in municipalities having low capacity and high vulnerability – to identify areas that may require greater financial and organisational support. The new Agency for the Development of PHC could act as an inspectorate for PHC, for example to provide independent and external verification that standards are being met, identify good practice and support weaker centres to improve their standards.

Going forward, Brazil will have to establish strong oversight and good management, develop appropriate training, alongside strong data and information systems for the successful implementation of RAS. Appropriate payment models would be critical to incentivise providers from different health sectors to effectively work together. While this may come too early at the moment, the introduction of add-on payments or bundled payments is certainly something Brazil could consider in the future.

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Notes

¹ See <https://aps.Saúde.gov.br/biblioteca/index/MQ==/Mg==>

² See for example <http://www.ripsa.org.br/category/publicacoes-ripsa/>

³ Available at <https://biblioteca.ibge.gov.br/index.php/biblioteca-catalogo?view=detalhes&id=2101758>

5 Workforce challenges in primary health care in Brazil

Brazil is struggling with a shortage and an uneven distribution of medical doctors across regions, compounded with a low credibility and recognition of the PHC specialty. To secure a greater number and distribution of primary care doctors and ensure a high-quality workforce, the first priority for Brazil is to implement a coherent workforce planning based on an objective assessment of present and future needs to govern health care human resources. In tandem, Brazil could look at the experiences of OECD countries to train more rural doctors and to provide both financial and non-financial incentives linked with a return of service obligation. A smarter use of nurses and community health workers is another option to cope with workforce gap. There are also opportunities to make the PHC specialty a mandatory requirement to be allowed to practise PHC, and to implement stronger requirements around continuous medical education.

5.1. Introduction

As many other OECD countries, Brazil is struggling with a shortage and an uneven distribution of medical doctors across regions. Practicing doctors per 1 000 population in 2019 were lower than the OECD average (at 2.3 compared to 3.5), while the distribution of medical doctor followed a North-South gradient where North and Northeast regions show a lower density of medical doctor. At the same time, the specialty of family and community medicine (FCM) is not attractive and undervalued in Brazil. Very few doctors working in PHC have undertaken specialist training. In 2018, only 5 486 medical doctors had a specialty training in FCM, representing 1.4% of all specialists in Brazil. While people living in North and Northeast regions have the highest health care needs, these regions gather less than 20% of all FCM specialists.

In recognition to these challenges, several programmes were implemented by the federal government to reduce the shortage of doctors in prioritised regions; and improve access to and quality of medical training in the country, including the More Doctors Programme (*Mais Médicos para o Brasil*). However, unfilled PHC positions is still the norm in Brazil, gaps in medical education remain between medical schools and geographic disparities in access to medical training persist. The North and Northeast regions offer the lowest levels of per capita medical undergraduate positions in the country.

This chapter examines workforce challenges for the PHC setting in Brazil. Section 5.2 describes the shape of Brazil PHC workforce in terms of the rates and ratios of medical doctors, nurses and community health workers. In Section 5.3, the chapters describes the organisation and quality of medical education in Brazil. In Section 5.4, the chapter then explores ways to improve the geographical distribution of doctors and overcome the issue of shortage of PHC doctors. Section 5.5 considers how to improve the recognition and credibility of the specialty of FCM and points out some key policy levers to strengthen the quality of medical education in Brazil.

5.2. The shape of Brazil's PHC workforce

Brazil's health workforce is characterised by a shortage and a misdistribution of medical doctors that is particularly acute in the North and Northeast of the country. Brazilian figures suggest that medical doctors do not perceive the specialty of PHC, and rural and remote areas, as attractive and prestigious. The shortage and misdistribution of PHC physicians is a common health workforce policy challenges OECD countries face. Concerns about income, opportunities for career development, isolation from peers, as well as educational options for children and professional opportunities for their spouse are leading factors of these challenges.

5.2.1. Physician density in Brazil remains lower than the OECD average, with persistent concerns over geographic distribution

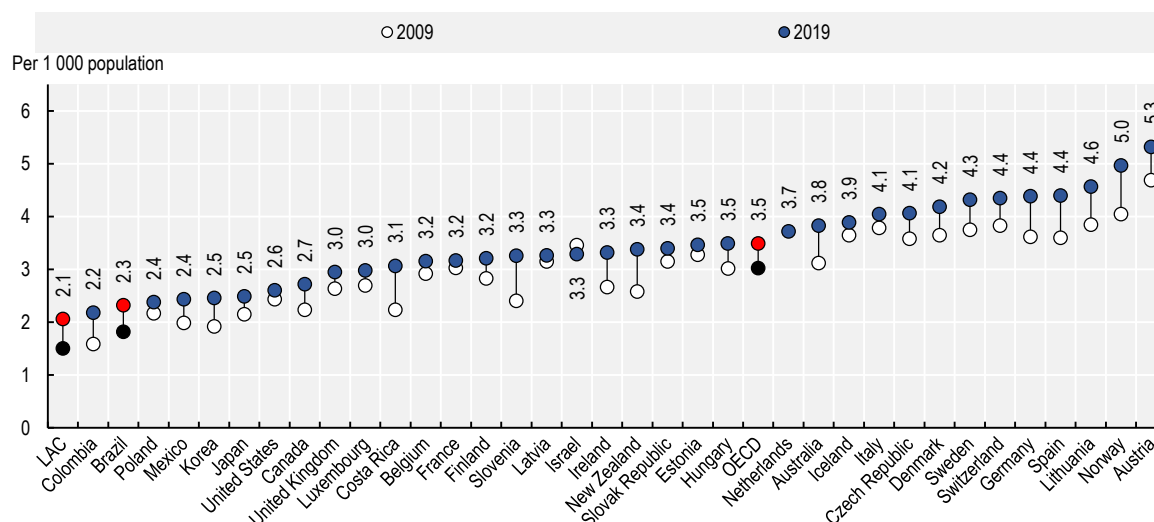
While Brazil has clearly recognised that a strong PHC is the foundation of an effective, efficient and responsive health system, the number of trained PHC physicians is well below OECD countries and of other LAC countries.

In Brazil, practicing doctors per 1 000 population in 2019 were very low compared to the OECD average, at 2.3 compared to 3.5 (Figure 5.1). This is among the lowest density, just above Colombia (with 2.2 doctors per 1 000 population) and the LAC average (2.1 doctors per 1 000 population). However, the number of doctors increased by 50% in Brazil over the past decade.

According to the Ministry of Health, there are around 60 000 medical doctors working in PHC in Brazil. Although a distinct specialty of FCM is established in Brazil, very few doctors working in PHC have undertaken specialist training in FCM. In 2018, only 5 486 medical doctors had a specialty training in FCM, representing only 1.4% of all specialists in Brazil (Scheffer et al., 2018^[1]). Medical doctors who specialise

in internal medicine also have some competencies applicable to FCM, but they also represent a small share of specialists (around 11.2%) in Brazil in 2018. Taken together, these figures suggest that Brazil struggles to attract physicians to specialise in PHC. The low attractiveness of the PHC specialties can be linked to a lower recognition, prestige, and remuneration level compared to other specialties.

Figure 5.1. Doctors per 1 000 population, 2009 and 2019 (or latest year available)



Source: OECD (2021^[2]) Health Statistics Database, <http://stats.oecd.org>; Brazilian Ministry of Health.

Regional differences in the distribution of all physicians is an important concern in Brazil. As in many OECD countries, the number of doctors per capita varies widely across regions. While overall there are 2.3 doctors per thousand population in Brazil, there are capitals with more than 12 doctors per 1 000 population (Vitória, in Espírito Santo for example), and regions with values below one doctor per 1 000 population (the Amazon Region in the North) (Scheffer et al., 2018^[1]). The differences between small and large municipalities in Brazil is comparable to other OECD countries.

The uneven distribution of doctors holds true when looking at some specialities, for example for the FCM specialist. Figure 5.2 indicates that some states in the North and Northeast regions have the fewest doctors per population, while the best-supplied region is the South and Southeast regions, followed by the Central-West (Scheffer et al., 2018^[1]).

On a state level, the FCM specialist per 1 000 population ratio ranges from 0.4 in Maranhão in the Northeast region to 1.6 in Rio de Janeiro. Overall, the North region of Brazil gather only 4.6% of all FCM specialists, while the South and Southeast regions gather almost 75% of all FCM specialists (Figure 5.3).

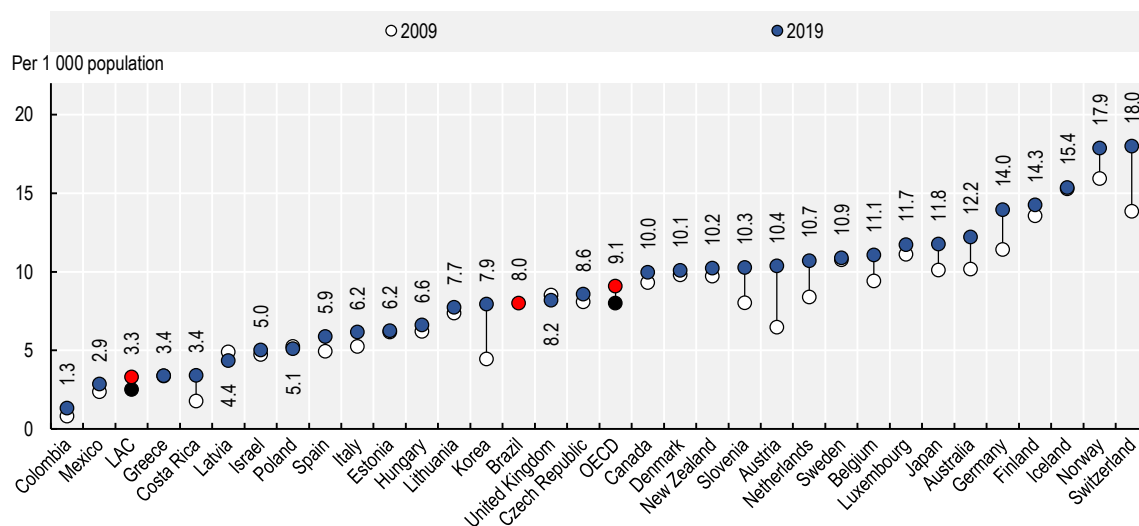
It is also important to note that geographical imbalances in medical doctor (and PHC physicians), is disproportionately experienced by people living in remote and rural areas who have the highest health care needs, and where social vulnerabilities are highest (Oliveira et al., 2017^[4]). This means that the PHC sector faces a double challenge of high health care demand and staff shortages, which can lead to inequitable access to care (Barbosa and Cookson, 2019^[5]). Shortages and geographical imbalances in workforce need to be addressed as a matter of priority as it can lead to lower health outcomes, lower health care quality and unmet needs for population living in underserved regions (see Chapter 2 and Chapter 4).

5.2.2. Brazil nursing workforce has significantly increased over the past decades

In 2019, Brazil had 414 712 nurses, and annually produced about 50 000 nurses in their undergraduate nursing programmes (Cassiani and Silva, 2019^[6]). In the Latin American region, Brazil is the only country that trained nurses in professional master's degree, which grew by 156% in the period 2011-20.

Brazil's nursing workforce has increased over the past decade to reach eight nurses per thousand population in 2019, moving closer to the OECD average of 9.1 (Figure 5.4). This is much higher than the LAC average of 3.3 nurses per 1 000 population. The increase in the volume of nurses is mainly explained by the implementation of new federal regulations, including the creation of specific PHC teams (Riverside team, Health Academy Programme, the Home care teams among other), requiring nurses.

Figure 5.4. Nurses per 1 000 population, 2009 and 2019 (or latest year available)



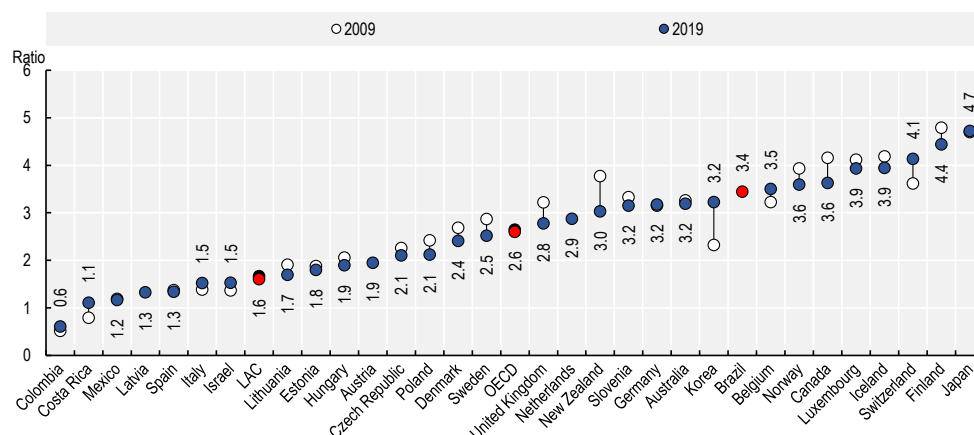
Source: OECD (2021^[2]) Health Statistics Database, <http://stats.oecd.org>; Brazilian Ministry of Health.

The skill-mix – the ratio of nurses to doctor – is relatively higher in Brazil (Figure 5.5) than the OECD and LAC average. Nurses outnumber doctors in Brazil at 3.4 nurses per medical doctor, compared to the OECD average of 2.6 and the LAC average of 1.6.

This is a welcome news to respond to the changing demand for health services. There is a need for a better health professional skill-mix in the context of rapidly ageing populations and rising burden of chronic conditions in Brazil (see Chapter 2). Currently, there is no discussion around the expansion of nursing role or implementing advanced functions for nurses, as it has been introduced with some success in other OECD countries such as Australia, Canada, the United States, the United Kingdom, some Nordic countries, and France more recently. In these countries, advanced practice nurses provide greater support in the care of chronically ill patients and those with complex morbidities, working in co-operation with PHC

doctors and specialists in teams and other health and long-term care settings. Improved interaction and communication between members of FHTs is critical for the success of task shifting and redistribution of roles.

Figure 5.5. Ratio of nurses to doctors, 2009 and 2019 (or latest year available)



Source: OECD (2021^[2]) Health Statistics Database, <http://stats.oecd.org>; Brazilian Ministry of Health.

5.2.3. Community health workers play a crucial role in PHC

Following the implementation of the Community Health Agents Program (PACS), CHWs started to play a crucial role in the Brazilian PHC system. CHW are frontline public health workers, often members of the communities in which they work and have therefore valuable knowledge of people needs and are trustful source of information (Grossman-Kahn et al., 2018^[7]; Wadge et al., 2016^[8]). While a range of different health professionals are involved in PHC delivery (physicians, nurses, nutritionists, pharmacists or social workers, etc.) (see Chapter 2), there are up to 12 CHWs involved in each FHT. In 2020, more than 250 000 CWHs served 61% of the population, up from 60 000 in 1998. The population covered by CHWs is the highest in the Northeast and North regions (with at least 74% of the population covered by CHWs) (Figure 5.6).

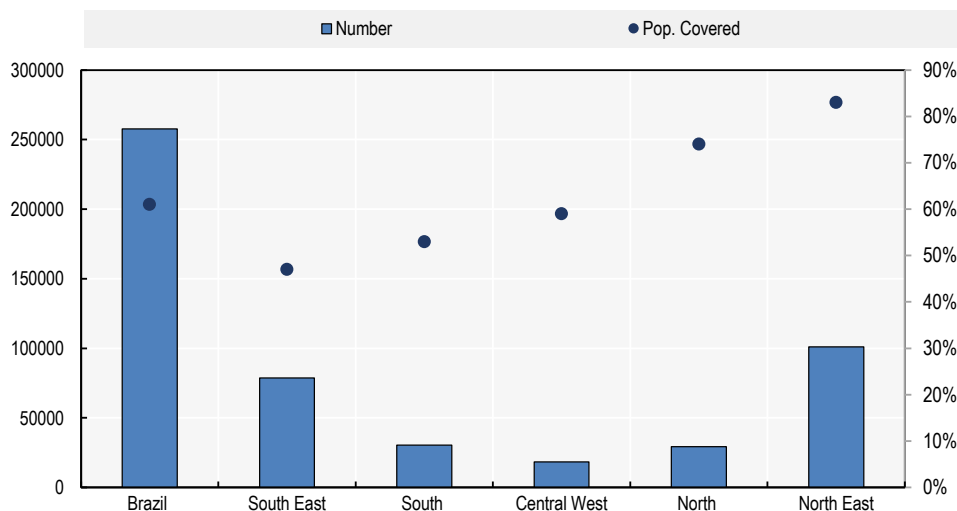
Unlike many OECD countries, in Brazil, CWHs provide a wide range of support for comprehensive PHC services including clinical triage, health promotion, social determinants and household data (Grossman-Kahn et al., 2018^[7]). They are expected to resolve many low-level problems such as medication review for patients chronically ill, while referring more complex issues to doctors and nurses of the FHT. CHWs complete between 10-15 home visits a day, and spend between 1-2 hours a day at the clinics doing paperwork, organising the waiting rooms, consultation appointments for other health professionals and offering educational sessions. They are responsible for registering the households in the areas where they work, and help families access other government services such as conditional cash transfers and social services. Within FHTs, each CHW is responsible for 150 families.

The minimum requirement to become a CHW is to get a secondary education, but more than 60% have a professional diploma. Turnover is low as it is a stable and respectful profession (Wadge et al., 2016^[8]).

Despite their importance for the provision of PHC services, studies have shown that CHWs do not have many technological supports to operate in the community. Evaluations show that CHWs are effective at improving specific health outcomes and access to health care (Grossman-Kahn et al., 2018^[7]), and that it can represent a cost-effective intervention to deliver essential health care services (Vaughan et al., 2015^[9]). Yet, available evidence also highlights that, in many remote and vulnerable municipalities, CHWs

are not equipped with mobile phones and tablet computers to enable remote diagnoses and real time communication with the PHC facilities (Wadge et al., 2016^[8]).

Figure 5.6. Number of community health workers by regions



Source: Ministério da Saúde (2020^[10]), Painéis de Indicadores : Agentes comunitários de saúde, <https://sisaps.saude.gov.br/painelsaps/acs>

Overall, the COVID-19 pandemic demonstrated the importance of having a skilled and sufficient health workforce to respond swiftly and effectively to the public health crisis. Health professionals (doctors, nurses and community health workers) are at the forefront of the crisis, and the pandemic has put huge pressure on them, affecting their physical and mental health. In Brazil, as well as in OECD countries, investments in human resources for health are critical (see Box 5.1).

Box 5.1. The COVID-19 pandemic demonstrated that investments in human resources for health are critical

Across OECD countries, the pandemic is making pre-existing shortages of doctors and nurses more visible and acute. Shortages of doctors and nurses result from a lack of investment in education and training programmes and in recruitment of new staff and retention of existing staff. These shortages, in turn, resulted in heavy workload and burnout among health workers.

As doctors, nurses and other health professionals mobilised on the frontline to respond to the COVID-19 pandemic, OECD health systems sought ways to increase the number of staff available during the peak of the pandemic and to make the most efficient use of their work. This included recalling inactive and retired health professionals, offering them some short training to update and upgrade their skills (Italy, Portugal, the United Kingdom for example). Other countries mobilised or established a “reserve list” of health workers to deal with the outbreak and reallocate staff across regions (France, Belgium, Ireland and Iceland). COVID-19 has also accelerated innovative roles of health professionals, removing pre-existing structural and institutional barriers and permitting a more efficient use of human capital in the health sector. In Canada, for example, pharmacists have been authorised to extend prescriptions beyond what they were previously allowed to do, and to prescribe certain medications.

Source: OECD/European Union (2020^[11]), *Health at a Glance: Europe 2020*, <https://doi.org/10.1787/82129230-en>.

5.2.4. Health workforce planning is at best very limited in Brazil

In Brazil, the federal agency called *Instituto Nacional de Estudos e Pesquisas Educacionais Anísio Teixeira* (INEP) monitors workforce trends to support policy planning. INEP is also responsible for monitoring and assessing the basic and higher education system, and for promoting the dissemination of indicators and results of evaluation of the education system. The Secretaria de Gestão do Trabalho e da Educação na Saúde (SGTES) at the Ministry of Health is responsible for formulation of policies for health workforce management, training and qualification of health workforce, regulations of health professionals and regulation of the decentralisation process. It was created as the main federal instrument to mobilise and manage health care workforce. It defines the National Human Resources Health Policy for the good performance of SUS, in partnership with the Ministry of Education.

However, there is a lack of knowledge of present and future needs for doctors and other health care professionals in Brazil (Oliveira et al., 2017^[4]). Health care workforce planning is not based on an objective assessment based on the characteristics of the professionals (age, and sex), the work processes (productivity, workload), the characteristics of the health system in place (coverage and type of services offered), and population health needs (such as socio-economic and epidemiological profiles).

Rather, existing estimates are built on feasibility concerns, using data comparing ratio of doctors per thousand population between Brazil and other countries to determine a desirable ratio and create targets for improvement (Oliveira et al., 2017^[4]). A fixed minimum ratio target can be defined, but without considering the contextual factors described above to make a forecast of anticipated needs and demands, and a simulation of the necessary profiles and possible reform scenarios that would lead to a set of human resources better suited to meet the increasing demand for care.

Box 5.2. The workforce planning in Italy is based on a bottom-up approach

In Italy, there is a bottom-up approach for human resources planning in the Italian Servizio Sanitario Nazionale. Primary responsibility for health workforce planning is at the regional level, with information then fed back to the Ministry of Health, which brings together the data and forecasts from the regional levels, and analyses and validates the results to make appropriate recommendations to the Ministry of Education concerning entry to medical, nursing and other health-related education programmes.

The main objective of the health workforce planning in Italy is to ensure a suitable number of health care professionals in order to satisfy demand and to avoid workforce imbalances in the Servizio Sanitario Nazionale. The Ministry of Health and the regions then agree on the number of students to enter related education and training programmes.

Source: OECD (2014^[12]), *OECD Reviews of Health Care Quality: Italy 2014: Raising Standards*, <https://doi.org/10.1787/9789264225428-en>.

This is unlike many OECD countries such as Italy or the United States, which have strong bases for workforce planning. In the United States, the National Centre for Health Workforce Analysis offers health workforce research, data, tools and projections to inform programme planning and policy making (HRSA, 2021^[13]). In Italy, national needs in human resources are established based on regionally reported needs to establish the NHS human resources planning (Box 5.2). It is based on a bottom-up approach where information from the Regions is fed back to the Ministry of Health to make recommendations to the Ministry of Education (OECD, 2014^[14]). Experiences from OECD countries can be useful for Brazil in its efforts to define a clearer workforce management policy to govern health care human resources. Brazil already collects municipality-level data on a monthly basis, meaning that the pre-requisite data is already there to inform workforce management. A smarter use of those data is warranted to better understand workforce

challenges in Brazil, including for example the shortage and uneven geographic distribution of doctors. Greater co-ordination between municipalities and federal level could help in this direction.

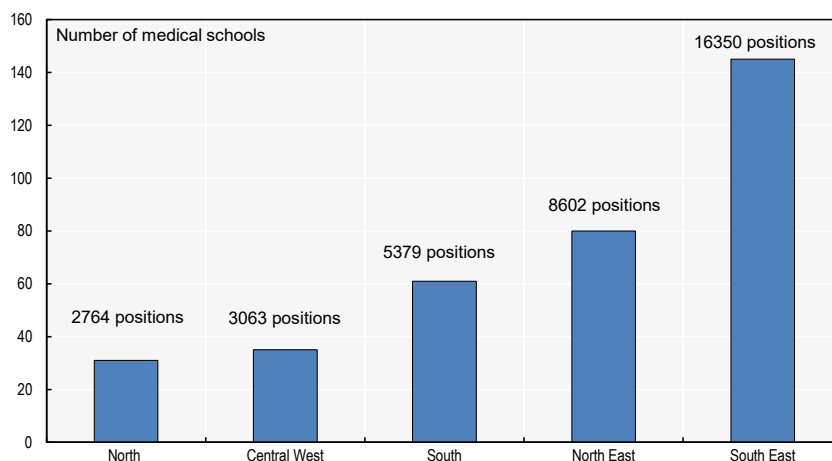
5.3. The organisation and quality of medical education in Brazil

Over the past decades, the number of medical schools has grown exponentially, driven mainly by the opening of private institutions. While some programmes have been set up to improve the medical curriculum toward PHC and achieve consistent quality across curriculums, gaps in medical education remain between medical schools and geographic disparities in access to medical training persist. The North and Northeast regions offer the lowest levels of per capita medical undergraduate positions in the country. Last but not least, the specialty of family and community medicine is undervalued in Brazil, and most doctors working in PHC in Brazil do not have a specialist qualification.

5.3.1. The number of private medical schools has grown exponentially in Brazil

In Brazil, the distribution of medical schools is uneven across the country. In 2020, the Southeast region had the highest distribution of medical schools among all regions (with 145 medical schools and 16 350 positions) (Figure 5.7). The same region concentrates around 45% of all positions in Brazil, with only one-fourth being offered in public institutions. At the other end of the scale, the North region concentrates 8% of all positions in Brazil.

Figure 5.7. Medical schools and positions by Regions, 2020



Source: Escolas Médicas do Brasil (2021^[15]), Escolas Médicas do Brasil, <https://www.escolasmedicas.com.br/index.php>.

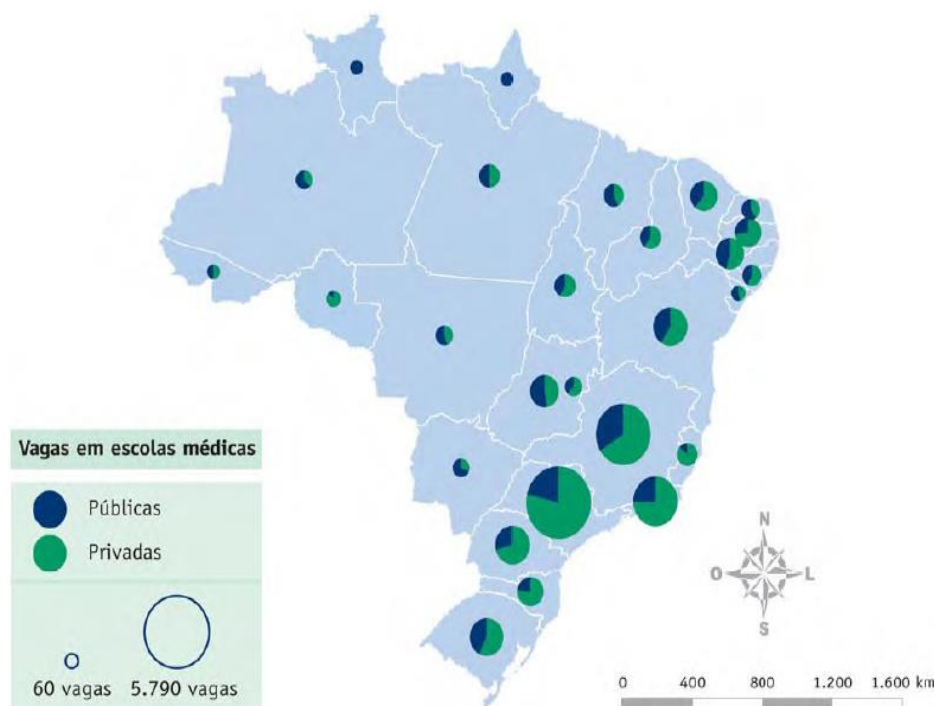
Unlike many OECD countries, medical education in Brazil has been marked by privatisation (Scheffer and Dal Poz, 2015^[16]; Scheffer et al., 2018^[1]). In addition to being larger in number, private medical schools also offer more places. In 2020, there were 352 medical schools, offering more than 35 000 training positions. Of these schools, 41% were public and 59% private (Escolas Médicas do Brasil, 2021^[15]). Public medical schools are free, while tuition at private medical schools varies from USD 700 to USD 2 500 (BRL 3 600 to BRL 13 000) per month (Escolas Médicas do Brasil, 2021^[15]). The country has around 14 places in medical schools per 100 000 population.

Throughout the country, public training positions represent 35% of the total training positions. In the South, 33% of training positions are public, in the Northeast 42% are public; in the North 48% are public, and in the Central-West 53% are public. Figure 5.8 presents the distribution of public and private medical schools

by States. In the State of São Paulo – which gathers 20% of training positions – only 21% are public. At the other end of the scale, the States of Roraima and Amapá only have public medical positions.

Private medical schools target niche markets, and generally offer education in more restricted and prestigious areas of knowledge (Scheffer and Dal Poz, 2015^[16]; Scheffer et al., 2018^[11]). Given that private medical schools charge high entry fees, the current admission system fosters inequality of access to medical school, favouring students from higher socio-economic background that do not specifically wish to work in PHC area and in rural areas.

Figure 5.8. Distribution of public and private medical schools in Brazil, 2018



Source: Scheffer et al. (2018^[11]), *Demographia Médica No Brasil 2018*.

5.3.2. Medical education in Brazil lasts six years

Undergraduate programme in Brazil is similar to many OECD countries. Physicians trained in Brazil follow an undergraduate programme lasting six year. The curriculum is divided into three cycles of two years (Conselho Federal de Medicina, 2020^[17]). During the first two years, the *ciclo básico* instruct basic scientific education with activities integrated to several medical specialties, allowing the practical application of such content. The next two years is called *ciclo clínico* with continued courses alongside clinical experience in medical setting (generally hospital setting). The last two year – *ciclo do internato* – focuses on clinical practice, through training in teaching hospitals and clinics. During the *ciclo do internato*, physicians actively participate to the medical practice under the direct supervision of medical schools. There are four six-month clinical rotations through internal medicine, surgery, paediatrics/obstetrics/gynaecology and public health, of which the latter includes exposure to PHC. In Brazil, graduate students are awarded the title of physician (Médico), allowing them to register with the Regional Council of Medicine (*Conselho Regional de Medicina*).

Unlike many other OECD countries (such as Canada, France or Italy), there is no national examination to be placed on a national register and be allowed to practise. Some states have made progress towards the

introduction of an Examination in Medicine, to be registered and allowed to practise. For example, the Regional Council of Medicine of São Paulo (*Conselho Regional de Medicina do Estado de São Paulo*) established that physicians who graduate from 2012 must pass a test to obtain professional registration.

Following the licensing, physicians can work as a “generalist” or choose among various professional path depending on the postgraduate specialisation programme they attend. Specialisation can last two to five years, and take place in a medical school. There are two types of specialisation programme. First, direct access programme where any graduates can apply to examinations for these specialties regardless of training or prior experience. Second, prerequisite programme, where doctors should have already completed a specialty or achieved expertise.

As of February 2019, the Federal Council of Medicine, the Brazilian Medical Association and the National Commission of Medical Residency Brazil recognise 54 specialty programmes (Scheffer et al., 2018^[1]). In 2018, among the total number of medical doctors, 169 479 (37.5%) were generalists with no specialty training and 282 298 (62.5%) had a specialty training. Four specialty training concentrate 40% of all specialists in 2018 (Clínica Médica, Pediatria, Cirurgia Geral, and Ginecologia e Obstetrícia).

The specialty of FCM is not recognised as a knowledge area by the Federal Agency for Support and Evaluation of Graduate Education. However, the specialty of community general medicine was officially recognised in 1981 by the National Commission on Medical Residency (Giliate Cardoso Coelho, Valeska Holst and Aristides, 2019^[18]). The Brazilian Society of Family and Community Medicine (SBMGC) was also created at that time, followed by the creation of the Brazilian Society for Family Medicine (SOBRAMFA) (Box 5.3). Since 2001, the specialty is called Family and Community Medicine, with medical residencies lasting two years. In 2018, there were more than 100 FCM residency programmes in the country, with around 1 600 training positions each year (representing 4.4% of all training position) (Scheffer et al., 2018^[1]). Although few in number, some post-graduate programmes in Family and Community Medicine are well known in Brazil such as the Conceicao Hospital Group (with 22 training positions each year), or the Municipal Health Department of Fortaleza in the State of Ceará (with more than 50 training positions each year) (Justino, Luzón Oliver and Palhano de Melo, 2016^[19]). Most medical residencies, master’s and PhD degrees in FCM are located in the Southeast and South Regions (Fontenelle et al., 2020^[20]).

While the number of specialists in FCM increased over time, recent evidence suggests that FCM is still the last option among physicians (Fontenelle et al., 2020^[20]). As earlier mentioned, in 2018, only 5 486 medical doctors had a specialty training in FCM, representing only 1.4% of all specialists in Brazil (Scheffer et al., 2018^[1]).

Most doctors working in PHC in Brazil do not have a specialist qualification. This is unlike many other OECD countries where medical doctors are required to have a specialty training to be allowed to practise as a PHC doctor. In Canada, for example, physicians must have additional specialist training in family medicine to work in PHC (Ponka et al., 2019^[23]). In the United Kingdom, participation in a three-year course is compulsory to practise family medicine. Turkey introduced specialised family medicine training as part of the *Health Transformation Programme* launched in 2005. The reforms sought to reinvigorate the speciality of family medicine, which was first defined in 1983 but failed to embed itself extensively in PHC provision. The reforms defined the Family Medicine core team as comprising a family physician, nurses and professional assistants, to whom a list of named patients was assigned. Family physicians across Turkey are required to deliver a defined set of services, to work to a standard set of norms and are paid according to national terms and conditions. Significant investment has been made to improve working conditions, with more generous salaries than before the reform. In 2018, Turkey’s PHC/generalist doctors now comprises 32% of all doctors, above the OECD average of 23% (OECD, 2019^[22]).

Box 5.3. The Brazilian Society of Family and Community Medicine (SBMGC) and the Brazilian Society for Family Medicine (SOBRAMFA)

Founded in 1981, the Brazilian Society of Family and Community Medicine aims to promote the principles and values of Family and Community Medicine as an academic discipline, as well as to promote patient-centred care in the clinical practice and the medical education.

The Brazilian Society of Family Medicine is organised around three priorities:

- Prepare future professionals based on patient-centred approach
- Integrate medical students to health services and connect them to the real needs of the population
- Train teachers, educators, and managers to promote academic incorporation of the specialty.

The Brazilian Society for Family Medicine (SOBRAMFA), founded in 1992, also organises several activities to spread the family medicine value among medical students at congresses, academic meetings, family medicine seminars and international meetings. It organises Mini-Fellowship in Family Medicine programme, where students experience family medicine practice by managing patients under the supervision of the SOBRAMFA faculty and residents in a range of practice setting. For example, the SOBRAMFA develops guidelines, promote research and provide training and continuous medical education (such as the Miles Programme).

The internship for medical students in undergraduate programme, and the specialty programme focus on follow-up care for chronic patients, health prevention, treatment of multiple comorbidities. It also focuses on geriatric care (care for the elderly who live in communities), palliative care (learning how to care for patients without therapeutic possibility, discuss approach to death, deal with disabilities, and symptom treatment), understanding patient's pathway, learning team work and improving communication, and work in palliative care team.

Source: Janaudis et al. (2009^[24]) Academic Family Medicine in Brazil ; SBMFC (Janaudis et al., 2009^[24]; SBMFC, 2021^[25]).

In the area of nursing, Brazil has several degrees ranging from bachelor degree, master degree to doctoral professional degrees (Cassiani and Silva, 2019^[6]; Neto et al., 2018^[26]). It is reported that around 50 000 nurses are trained every year (Cassiani and Silva, 2019^[6]). An evaluation of nursing training found that the training had a hospital-centric model, with little interdisciplinary integration (Magnago et al., 2017^[27]). This is not appropriate to strengthen people-centred primary health care.

CHWs are most often selected by local health committees and have to be literate. The training of community health workers is conducted in regional health schools operated by the national Ministry of Health using curriculum approved by the Ministry of Education (CHW Central, 2018^[28]). The curriculum consists of 8 weeks of training from local nurses, followed by 4 weeks of supervised field training. These include training on home visits and on how to conduct a family census, but also on specific priority health care interventions. CHWs are not trained to address health needs, but rather they are trained to recognise health needs and encourage communities to seek care (CHW Central, 2018^[28]; Grossman-Kahn et al., 2018^[7]). CHWs also receive monthly and quarterly ongoing training.

5.3.3. National evaluation tools and voluntary accreditation facilitate evaluations of the quality of medical education in Brazil

Each medical school in Brazil decides its own curriculum and educational objectives, and implements its own summative and formative assessments of students based on skills, abilities, and curriculum content.

To promote changes in the medical curriculum toward PHC, the Ministry of Education has introduced the Program for the Encouragement of Curricular Changes in Medical Courses (Promed), the National Program for the Reorientation of Health Professional Education (Pró-Saúde), and the National Guidelines for the Curriculum of Health Professional Education. The National Curricular Guidelines of Medical Courses introduced in 2001, then updated in 2014 to support the More Doctor Programme, intends to ensure homogeneity between medical schools across the country (Ferreira et al., 2019^[29]).

The curriculum's competencies target three axes: health care, health management and health education, with a focus on collective health and comprehensive care. The updated guidelines aim at moving away from hospital practice, increasing the inclusion of theoretical and practical activities related to collective health and PHC, and improving uptake of activities related to social determinants of health. The overarching objectives encompass (1) promoting patient-centred care, (2) reliance on inter-professional teamwork, and (3) developing shared individual therapeutic plan with the patient (Ferreira et al., 2019^[29]). With these programmes, undergraduate students are supposed to be exposed to a more community-based curriculum, although it is difficult to appreciate the impact of these initiatives on the skills of the students (Scheffer and Dal Poz, 2015^[16]).

National tools have also been set-up to evaluate medical schools as a result of the sharp increase in medical schools from 1990. The National System for the Evaluation of Higher Education (SINAES, *Sistema Nacional de Avaliação do Ensino Superior*) is responsible for evaluation system to renew, supervise, accredit and reaccredit medical schools (INEP, 2015^[30]). The evaluation system consists of three main components:

- Institutional evaluation: performed by self-evaluation and external evaluation by institutional evaluators accredited by the INEP;
- Evaluation of the course and curriculum by teachers, co-ordinators, students, and external evaluators;
- Evaluation of the students entering and finishing the course: through the National Exam of Student Performance (ENADE, Exame Nacional de Desempenho dos Estudantes), which is performed by each schools (Ministerio da Educação, 2021^[31]). ENADE covers seven areas of knowledge including general medicine, paediatrics, gynaecology and obstetrics surgery, public health, medical ethics and basic sciences. The test results are used to identify areas of good performance and weaknesses in all medical schools. The ENADE gives an indication of the development of competences and skills foreseen in the National Curricular Guidelines of Medical Courses.

The Sistema de Acreditação de Escolas Médicas (Accreditation System of Medical Schools, SAEM) has also been developed by the Conselho Federal de Medicina (CFM, Brazilian Federal Council of Medicine) in 2016 to reaffirm its commitment toward high quality standards for medical education (SAEME, 2021^[32]). The accreditation is voluntary and complements the government evaluation procedure. The SAEM is based on the WHO and the WFME Guideline for Accreditation of Basic Medical Education. The accreditation process consists of the completion of online questionnaire covering five standards (Educational Management, Educational Programme, Academic Staff/Faculty, Students, and Educational Resources), data analysis by an evaluation committee; and a visit to the medical school by a team composed of four evaluators, one of them being a student. After this process, the evaluation committee issues a report to the educational institution with the result, which can be classified into three categories: Accredited, Accredited with recommendations of improvement, and Non-Accredited.

National tools and voluntary accreditation allowing to evaluate the quality of medical education is paramount to ensure consistency in the skills and competences of medical doctors across the country, but also to make sure the supply of medical schools and their postgraduate activities reflect the evolving Brazilian health needs. However, the CFM recently noted that, in spite of government initiatives, there are still inadequate academic programmes, inadequate internship offers and that gaps in medical education remain between medical schools (AML, 2019^[33]). In addition, comparing the performance of public and

private medical schools, evidence suggests that public medical schools consistently performed better than the private undergraduate medical courses about the ENADE (Scheffer and Dal Poz, 2015^[16]). Reinforcing mechanisms to ensure high quality of medical education is thus warranted in Brazil.

5.4. Improving the distribution and supply of PHC doctors in Brazil

To address the shortage of PHC doctors and the uneven geographic distribution of doctors, Brazil implemented several federal programmes including the More Doctors Programme. These efforts contributed to improvements in access to and quality of medical training in Brazil, but geographic disparities in access to medical training persist and unfilled PHC positions remains the norm.

There are three main policy levers to help address the shortages of medical doctors and improving its distribution:

- Training more rural doctors;
- Providing financial and non-financial incentives, while implementing regulations to restrict the freedom of medical doctors (the “stick and carrot approach”);
- Promoting innovations in health service delivery, through task-shifting and the use of telemedicine.

They are not mutually exclusive, and may not offer the panacea when implemented in isolation. Previous studies show that these policy levers may have a greater impact if they are used in combination (OECD, 2016^[34]).

5.4.1. Large-scale PHC programmes to address maldistribution of PHC physicians

The More Doctors Programme partly addressed the imbalances in the distribution of health workers

Established in 2013, the More Doctors Programme (MDP) is a large-scale health system intervention that aims to strengthen the provision of PHC services in underserved communities, within the framework of the Family Health Strategy (see Chapter 2). The MDP was designed by the Ministry of Health as a supply side intervention with three prongs that complement each other, with a particular emphasis on strengthening PHC provision in underserved communities:

- Transfer of funds to municipalities to strengthen PHC infrastructure;
- Improving access to and quality of medical school training; and
- Recruitment of Brazilian and foreign physicians to work within FHTs.

At the outset, the first two components were envisioned as long-term investments in the overall organisation of the Family Health Strategy. Broadly, the first component aimed at improving the availability and quality of work environments in which the FHT provide care (Santos et al., 2017^[35]). The second component focused on enhancing access to and quality of medical training, particularly in communities that have been lagging in attracting and retaining health workers (Figueiredo et al., 2021^[36]). In comparison, the third component was initially envisaged as an emergency provision to fill the immediate gaps in PHC physician availability through a large-scale physician recruitment initiative, particularly targeted at municipalities that had difficulty recruiting and retaining doctors (Santos et al., 2017^[35]).

The MDP came on the heels of previous nationwide initiatives that attempted to address the imbalances in the distribution of health workers. Prior to the MDP, Brazil introduced several programmes to improve access to medical training in underserved communities, including the 2001 Programme of the Interiorisation for Health Work (*Programa de Interiorização do Trabalho em Saúde*) and the 2011 Programme for Valuing PHC Professionals (*Programa de Valorização dos Profissionais da Atenção –*

PROVAB) (Figueiredo et al., 2021^[36]). While these initiatives did not eliminate the underlying drivers of the geographic imbalances in distribution of health workers, they provided valuable experiences and the know-how that informed the design and implementation of the MDP.

The new stream of funding led to a laudable expansion of the Family Health Strategy infrastructure

The first component involved creating a new stream of funding to strengthen the Family Health Strategy physical infrastructure. Municipalities could use the allocated MDP funding to increase the availability and the quality of the existing Family Health Strategy infrastructure on the ground. To receive funding, each municipality was required to submit an application to the MOH for review. Upon approval of their application, municipalities could choose to either refurbish the existing PHC clinics and purchase basic equipment (e.g. autoclaves) or build new facilities (Santos et al., 2017^[35]).

Relatively little evidence exists on the impact of this component on the provision of PHC services. Though limited, evidence suggests that the MDP successfully channelled additional resources for strengthening the PHC infrastructure, with the MDP resources leading to the construction of 3 496 new basic health units and refurbishment of 3 417 units by 2015 (Santos et al., 2017^[35]). According to Biernath (2020), more than 26 000 PHC facilities were built or renovated throughout Brazil following the implementation of the MDP (Biernath, 2020^[37]). The administration of this component was recently halted following a federal funding freeze (Hone et al., 2020^[38]).

The MDP contributed to improvements in access to and quality of medical training in Brazil, but geographic disparities persist

Despite previous attempts to improve equitable access to medical training, the geographic distribution of medical schools prior to the MDP were marked by notable disparities. For instance, in 2013, nearly 41% of all medical schools were located in the Southeast Region, compared to only 8.2% in the North and 9.3% in the Central-West regions (Santos et al., 2017^[35]). Compounding this challenge, the undergraduate medical curriculum did not adequately emphasise the diverse health needs of the Brazilian population (e.g. population groups residing in rural and remote areas) and PHC competencies in line with the priorities of the SUS (Carvalho, Marques and Silva, 2016^[39]).

In recognition, the second component consisted of a set of initiatives that aim to enhance access to and quality of medical education, with an explicit emphasis on alleviating geographic disparities in medical training. The MDP introduced multiple changes in medical education at the same time. First, the education component of the MDP involved the development of new frameworks to regulate the location of new public and private medical schools. Next, the MDP aimed at expanding the number of positions in medical undergraduate institutions by adding nearly 11 500 new positions by 2017 (Santos et al., 2017^[35]). Finally, the Programme led to the updating of the national undergraduate curriculum for medical training with more emphasis on PHC provision in line with the SUS priorities.

Early evidence suggests that the MDP may have contributed to improvements in access to medical training. Evaluating the impact of MDP on medical education in Brazil is complicated by the length of time required before the potential impacts on health system performance can become discernible. Yet, data obtained from the Higher Education Census suggests that the MDP achieved its medium-term objective of expanding the number of new positions by 2017. Mirroring this, an estimated 92% of the new medical undergraduate vacancies created from 2010-18 took place after the MDP was launched in 2013 (Figueiredo et al., 2021^[36]).

Despite these improvements, geographic disparities in access to medical training persists. Today, the North and Northeast regions continue to offer the lowest levels of per capita medical undergraduate vacancies in the country (Figueiredo et al., 2021^[36]).

The recruitment of PHC physicians contributed to enhancing health system performance, but challenges in the implementation of MDP diluted beneficial impacts

The third component involves the roll out of one of the world's largest PHC physician recruitment programme. At its height, the MDP recruited more than 16 000 physicians both from within Brazil and abroad to work exclusively within the FHT. The majority of the MDP physicians were recruits from across 85 countries, mostly from Cuba (Santos et al., 2017^[35]). All MDP recruits were selected from physicians who had prior experience in practicing family medicine, but without recertification requirement.

All Brazilian municipalities are eligible to request physicians from the MDP. To receive MDP physicians, municipalities are required to submit an online application to the Ministry of Health for review and approval. The Ministry of Health has the discretion to decide on the number of MDP physicians that will be deployed in the applying municipality. To inform the decisions over MDP physician allocations, the Ministry of Health developed a set of criteria aiming to identify underserved communities across Brazil in congruence with the aims and objectives of the Programme (Box 5.4).

Provider contracting arrangements for MDP physicians differ from contracts typically used by municipalities to hire non-MDP physicians. In general, the FHP physicians are hired directly by municipal governments, with little standardisation of contractual terms across physicians or municipalities. In comparison, the MDP physicians are hired directly by the Ministry of Health through contracts that can be renewed in every three years. Municipal governments are responsible for covering costs associated with transportation, lodging and food. This contracting structure was embedded in the MDP to reduce potential financial barriers that may disincentivise municipalities from joining the Programme. Importantly, municipalities are not allowed to substitute Brazilian physicians that are already practicing within the FHT in their communities with MDP physicians. However, no mechanism has been put in place to monitor any potential physician substitution.

It is not mandatory for the MDP physicians to take the national competency exam that is required for other physicians to practise medicine in Brazil. Instead, the MDP physicians are required to complete a mandatory three-week training course provided by the Brazilian health authorities before they could be deployed in municipalities. In addition, all MDP physicians are required to attend brief training sessions organised by Brazilian authorities in regular intervals. Online-learning courses are also available to the MDP physicians. Once deployed in municipalities, the MDP physicians are assigned to advisors, who are selected from experienced Brazilian health care professionals.

To date, the MDP has been associated with improvements in physician availability, though there is evidence of substitution of Brazilian physicians with the MDP recruits. Hone and colleagues (2020) suggested that the MDP was associated with a 12.2% increase in the number of PHC physicians by its fifth year of implementation, with more pronounced effects among municipalities that had lower levels of physician availability at the start of the Programme (Hone et al., 2020^[38]). Despite this, evidence suggests that the MDP also had unintended consequences for human resources for health, with the MDP leading to the substitution of Brazilian physicians who were already working in communities. In their analysis, Hone and colleagues quantified that, in the absence of any substitution of Brazilian physicians with MDP recruits, the MDP could have led to a 30% increase in PHC physician availability, more than double that of the current improvements in physician availability (Hone et al., 2020^[38]). This is despite the design feature of the MDP that explicitly prohibited any replacement of Brazilian physicians with MDP physicians.

The impact of MDP on adult and child mortality remains mixed. Hone and colleagues found that the MDP was associated with a modest 1.4% reduction in adult mortality amenable to health care services (Hone et al., 2020^[38]). Earlier studies suggested that the MDP fell short of reducing maternal and child mortality (Carrillo and Feres, 2019^[45]; Mattos and Mazetto, 2019^[46]), despite an attributable rise in the number of FHT consultations attended by physicians (Mattos and Mazetto, 2019^[46]). One potential explanation relates to the significant bottlenecks in access to the secondary and tertiary care, suggesting that further progress in maternal and child health may require further improvements in co-ordination of care between different

levels of care (the Brazilian Network for the Surveillance of Severe Maternal Morbidity study group, 2014^[47]).

Box 5.4. Defining underserved municipalities within the framework of MDP

The MOH ranks municipalities based on a set of explicitly defined criteria to inform the distribution of MDP physicians (Hone et al., 2020^[38]). Criteria used for the MDP are similar to those used in PROVAB, with the two initiatives sharing similar, readily available data sources at the community level (e.g. municipality-level or census tracks within municipalities). In the first five years of the MDP, the Ministry of Health revised the prioritisation criteria twice, but in each iteration, the selected criteria aimed at capturing the socio-economic, health systems and geographic determinants that may reveal the local needs in each community. These indicators included: the proportion of population living in extreme poverty, population size, PHC infrastructure and health worker supply, as well as geographic location (Özçelik et al., 2021^[40]).

A substantial fraction of Brazilian municipalities enrolled in the MDP in a relatively short time span, though the rollout of the Program has not been without challenges. The share of municipalities that received at least one MDP physician expanded from about 20% in 2013 to 69% in 2017 (Özçelik et al., 2021^[40]). Over time, participation improved among municipalities that were considered priority for the purposes of the MDP. By 2017, about 76% of priority municipalities had at least one MDP physician. Priority municipalities that did not enrol in the MDP had smaller population density and higher rates of extreme poverty compared to priority municipalities that participated in the MDP. Importantly, this finding is in congruence with earlier evidence suggesting that less densely populated communities with higher levels of poverty also lagged in the expansion of the Family Health Strategy prior to the MDP (Andrade et al., 2018^[41]).

It might be warranted to improve the community targeting methods used by the MDP to maximise the potential benefits of the Programme. As experienced by other OECD countries (the United States, Australia, and Canada), the community targeting method has large implications for human resource planning, and this might not be enough recognised in Brazil. In Australia, for example, communities with a shortage of medical practitioners are designated as the *Distribution Priority Areas* on the basis of their demographic and socio-economic status, population size and measures of geographic remoteness. Foreign-trained general practitioners are required to work at least 10 years in communities with DPA designation. In Canada, criteria used to define underserved communities consider physician to population ratios, demographic and socio-economic status and geographic accessibility. In The United States, the Health Professional Shortage Area designation is determined based on a composite score derived as a combination of health care service provider density, level of poverty, infant mortality rate and the age structure of each community.

Source: Özçelik et al. (2021^[40]), Assessing the performance of beneficiary targeting in Brazil's More Doctors Programme, <https://doi.org/10.1093/heapol/czaa137>; Andrade et al. (2018^[41]), Transition to universal primary health care coverage in Brazil, <https://doi.org/10.1371/journal.pone.0201723>; Hone et al. (2020^[38]), Impact of the Programa Mais médicos (more doctors Programme) on primary care doctor supply and amenable mortality: quasi-experimental study of 5 565 Brazilian municipalities, <https://doi.org/10.1186/s12913-020-05716-2>. And information taken from Canada (Ministry of Health, 2018^[42]), Australia (Australian Government, 2021^[43]), and the United States (HRSA, 2021^[44]).

Most evidence suggests that the MDP led to improvements in access to care. One study that evaluated the impact of the MDP in its first two years found that the Programme was associated with increases in appointments, consultations, medical referrals and home visits by 5.9%, 9.4%, 12.3%, and 29.7%, respectively (Mattos and Mazetto, 2019^[46]). This study also found that the MDP led to a 4.6% reduction in

general hospitalisations. Similarly, the MDP has been shown to improve patient satisfaction (Comes et al., 2016^[48]).

The MDP has also been linked with improvements in the quality of care, as measured by reductions in hospitalisations considered to be preventable by timely and adequate access to health care services. Earlier assessments concluded that the MDP resulted in a decrease of about 23 000 hospitalisations due to preventable causes in its first three years, corresponding to about USD 6 185 019.85 in savings from treatment costs (Fontes, Conceição and Jacinto, 2018^[49]). The beneficial impacts of the MDP have been shown to accumulate and grow over time. Evidence also shows that the MDP led to reductions in adult hospitalisations primarily due to infectious gastroenteritis, bacterial pneumonias, asthma, kidney and urinary infections, and pelvic inflammatory disease (Maffioli et al., 2019^[50]), though it fell short of reducing hospitalisations due other preventable conditions like hypertension (Özçelik et al., 2020^[51]).

Evidence on the contributions of MDP for narrowing geographic disparities in population health remains mixed, suggesting that further improvements in community targeting methods may be needed to reap the potential benefits. Hone and colleagues (2020) concluded that the MDP fell short of yielding greater health gains in priority municipalities (Hone et al., 2020^[38]). In concordance, another recent study also showed that there has been no discernible differences in hospitalisations due to hypertension between urban and rural municipalities as defined by the Programme (Özçelik et al., 2020^[51]). This study also showed that the MDP was linked with reductions in hospitalisations due to cerebrovascular diseases only in urban municipalities. Both studies suggested that further improvements in the community targeting methods used by the MDP may be considered to maximise the potential benefits of the Program.

The new programme Médicos pelo Brasil will gradually replace the MDP

A new programme called Doctors for Brazil (Médicos pelo Brasil, law No. 13.958 of 18 December 2019) was initiated by the new federal government administration. Its purpose is to increase the provision of medical services in places of difficult delivery or of high vulnerability, along with promoting the training of doctors specialising in family and community medicine.

The *Médicos pelo Brasil* programme is expected to gradually replace the *Mais Médicos* programme, in particular by hiring Brazilian trained doctors who have already shown to be scarce or not wanting to go to isolated or underserved areas.

Médicos pelo Brasil and *Mais Médicos* will operate in parallel to guarantee continuity in the supply of health care services. This type of changes represent a governance challenge for the Brazilian health system, which should develop a strategic vision to preserve and amend policies or programmes that have shown to improve outcomes. The MDP should inform the design and implementation of the *Médicos pelo Brasil* programme.

Based on prioritisation criteria, the new programme will expand the number of PHC doctors to 7 500 in the poorest areas of the country, with 60% of the professionals to be hired in the North and Northeast regions. In total, 4 823 municipalities are expected to be part of the programme. To maximise the potential benefits of the new programme, it will be critical to make the prioritisation criteria clear and transparent.

Médicos pelo Brasil has a stronger focus on quality than the MDP, and positive steps have been taken so far:

- Medical licenses have to be recognised for all medical doctors who apply to the programme, either through the registration process for Brazilian students or the revalidation process for foreign-trained medical doctors. The recognition process is the responsibility of the Medical Council and Regional Councils of Medicine.
- The *Médicos pelo Brasil* programme offers two-year specialty training in Family and Community Medicine so that all contracted doctors become specialists in PHC. At the end of the training,

specialists in PHC will have to present a “*Trabalho de Conclusão de Curso*”, an analysis of the local health needs of the catchment area where medical doctors work so that improvement and evaluation strategies are being implemented.

- PHC doctors are expected to remain in the Family Health Units where they did the specialty training, and receive a 2-year training grant, which varies according to the municipalities’ level of vulnerability. The grant is BRL 12 000 net monthly, and an additional BRL 3 000 bonus for remote locations (those classified as rural and intermediate) or an additional BRL 6 000 for indigenous districts, as well as riverside and river locations. Once qualify as specialist in Family and Community Medicine, they are hired under the Consolidation of Labor Law (CLT) which supposes better working conditions and security at work compared to the *Mais Médicos* programme.

With regards to the tutorial programme, only specialists in family and community medicine or in internal medicine can be selected. Tutors are also hired under the CLT. They will be responsible for serving the population registered to the FHT, and for supervision of other medical doctors who are enrolled in the specialisation training. There are four salary scales, with progression every three years: The first salary level can reach up to BRL 21 000 and, gradually, up to BRL 31 000, considering performance bonus and additional grants depending on the level of vulnerability.

All these changes are promising, representing improvements compared to the Mais Medicos Programme and real opportunities to increase the attractiveness of the PHC specialty (Table 5.1). Strategic thinking toward the community targeting methods is required to make sure that eligible municipalities are effectively enrolled to the new programme.

Table 5.1. Médicos pelo Brasil programme has a greater focus on quality

Mais Medico	<i>Médicos pelo Brasil</i>
Scholarship without employment link and for a determined time	Contracting by the CLT with the right to FGTS holidays
Does not require any Brazilian diploma or revalidation	Requirement to have a recognised Brazilian licence or to undertake revalidation of foreign diploma
No requirement to have a formal specialisation degree in PHC	Requirement to pass the specialty training in Family and Community Medicine.
Foreign Medical doctors are not allowed to bring the family to live in Brazil	Foreign Medical doctors are allowed to bring the family to live in Brazil

Source: Brazilian Ministry of Health.

The next two sections present policy responses that could help tackle the uneven distribution of PHC physicians in Brazil.

5.4.2. Changing regulations can help address shortages of medical doctors in some remote areas

Training more rural doctors to boost their numbers

As the MDP, the new *Médicos pelo Brasil* programme sets very promising objectives to improve the supply and the distribution of PHC doctor across the country. As part of the new programme, it will be important to train rural PHC doctors to improve retention over the longer term. International evidence confirm that medical education in rural location led to increases in the number of medical graduates that will work in rural place over the long term (McGirr, Barnard and Cheek, 2019^[52]). In congruence, experiences from OECD countries also highlight the importance of having medical studies for PHC professionals be located in local regions such that students are close to their residencies and are incentivised to stay upon successfully completing their training (OECD, 2016^[34]).

Box 5.5. The Rural Clinical School Programme in Australia

The Rural Clinical Schools (RCS) Programme was launched in 2000 to enable medical students to undertake the majority of their clinical training in regional areas.

The RCS aim to deliver significant components of the medical curriculum in a rural environment, with students undertaking a year or more of their medical training in a rural location. It entails the following targets (McGirr, Barnard and Cheek, 2019^[52]):

- 25% of all medical students to have at least one year of rural clinical training by graduation
- 25% of government-supported medical students to be recruited from a rural background
- All government-supported medical students to have at least 4 weeks of rural placement.

The programme is structured around four broad objectives (Australian Department of Health, 2008^[53]):

- Encourage medical students (and medical professionals) to take up a career in rural practice
- Encourage rural health professionals to take up academic positions
- Improve the range of rural health care services in rural communities across Australia
- Strengthen the health workforce in rural communities across Australia.

As of 2018, there were currently 18 RCSs in Australia (McGirr, Barnard and Cheek, 2019^[52]). The Rural Clinical Schools Programme has allowed to develop teaching and learning facilities, and student accommodation in dozens of rural and regional locations across Australia. There are positive evidence showing that RCS programme improve retention and recruitment, but also improving the range of quality of health care services in rural area. With this regards, particular specialties that were not available in the community have become available because of a successful appointment by the programme.

Source: McGirr, Barnard and Cheek (2019^[52]), The Australian Rural Clinical School (RCS) programme supports rural medical workforce, <https://doi.org/10.22605/RRH4971>; Australian Department of Health (Australian Department of Health, 2008^[53]), Rural Clinical Schools Programme.

Prior experiences from OECD countries, such as Australia, Canada, England or Norway, offer many opportunities for Brazil. For instance, in Australia, the Rural Clinical School (RCS) Programme was launched in 2000 to enable students to undertaken their training in rural areas. The RCS Programme supports 18 rural clinical schools, where it is required that 25% of students enrolled in the medical course have a rural background (McGirr, Barnard and Cheek, 2019^[52]). Recent evaluations showed that the RCS Programme was effective in increasing both the recruitment and retention of medical practitioners in rural areas over the long term (McGirr, Barnard and Cheek, 2019^[52]). Results show that students who have participated in RCS Programme were 1.5 times as likely to continue practicing medicine in regional and rural locations five years after graduating from medical school (Box 5.5). Other OECD countries established medical programmes in rural localities, such as Canada with the Northern Ontario School of Medicine, England with the Exeter and Plymouth medical school, or Norway with the Tromsø's medical school.

Implementing a “stick and carrot” approach

It is important to implement a “stick and carrot” approach to attract and retain PHC doctors in rural areas. This approach consists of combining financial and non-financial incentives for medical doctors with regulations which restrict the freedom of doctors.

Offering financial incentives to compensate for some of the disadvantages in less attractive regions is an important policy option already implemented in many other OECD countries such as in Chile, Germany, Canada or France. Such financial incentives compensate for longer working hours, a less advantageous “business environment” for doctors in underserved areas. They can be either one-off or wage-related financial incentives that are directly directed to the PHC doctors:

- First, there are **non-wage related payments**, which are used to encourage physicians to move to or stay in underserved areas. It is a one-off payment to help facilitate a location choice, and which can be linked with a return of service obligation. Such policies exist in some Canadian provinces. In Ontario, the Northern and Rural Recruitment and Retention Initiative offers grants of between CAD 80 000 and CAD 117 600 for a practice opening in a rural area paid over a four-year period (Ontario Ministry of Health, 2021^[54]). The applicant must commit to establish a full-time practice in a community in Northern and very rural areas, and in which they have never previously practiced full-time. The higher the rural index (the more Northern and rural the area), the higher the value of available grant. To encourage both recruitment and retention, the proportion of the grant paid out will be highest in year 1 and year 4. The payment schedule is as follows: Year 1: 40%, Year 2: 15%, Year 3: 15%, Year 4: 30%. The province of New Brunswick also offers recruitment incentives to Medical Residents and newly recruited family physicians looking to establish a full-time community-based practice in designated fields. Family Medicine New Brunswick is a new programme that aims to improve patient access, increase collaboration between physicians using technology, and create a better work-life balance for physicians. For the first year, a new physician joining the programme will receive a guaranteed minimum remuneration of CAD 175 000 (New Brunswick Canada, 2021^[55]). Similar policies exist in Germany, where states (Länder) offer financial incentives for GPs opening their practice for the first time. In some states, the one-time payment comes with a return-of-service obligation of five to ten years (OECD, 2016^[34]).
- Second, there are **wage-related financial incentives**, where physicians in underserved areas receive financial incentives linked to income. These payments aim to compensate rural GPs for a smaller number of patients, longer working hours or challenging working conditions. In the Canadian province of Ontario, the Northern Physician Retention Initiative enables eligible physicians (general practitioners and specialists) in Northern Ontario to receive the equivalent of CAD 7 267.46 retention incentive paid at the end of each fiscal year in which they continue to practise full-time in Northern Ontario (Ontario Ministry of Health, 2021^[56]). In France, the Contrat d'Aide à l'Installation des Médecins offers a one-off payment of between EUR 50 000 and 31 250 to help physicians set-up a practice in an underserved region. The payment schedule is 50% upon signature of the CAIM, and 50% one year later (Assurance Maladie, 2020^[57]).
- Third, countries can **combine financial and non-financial incentives** to attract and retain physicians in underserved communities. For instance, Chile provides a promising example, where several incentives are combined to attract physicians to work in rural PHC hospital and health centres for a minimum of three years to a maximum of six years (Pena et al., 2010^[58]). The programme, called the Chilean Rural Practitioner Programme, entails 1) monetary compensation (direct and indirect financial incentives such as additional compensation depending on the level of isolation, performance bonus, installation and transport costs), 2) education incentives (four week clerkship with physicians, 3) management, environment and social support interventions (paid continuous medical education, additional days for personal reason, and additional week off) and 4) external incentives (better equipment, better connectivity, and internet and mobile phone access) (Pena et al., 2010^[58]). The programme has been found successful: the number of applicants exceeded the number of available positions, high retention rates, and high satisfaction rates from applicants (Pena et al., 2010^[58]). While dropouts from the programme are exceptional, around 60% of participants stay for the maximum period (six years).

As part of the *Médicos pelo Brasil* programme, physicians in underserved areas will receive financial incentives of BRL 3 000 when working in “rural and intermediate” areas, and of BRL 6 000 when working in indigenous districts, as well as riverside and river locations. As done in other OECD countries, it could be important to link financial incentives with a return of service obligation of between two and four years. Combining financial and non-financial incentives, and implementing some regulations to restrict the freedom of doctors, as experienced by OECD countries, could also be a good solution to recruit and retain doctors in underserved areas. The federal government will need to ensure that the various types of financial incentives are received by the medical doctors, and not distributed at the discretion of the municipalities.

Restricting the choice of location

A more stringent regulatory measure to redress the geographical imbalance of doctors would be to restrict the choice of practice location for PHC doctors. Brazil could, for example, learn from Turkey, Germany, Norway or some provinces in Canada (New Brunswick, Quebec) where interventions to restrict the choice of location of newly medical doctors have been established. At any rate, such regulatory approach requires careful and coherent workforce planning.

In Germany, self-employed doctors are not allowed to set up their practice and serve public patients in a catchment area that is considered ‘overserved’ based on a patient-to-doctor ratio. Physicians need to obtain a practice permit to be reimbursed by the statutory health insurance (KBV, 2021^[59]). The number of these permits is controlled by the National Association of Statutory Health Insurance Physicians through its 17 state associations. It is mandated by the government to guarantee medical service coverage of the population based on a quota agreed within the self-administration of the German health care system. The service coverage is measured based on the ratio between physicians and inhabitants in each of the 395 planning regions. For GPs, 100% coverage is achieved when the ratio of GP to inhabitant reaches 1:1 617. If the coverage of a region exceed 110%, no further permits are issued (OECD, 2016^[34]). In Canada, in the province of Alberta, the Bill 21 proposed that after April 2022 new doctors in the province will be required to obtain practice permits from the government upon completion of residency training. These permits will restrict new doctors to work only in certain regions. Turkey restricts service location choice of new medical graduates who want to work in the public sector. Recent medical schools graduates are required to work for the states for a period of 300 to 500 days after graduation. After this service period however, physicians wants to secure employment in urban areas (Phcpi, 2018^[60]).

Such regulatory approaches can be controversial with fears from professional association that designating where doctors should work impede recruitment. In this case, it is important to provide strong financial compensation to medical doctors.

5.4.3. Changing service delivery can help reduce geographical inequalities in access to care

There are several interventions on the supply side to promote the use of innovative health service delivery to provide adequate level of access to PHC in areas with fewer physicians (e.g. through telemedicine or telehealth) or shifting some health service provision from physicians to other health care providers. The strategy is not to train more doctors or to rely on more doctors, but rather to rely on smart information and communication technology and to maximise use of human resources already living in areas where there are shortages of medical doctors. This could constitute a good complement for usual care provided by medical doctors.

Expanding the role of nurses or community health workers

Unlike many other OECD countries, Brazil has not yet experimented with changing scope of practice and task shifting among health workers. More strategic use of the existing rural health workers other than

physicians who are already residing in areas where there are physician shortages will be crucial to meeting health care needs in more remote areas.

Core PHC functions, including comprehensiveness, continuity and co-ordination, can be discharged to health workers other than PHC doctors. In recognition, many OECD countries are moving to change the scope of practice of health professionals as a means of coping with health workforce shortage, which typically involves developing new roles for nurse practitioners and community health workers (OECD, 2020^[61]).

International evaluations of task shifting provide conclusive results. Nurse practitioners with proper education and training generally show that they can improve access to services, reduce waiting times and deliver the same quality of care as doctors for services such as routine follow-up of patients with chronic conditions (Maier, Aiken and Busse, 2017^[62]). Recent systematic review found that PHC teams that enable nurses to play a more advanced role in service provision (e.g. providing patient education, co-ordination, prevention advice or drug prescription) achieve better patient outcomes and greater patient satisfaction, while reducing hospitalisation (Matthys, Remmen and Van Bogaert, 2017^[63]). Moreover, another recent systematic review showed that scaling up reliance on nurse practitioners yielded greater access to care among rural and underserved populations without deteriorating the quality of care (Yang et al., 2020^[64]). In low and middle income countries, the implementation of task-shifting has been a key lever to fill the gap in PHC provision due to a lack of PHC doctors (Afobali et al., 2019^[65]).

Among OECD members, some countries such as Australia, and more recently France have sought to improve access to PHC by expanding advanced education programmes for nurses. In Australia, remote area nurses are recognised as the backbone of rural and remote PHC. They can work as part of a small team or work independently. With a rotation system, they are available 24 hours a day, seven days a week. They are able to work across diverse care settings, including acute, emergency, aged, palliative, mental health, family and community health care areas (OECD, 2020^[61]). Some Jurisdictions in Australia, including Queensland, and Victoria, have also implemented the Rural and Isolated Practice Endorsed Registered Nurse scheme. The scheme permits approved nurses to provide a limited range of medicines, where there is little or no access to PHC. In France, the National Health Strategy 2018-22 promotes more advanced roles for nurses to improve access to care, particularly in areas where there are relatively few doctors (OECD, 2020^[61]). The position of advanced practice nurses was legally created in 2018. These nurses provide support in the care of chronically ill patients and those with complex morbidities, working in co-operation with GPs and specialists in PHC teams and other health, and long-term care settings. Their tasks include prevention and screening activities, prescription of complementary exams and renewal or adjustment of medical prescriptions.

Brazil has amongst the most extensive education networks for nurse training in the LAC region. According to the Department of Management of Health Education in Brazil, the number of nursing schools amount to 790 in 2020, with several specialisation programmes being offered. In 2019, there were around 24 master's degrees programme in nursing, constituting powerful training spaces for developing advanced practice nurses (Cassiani and Silva, 2019^[6]; Neto et al., 2018^[26]).

Brazil can capitalise on its existing nurse training network to expand and recognise the role of advanced practice nurses, which in turn, can help alleviate the persistent workforce shortages in PHC. Advanced practice nurses in Brazil could contribute to the management of care of patients with mild acute diseases and chronic disorders, complementing the already practicing PHC doctors. Alternatively, they can be deployed in remote and underserved areas to help enhance access to qualified health professionals. Such a policy would require a thorough analysis on current nursing curriculums to further invest on core competencies including team practice, care co-ordination between and across health sectors, clinical and professional leadership, quality and safety management among other (Cassiani et al., 2018^[66]).

Beyond expanding the role of nurse practitioners, it is also warranted to make the most of community health workers within the Brazilian PHC team. Community health workers already play a crucial role in

PHC provision in Brazil. They are trusted members of the communities in which they work and have valuable knowledge of local health needs. In Brazil, there is scope to expand further their role toward more preventive activities and care management. In Alaska (United States), CHWs are the first point of contact with the health care system for the population living in very remote villages (Box 5.6). There, they provide preventive treatment, chronic care and emergency visits.

In a systematic studies review (Hartzler et al., 2018^[67]), community health workers are found to be able to perform three main functions:

- providing clinical services, such as assessment of vital signs, lifestyle advice, and routine examinations aided by remote communication with physicians;
- linking patients with community-based services, such as referrals for transportation or food assistance; and
- providing health education and coaching, to help patients achieve health goals and increase self-efficacy.

With serious workforce gap in Brazil, community health workers could play a bigger role, notably by taking a patient's medical history, conducting basic physical exams, ordering tests, counselling on preventive care. To date, CHWs in Brazil motivate patients to get care and bring test results to patients, check whether patients are taking medications, schedule appointments for individuals with other health professionals and encourage community members to seek care at the clinic (Wadge et al., 2016^[68]).

There is further scope to train CHWs in Brazil to directly address health needs and provide some clinical services, aided when necessary by remote communication with PHC doctors. The lack of opportunities to learn and advance in the community health worker role has already been noted elsewhere (Grossman-Kahn et al., 2019^[69]). So were the lack of recognition of their work, the poor communication and co-ordination between community health workers and the basic health units, as well as the lack of resources at the basic health units (Grossman-Kahn et al., 2019^[69]). Investments in CHW trainings, in streamlined IT systems and improved communication between CHW and other staff from the FHT will help expanding CHW roles.

Box 5.6. CHWs are the first point of contact with the health care system in remote part of the United States

In the United States, community health workers have a key role to address social determinants of health and to promote access to care for vulnerable and hard to reach populations (Hartzler et al., 2018^[67]). Community health workers, for example, provide PHC services in remote Alaskan villages, whose population would otherwise have no access to appropriate health care delivery (Golnick et al., 2012^[70]). They are the first point of contact with the health care system for the population living in these very remote villages. They work under the supervision of community health practitioners, and there is an integrated referral system that includes physicians, regional hospitals and a tertiary hospital (Golnick et al., 2012^[70]). The range of PHC services delivered by community health aides mostly includes care for chronic and preventive treatment, and emergency visits (often for respiratory distress and chest pain).

Furthermore, community health workers played a critical role as frontline health care workers during the first wave of the COVID-19 pandemic. In New York for example, community health workers provided educational sessions to patients with chronic conditions or at risks for chronic conditions to help them manage their conditions. They have also made home-visits, conducted wellness checks over the phone, helped people enrolling in online patient portals and prepared them for tele-health appointments. They served as support in navigating the health care systems, and mitigating fear and correcting misinformation in disadvantaged communities.

Source: Golnick et al. (2012^[70]), Innovative primary care delivery in rural Alaska, <https://doi.org/10.3402/ijch.v71i0.18543>; Hartzler et al. (2018^[67]) Roles and Functions of Community Health Workers in Primary Care, <https://doi.org/10.1370/afm.2208>.

Greater use of teleconsultations needs greater funding and new governance model

Improved use of alternatives to face-to-face consultations is another avenue for consideration to cope with PHC doctor shortages and geographical imbalances in Brazil. Telemedicine is used internationally to make PHC services available to patients closer to their home or work. It facilitates communication between patients and medical staff, as well as the transmission of medical records and other data between different locations (OECD, 2020^[61]).

Teleconsultation is a very promising way to improve access – both timely and geographically – and to relieve pressure on PHC physicians. Previous studies have shown that telemedicine improves access to care, reduces travelling costs and leads to better equity for rural and indigenous population (Oliviera Hashiguchi, 2020^[71]). Nearly 90% of international studies concluded that telemedicine interventions were at least as effective as face-to-face interaction (Oliviera Hashiguchi, 2020^[71]). Benefits can be found for preventing and controlling PHC sensitive conditions, like glycaemic control in diabetic patients, fewer hospitalisations for patients with chronic heart failure, effective pain management and effective management of health risk factors including exercise and nutrition (Oliviera Hashiguchi, 2020^[71]).

As already detailed in Chapter 6, telehealth has been in use in Brazil for quite some time now, especially in remote areas like the Amazon (Santos et al., 2016^[72]). In addition, the Brazil Telehealth Network Programme seeks to expand and improve PHC provision in regions with high social vulnerability, where the supply of health care services is very limited. The programme provides health care professionals that are part of the network both synchronous (real-time) and asynchronous teleconsultations, telediagnosics, second opinions and telelearning (Ministério da Saúde, 2020^[73]).

Telehealth services remain under-utilised in Brazil, despite evidence that points to beneficial effects. Available evidence shows that so far implementation of telehealth in PHC in Brazil has been associated

an improvement in child care, diabetes mellitus care, hypertension and women's health (Dos Santos et al., 2019^[74]). Despite this, from 2016 to July 2019, just under 3.5 million telehealth services across all levels of care (from tele-consultations to tele-learning sessions) had been registered in Brazil. Of these, 50% had taken place in the Southeast region and 36% in the South region. Only 1% of activities took place in the North region. Out of 26 Brazilian states, only 18 have telehealth centres participating in the network.

In many ways, telehealth initiatives in Brazil are promising, but several factors impede the development of telehealth. These impediments include the lack of funding, the lack of training, the lack of equipment for FHT and a fragmentation of governance mechanisms (see Chapter 6 for more details). Argawal et al. (2020^[74]) noted that Brazil lacks guidelines around reimbursement models and regulatory frameworks for telemedicine.

There are successful experience across the OECD, where telemedicine services have been used to expand access to care among rural and remote patients. Canada and Australia are two examples, where large investments have been made to make access to telehealth services a reality:

- In **Ontario, Canada**, the Telemedicine Network includes 120 virtual care systems to connect communities to care across the province (Ontario Telemedicine Network, 2021^[76]). In 2017, the Ontario Telemedicine Network supported more than 896 000 visits (Agarwal et al., 2020^[75]). The province of Saskatchewan has also been using telehealth technology for over a decade to reach rural and remote locations (Ontario Ministry of Health, 2021^[54]). Patients can receive PHC by visiting the nearest telehealth site and meeting with a professional in a virtual exam room. Telehealth can connect to available diagnostic peripherals, such as stethoscopes, vital signs monitors and ultrasound equipment, making real-time diagnosis and patient monitoring possible. In rural and remote areas, telehealth services are particularly used for mental health and addiction appointments, for prescription refills and chronic disease management. In addition, Saskatchewan began piloting remote presence technology for a number of North communities in 2017 (OECD, 2020^[61]). The project allows the PHC team in the community to have access to expertise (such as physician, nurse or pharmacist, etc.) on demand. Overall, previous experiences with telemedicine in Ontario (called Champlain BASE) have been shown to be cost-effective for the provision of specialist care. The cost of traditional referral to a specialist provider was, on average, CAD 133.60 per case; versus CAD 47.35 per case for a teleconsultation service on average (Agarwal et al., 2020^[75]).
- **Australia** has also made large investment in telehealth services to be used for specific health conditions such as chronic conditions and dermatology for people (OECD, 2015^[77]). In the Northern Territory, PHC doctors can be anywhere in Australia, and assist remote nurses and Aboriginal health practitioners to manage chronic disease patients appropriately by reviewing pathology and assessment results, then have case discussions with the local team. They also monitor and advise on other pathology testing. The medical practitioners consist of a group of general practitioners, who have usually previously worked in remote Northern Territory, and have moved away but can continue to provide quality care for remote patients through the use of a web-based electronic patient record with a unique patient identifier. Remote nurses and Aboriginal health practitioners can then contact a doctor who understands the conditions and circumstances they are providing care in. The Australian College of Rural and Remote Medicine also developed Tele-Derm, an online resource enabling rural doctors to receive advice on the diagnosis and management of skin disease. A rural doctor submits a photo of a skin condition, together with information on the patient's history and a possible diagnosis. A dermatologist responds usually within two days with a diagnosis and treatment options. PHC doctors can also access online case studies and education opportunities.

To address workforce shortages through a wider use of telehealth, Brazil will need to establish a governance structure that better support municipalities with insufficient resource capacity. Greater financial

support and adequate training will be required to avoid compounding existing social divides (see Chapter 6).

5.5. Increasing the recognition of the PHC specialty and ensuring high quality of medical education

Beyond addressing the uneven geographic distribution of PHC physician, there is an urgent need to increase the recognition of the PHC specialty and to ensure high quality of medical education in Brazil. Actions are needed in three broad areas:

- Standardising the contractual arrangement of PHC workers, with a more consistent salary scale across the country;
- Improving the academic incorporation of the family and community medicine specialty in graduate and post-graduate programmes to improve its credibility and recognition;
- Introducing quality measures to medical education to secure a high quality PHC workforce through a national licensing examination, making the specialty training in PHC compulsory for all medical doctors wishing to practise PHC, and implementing stronger requirements around continuous medical education.

5.5.1. Standardising the contractual arrangement for PHC workers

There are avenues for Brazil to improve the attractiveness of the PHC profession through a greater standardisation of contractual arrangement outside of the *Médicos pelo Brasil* programme. As each municipalities define the level of salaries, bonuses and other payment mechanisms, there is wide heterogeneity in these arrangements between municipalities. In Brazil, stronger oversight and regulation from the federal government is warranted to ensure consistency across the country. It could be important to make sure that small municipalities offer similar contracts as bigger municipalities, with a more consistent salary scale across the country. The new Agency of the Development of PHC could have a role to play in this direction.

Across OECD countries, Turkey implemented a similar strategy with the Health Transformation Programme launched in 2005. The overarching objective was to strengthen family medicine and PHC, notably through establishing new contractual arrangements for family physicians. Before the Health Transformation Programme, contractual arrangements and roles of family physicians were loosely defined. With the implementation of the health reform, family physicians have started to be paid according to national terms and conditions, and with more generous salaries and improved working conditions.

5.5.2. Improving the credibility and recognition of the specialty of PHC in Brazil

While the Family Health Strategy has made access to PHC a priority, the country has not invested well in building a strong credibility and recognition of the specialty of family and community medicine. In Brazil, the FCM specialty is the last option among physicians. Available evidence shows that the specialty of PHC is the first option for only 1.5% of recent graduates among the 55 specialties (Scheffer et al., 2018^[1]). It is important to add that the specialty is not recognised as a knowledge area by the Federal Agency for Support and Evaluation of Graduate Education, meaning that funding research and investment in this research area has traditionally been very low.

The low degree of academic incorporation of the family and community medicine specialty in graduate and post-graduate programme has been found to impede building a strong recognition of this specialty in Brazil (Wenceslau, Sarti and Trindade, 2020^[78]; Fontenelle et al., 2020^[79]). Few researchers build their carrier

around PHC, and perhaps as a consequence, PHC departments in universities and medical schools are very rare.

A strong academic incorporation is however critical to allow transmission of the specialties' value and allow a practice to be recognised by other health care professions. Without such promotion, it is very difficult to promote FCM among the students and to encourage them to choose this specialty as their future career. Low academic incorporation also reduces scientific production in the area of FCM, and limits it to collective health (Wenceslau, Sarti and Trindade, 2020^[78]; Fontenelle et al., 2020^[79]). Arguably, the specialty of family and community medicine is undervalued in Brazil, explaining why there is still bottleneck in filling physician vacancies in FHTs. Even with the implementation of the More Doctors programme, which has managed to allocate more than 16 000 new professionals to PHC since 2013, unfilled medical positions in FHTs are still common (Giliate Cardoso Coelho, Valeska Holst and Aristides, 2019^[18]).

Serious efforts are needed to counter the misconception that PHC is merely health care for the poor or the marginalised people, that it is a profession without much qualification, a second-rate profession (Justino, Luzón Oliver and Palhano de Melo, 2016^[19]). The knowledge, skills, responsibilities and core competencies of the family and community medicine speciality should be evident to all population groups and from all other health care professionals. The creation of academic departments of family and community medicine in all medical schools is a prerequisite to undertake research in PHC, develop clinical guidelines specific to PHC, as well as teach the specialty. Dialogue, inter-disciplinary collaboration with other specialists and mutual respect between health professionals are also key to support this.

When good family and community medicine curriculums are in place for both students and residents, it becomes possible to attract students to this specialty (Blasco et al., 2008^[80]). Family physicians who are recognised for their academic competence, leadership and who teach PHC practice with determination can awake a sense of vocation in their students, and will also win prestige among other health care professionals. Strong academic incorporation is also a prerequisite to develop and encourage further continuing medical education for all medical doctors who practice currently with no specialty training, helping to maintain and improve the quality of care.

Brazil could build up from experiences that have allowed to make good steps towards incorporating PHC as a core scientific area in some federal universities. For instance, the Master's Program in PHC at the Federal University of Rio de Janeiro, or the Family Health Master's Programs at the Federal Universities of Ceará and Mato Grosso do Sul. At the same time, Brazil could also build from successful experiences of network of Family Health Professional Master's, which attracted a high number of professionals: the programme of the Family Health Training Northeast Network (RENASF) created in 2009 (RENASF, 2012^[81]), and the Family Health Professional Master's (ProfSaúde) created in 2016 (ProfSaúde, 2019^[82]). Such good local initiatives could be consolidated by the federal government to design a National Family and Community Medicine postgraduate programme for Brazil.

5.5.3. Introducing quality measures to medical education

Implementing a national licensing examination

While the Ministry of Education in Brazil regulates medical education nation-wide, more could be done to secure standardisation of the medical curriculum in Brazil. The inadequate academic programme and gap in medical education between medical schools is an area of concern.

To achieve greater standardisation of the medical curriculum and teaching across the country, Brazil could introduce a national test at the end of the university period. The implementation of a standardised national exam would replace each final exam in medical schools. The national licensing examination will not replace accreditation performed by the INEP or the CFM, but rather complement it.

A national licensing examination would assess both knowledge and clinical skills of undergraduate students. This is a necessary step to improve the public trust and confidence in physicians. It will make transparent that all medical doctors in Brazil are competent in their clinical skills and medical knowledge. There are several advantages of establishing a national examination (Bajammal et al., 2008^[83]):

- First, the use of a standardised national exam is the best way to ensure any medical graduate or practicing doctor in Brazil has achieved at least a common standard of medical knowledge and clinical skills competencies. This would ensure that minimum standards and minimum competencies are met.
- Second, a national examination would act as a tangible and standard measurable outcome to identify areas in need for potential improvement in the curriculum and teaching methods. It could be used to benchmark medical schools across the country, and to correct any potential problem or deficiency in the curriculum or teaching method.
- Third, a national exam could be used as a reliable and transparent measure of candidates' qualification for post-graduate program's application. In other OECD countries such as France and Italy, allocations to specialist schools are determined by the results of the national examination. This approach is seen as a more meritocratic and transparent way to attribute specialist training places.

The national examination has been increasingly used across OECD countries. This is a mandatory requirement to practise medicine in Canada, France, Italy, and just recently in the United Kingdom. While the United Kingdom has relied on external examiners and inspection carry-out by the General Medical Council to assure quality across the United Kingdom medical schools, the country is introducing a Medical Licensing Assessment which will test the core knowledge, skills, and behaviours needed to practise medicine (Box 5.7). The overarching objective is to drive up high standards for medical schools in the United Kingdom (CMC, 2020^[84]). Brazil could learn from the United Kingdom's experience in building the Medical Licensing Assessment, and the role played by the General Medical Council.

At the same time, it will be important to ensure that doctors who qualified abroad pass a national licensing examination in order to be eligible to practise in Brazil. This approach is taken internationally – for example in the United States, Canada, France or the United Kingdom. In these countries foreign medical graduates must pass licensing examinations in order to be eligible to practise in the country, and then be able to enrol in a residency programme and pass the post-graduate examination. This system ensures every foreign graduate has at least the minimum competences to function as a physician, according to the licensing body standards of the country. In the United Kingdom, international medical graduates have to pass the Professional and Linguistic Assessments Board (PLAB) test (GMC, 2021^[85]). The PLAB Test helps make sure doctors who qualified abroad have the right knowledge and skills to practise medicine in the United Kingdom. From early 2024, international medical graduates will also have to take the MLA if they are applying for registration with a license to practise in the United Kingdom (Box 5.5). In Australia, international medical graduate seeking general registration as a medical practitioner needs to undertake the Standard Pathway which consists of the completion of a written examination (CAT MCQ) and structured clinical examination of the Australian Medical Council (AMC, 2020^[86]).

With the new *Médicos pelo Brasil* programme, the Revalida system will be a compulsory requirement for foreign medical graduates. This is an important step going forward to ensure that all medical doctors who qualified abroad have the right knowledge and skills to practise medicine in Brazil. As in the United Kingdom and Australia, revalidation consists of a theoretical written test and a practical clinical skill test, covering all level of health (primary, secondary, clinical, surgical). It might be warranted to better target the Revalida Test toward specific clinical areas or level of care depending on the profile of each graduate to better recognise their qualifications.

Box 5.7. The United Kingdom is introducing a Medical Licensing Assessment

The United Kingdom has traditionally relied on external examiners – visiting medical educators from other organisation – and General Medical Council inspection to assure quality across UK medical schools. Medical schools used to set their final exam independently in line with the General Medical Council's outcomes for graduates. However, the United Kingdom is introducing a Medical Licensing Assessment (MLA) which will test the core knowledge, skills, and behaviours needed to practise in the country. Medical students graduating in the academic year 2024-25 will need to pass the MLA as part of their medical school degree before they can join the medical register. The MLA test will have two-part: an applied knowledge test that will be computer-based, and a clinical and professional skills assessment. The aim is to provide a standardised mean of assessing all UK doctors, which in turn will help ensure patient safety. The benefit of the new system is, for the first time, to demonstrate that graduates from each medical school have met an agreed standard of proficiency and are well prepared to practise medicine. The work entailed by the MLA will be overseen and regulated by the General Medical Council. This work will consist of:

- Defining the range of professional skills, knowledge and behaviours a candidate needs to have achieved to be ready to practise medicine in the United Kingdom;
- Approve procedures to compile test questions and papers, set standards and run exams;
- Take corrective action if through its quality assurance processes, it considers that standards are not met;
- Be responsible for using information and data from the exams to apply a consistent approach to the assessment of international medical graduates.

Under this agreement, medical schools will continue to develop and deliver their own curricula and prepare students for the MLA, which will be regulated by the General Medical Council.

A process of phased introduction will begin from 2021 with robust testing and piloting, and the assessment will be fully implemented for students graduating from UK medical schools from the academic year 2024/25.

Source: CMC (2020^[84]), <https://www.gmc-uk.org/news/news-archive/gmc-and-msc-welcome-agreement-on-medical-licensing-assessment-delivery-model>

The specialty of family and community medicine should be a mandatory requirement to be allowed to practise PHC in Brazil

As earlier mentioned, most doctors working in FHTs do not have a specialist post graduate training in FCM. Brazil is currently aiming for each FHT to have a FCM specialist with the implementation of the new Doctors for Brazil programme. The new programme will make the specialty training in PHC compulsory for all medical doctor wishing to practise PHC. Arguably, this new programme will help deliver more effective and patient-centred care. The benefits of specialist PHC has already been demonstrated elsewhere: PHC specialists place a stronger emphasis on preventive interventions and early management of health conditions, and are more likely to address the majority of patient's need, as well as to help them to co-ordinate care with other health services contributing to better quality and responsiveness (OECD, 2020^[61]).

Currently, at least 22 OECD countries (Australia, Austria, Belgium, Canada, the Czech Republic, Denmark, Estonia, France, Greece, Israel, Italy, Ireland, Latvia, the Netherlands, Norway, Portugal, Slovenia, Spain, Sweden, Switzerland, Turkey and the United Kingdom) have introduced a post-training

requirement to practise PHC. Such a professionalisation of PHC specialists goes along with the formation of professional organisations, scientific research in PHC, or the development of guidelines. This is warranted for driving up quality and patient-centred PHC in Brazil.

Requirements around continuous medical education for PHC medical doctor could be implemented

Continuing medical education (CME) includes the acquisition of new knowledge, skills and approaches considered useful for developing competent and experienced medical practice. It helps medical doctors maintain their competence and acquire new skills throughout their careers. This is critical to keep practice up-to-date, and to make sure medical doctors possess the skills useful and needed in daily practice to respond to evolving patients' needs.

In Brazil, the Política Nacional de Educação Permanente em Saúde was instituted in 2004 as the main mechanism to encourage continuing professional development. It is based on participatory teaching and learning approaches at the practice level. The overarching objective being to improve the skills and fill gaps in competence according to patients' health needs in daily practice, rather than updating medical knowledge using didactic approaches. While continuing professional development is a good practice to continuously and actively improve the skills and competences of medical doctor, the uptake of this activity within FHTs remains unknown.

In addition, CME is voluntary in Brazil. There is no specific requirement for each medical doctor. The take-up of CME relies upon professional and individual choice, without any regulatory approach, peer evaluation requirement or re-certification requirement. The current system does not guarantee high standard of competencies for PHC doctors, and does not ensure their fitness to practise throughout a career. So far, it is impossible to know whether and how many PHC medical doctors have undertaken CME.

A number of bodies such as the CONASS, CONASEM, the Brazilian Medical Association, the SBMGC or the SOBRAMFA organise several training, conferences and internet-based learning model. These are all valuable approaches. However, a more formal CME framework, which makes clear the expectation upon PHC doctors and supports them to meet it, will constitute a marked improvement. Brazil could introduce strong requirements around CME for PHC physicians, as experienced by other OECD countries including Norway, the Netherlands or the United Kingdom (OECD, 2014^[12]). In the United Kingdom and the Netherlands, CME for PHC medical doctor is compulsory and linked to re-certification. PHC physicians need to demonstrate that they have regularly participated in CME activities and there is a peer evaluation regarding professional skills. To be successfully re-certificated, PHC physicians need to achieve at least 50 CME credits per year. Norway use financial incentives to increase compliance with CME requirements. Specialists in PHC are able to charge higher fees for each consultation than regular GPs if they have followed a number of CME courses. This gives strong incentives to comply with national requirement, which in turn improve their competencies and performance.

Experiences from OECD countries could be helpful to inform Brazil on ways to implement requirements and mechanisms to secure compliance to CME requirements. The Agency for the Development of PHC and municipalities could for example add a requirement to demonstrate the attendance or completion of some specific training in the local contracting of PHC doctors. Using the contracting arrangement could actively push PHC doctors toward compliance with CME requirements, and encourage quality improvement. If strong CME requirements are introduced in Brazil, the country should also ensure that CME is directed towards some of the key challenges of the health care system, for example on prevention of risk factors for health and management of chronic conditions.

5.6. Conclusion

In Brazil, problems of shortage of PHC doctors and regional disparities in supply are acute. They are leading factors of lower health outcomes, lower health care quality and can result in unmet health care needs for Brazilian's most remote population. While the implementation of the More Doctor Programme in 2013 led to positive improvements with greater investment in PHC infrastructure, better access to medical training, and increased number of recruitment in underserved communities, more needs to be done to increase and ensure a more even distribution of PHC medical doctors. The new federal programme *Medicos pelo Brasil*, which has a stronger focus on quality than the previous programmes, aims at expanding the number of PHC doctors to 7 500 in the poorest areas. To meet this ambitious target, strategic thinking toward the community targeting methods will be required to make sure that all eligible municipalities are effectively enrolled to the new federal programme.

At the same time, and as informed by the experiences of OECD countries, it will also be important to increase the number of rural trained doctors and to provide both wage- and non-wage related financial incentives – linked for example to a service obligation of between two and four years – to recruit and retain doctors in underserved areas. Restricting the choice of practice location is another strategy to ensure a more even distribution of medical doctors. At any rate, such solutions require careful and coherent workforce planning in Brazil, based on an objective assessment linked to socio-demographic characteristics of medical doctors, their work processes and population health needs.

A smarter use of existing local workforces (nurses and community health workers), and changes in scopes of practice, are innovative ways that Brazil might want to consider in the future. The country should also continue to expand the use of telehealth to facilitate access to PHC in places with fewer physicians. To this end, it will be crucial to establish a governance structure that better supports municipalities with insufficient resource capacity to avoid compounding existing social divides.

While much policy attention has been directed to improving access to PHC, little attention has been devoted to building a strong credibility and recognition of the PHC specialty, and to ensuring high quality and well-skilled medical doctors. Actions to improve the academic incorporation of the FCM specialty in graduate and post-graduate programmes are needed through for example the development of additional family and community medicine departments in medical schools. To secure high quality PHC medical doctors and ensure their fitness to practise throughout their carrier, there are high opportunities to make the specialty training in FCM compulsory for all medical doctors wishing to practise PHC, and to implement stronger requirements around continuous medical education and recertification.

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6 The digital transformation of primary health care in Brazil

Brazil has made strides towards a digital transformation of PHC, building on more than a decade of policies to digitalise health care and make better use of health data, and with key investments in networks, data, interoperability and skills. The COVID-19 pandemic has only provided further impetus. Yet, progress towards effective use of digital PHC has been slow, and fundamentally unequal, with significant inequalities in the use of digital technologies and tools among health workers and citizens. Other major barriers include human and technical capacities in municipalities, with potential diseconomies of scale resulting from setting responsibilities for digital health at municipal level. Tackling these challenges requires: 1) digitalising all PHC units and teams; 2) promoting inclusive connectivity for all Brazilian citizens, especially the most vulnerable; and 3) establishing a governance structure with clear well-funded mandates at the right levels of government.

6.1. Introduction

A digital transformation of health care in Brazil is well underway, propelled by a broader strategy for a digital transformation of the public sector. For some time now, the Government of Brazil has been increasing its use of digital tools and data to make its public institutions more efficient, more transparent and more responsive to citizen and business demands (OECD, 2018^[1]). Clear governance and a focus on connectivity, interoperability, open government data and citizen-driven service delivery have resulted in Brazil recently scoring above the OECD average on the 2019 Digital Government Index (OECD, 2020^[2]).

Brazil has made significant inroads with respect to building a foundation for digital PHC, from strategies at various levels of government to essential investments in networks, data, interoperability and skills. However, progress towards effective use of digital technologies in PHC has been slow, and fundamentally unequal. Contributing factors include both demand and supply side factors, but also disarticulated governance and insufficient and poorly targeted funding.

This chapter provides an overview of recent initiatives and developments in the digitalisation of PHC in Brazil. Over the last decades, Brazil has been making impressive progress towards a digital transformation of its public sector and its health care services, but progress towards effective use of digital technologies in PHC has been slow, and fundamentally unequal, as discussed in section 6.2. Moreover, as shown in section 6.3, millions of Brazilians are digitally excluded, and there is a risk that a digital transformation of PHC will only compound existing social divides. The chapter concludes with a discussion in section 6.4 of how Brazil's decentralised government, along with the country's heterogeneous regional demographics, socio-economic indicators and governance structures, create challenges to digitalising PHC.

6.2. Overall progress in health system digitalisation but less so in the digitalisation of PHC

The creation of the Brazilian Internet Steering Committee (*CGI.br*) in 1995 to provide strategic guidelines related to the use and development of the Internet in Brazil can only be considered foundational (OECD, 2018^[1]). The Committee established the basis for an ecosystem between citizens, the private sector and the public sector. Linked to the Brazilian Internet Steering Committee, and sitting under the Brazilian Network Information Centre (*NIC.br*), the Regional Centre for Studies on the Development of the Information Society (*Cetic.br*) was created in 2005 to monitor the adoption of information and communication (ICT) technologies in Brazil. It has conducted its ICT Households Survey annually since 2005, and has, since 2013, also undertaken the ICT in Health Survey focusing on the infrastructure, availability and use of ICT-based applications in health care facilities in Brazil. These surveys are invaluable and they provide many of the statistics and indicators used in this chapter. Unfortunately, the most recent results predate COVID-19 and thus may not fully reflect the current situation.

6.2.1. The government has made strides with a clear ambition for digital health services

Since 1995, and especially in recent years, Brazil has made impressive progress towards a digital transformation of its public sector. In 2019, more than 500 public services, spanning 29 different entities, were made available through digital channels, leading to reported annual savings of USD 66.9 million¹ (BRL 345 million) (gov.br, 2020^[3]). Under its current Digital Government Strategy 2020-22, the government is aiming to offer all of its more than 3 000 public services via digital channels by the end of 2022 (Diário Oficial da União, 2020^[4]). Among the existing digital transformation plans of 73 public entities is the Brazilian National Digital Health Strategy 2020-28 (*Estratégia de Saúde para o Brasil para 2028* or *ESD28*).

The *ESD28* (2020^[5]) builds on more than a decade of policies to digitalise health care and make better use of health data, going as far back as the National Policy for Health Information and Informatics of 2004 (*Política Nacional de Informação e Informática em Saúde* or *PNIIIS*). There has been an acceleration of efforts in more recent years, with the publication of the 2015 *PNIIIS*, which is currently being reviewed, the publication in 2017 of the e-Health Strategy for Brasil 2020 (*Estratégia e-Saúde para o Brasil 2020*), and the publication in 2020 of Brazil's National Digital Health Strategy Action, Monitoring and Evaluation Plan (*Plano de Ação, Monitoramento e Avaliação da Estratégia de Saúde Digital para o Brasil 2019-23*). The *ESD28* is also aligned with Brazil's National Health Plan 2020-23 (Ministério da Saúde, 2020^[6]).

Actions within the *ESD28* are built around three axes (Ministério da Saúde, 2020^[5]). The first axis seeks to build on existing programmes and initiatives to digitalise health care in Brazil, including the National Health Data Network (*Rede Nacional de Dados em Saúde* or *RNDS*), as well as the Connect SUS (*Conecte SUS*) and the *Informatiza APS* programmes (these are described in greater detail below). The second axis looks to develop the necessary conditions (e.g. organisational, legal, regulatory and governance structures) to spur effective collaboration in digital health. The third and final axis will establish a conceptual, normative, educational and technological forum (the so-called *Espaço de Colaboração*) to operationalise collaboration. Training and capacity building for health care workers is a key priority in the implementation of the *ESD28*.

The *ESD28* plan of action has seven priorities (Ministério da Saúde, 2020^[5]): 1) governance and leadership, 2) digitalisation of the three levels (primary, secondary and tertiary) of health care provision, 3) supporting health care quality improvements, 4) the patient as the protagonist, 5) human resource training, 6) an environment of interconnectivity, and 7) an innovation ecosystem. The *ESD28* also identifies the main actors and stakeholders, establishing five types of participation (leadership, partnership, sponsorship, contribution, engagement) that are expected of each actor for each action point (see Box 6.1).

Box 6.1. Digitalisation of PHC in the National Digital Health Strategy 2020-28

Action point 2.1.2 of the *ESD28* seeks to expand the digitalisation of PHC in Brazil. It is led by the Ministry of Health and the Health Secretaries of Municipalities and States, in partnership with the CONASSEMS and CONASS, with sponsorship from the Federal Government, the National Health Council and the CIT, plus contributions from Associations of Health Professionals, and engagement from Anvisa and ANS. The foundation of this action point is the Primary Health Unit Computerisation Programme (*Programa de Apoio à Informatização e Qualificação dos Dados da Atenção Primária à Saúde*) or *Informatiza APS*, which seeks to digitalise all PHC units and teams in Brazil. New financing schemes (part of *Programa Previne Brasil*) and implementation and training initiatives have been piloted in the state of Alagoas (more detail is provided in the text below). The timeline for full implementation of the action point is December 2022.

Source: Ministério da Saúde (2020^[5]), *Estratégia de Saúde Digital para o Brasil 2020-28*, http://bvsms.saude.gov.br/bvs/publicacoes/estrategia_saude_digital_Brasil.pdf.

There is a comprehensive push for a digital PHC in Brazil

The importance of connecting and digitalizing PHC practices is clear and the potential value added would be very significant (see Box 6.2). Digital health can help PHC be the first point of contact with patients, co-ordinate referrals and follow-up to secondary and tertiary care, and monitor and keep track of community and population-level health risks and conditions (Castle-Clarke and Imison, 2016^[7]). The COVID-19 pandemic has made clear there are enormous benefits to digitalising and connecting health care facilities and services.

Box 6.2. A digital transformation can help realise the full potential of PHC

Digital PHC is the use of digital technologies to support PHC (it is a subfield of digital health focusing specifically on PHC). Digital technologies (or information and communication technologies) are electronic tools, systems, devices and resources that generate, store, process and/or transmit data. Digital technologies include devices such as smartphones and computers, as well as intangible products such as software, platforms and algorithms, and can support PHC in a variety of ways.

At the centre of digital PHC is the **electronic health record** or EHR. The EHR is a longitudinal digital record of an individual patient that contains or virtually links together multiple electronic medical records from different institutions and health care settings. As it contains a history of contact with health care providers from any organisation at any level of care, the EHR plays an essential role in helping PHC providers co-ordinate care for individual patients.

An EHR that is complete, accurate, up-to-date, comprehensive, and that can be shared, enables a plethora of digital applications in PHC. An EHR that contains clinical (e.g. medications prescribed), sociodemographic (e.g. employment) and administrative (e.g. insurance) information allows supports many applications, including among many others:

- Targeted **alerts and reminders** to patients (e.g. for screenings or vaccinations).
- Electronic referrals, **ePrescribing** and certifications (e.g. for sick leave).
- Clinical decision making (e.g. medications to avoid given patient allergies).
- **Telemedicine** applications (e.g. video consultations and remote monitoring).
- Mobile health or **mHealth** applications (e.g. smoking cessation and mental health apps).
- Population-level statistics on the prevalence and incidence of diseases and risks.
- Research studies taking into account **real-world data** (e.g. pharmacovigilance).

It is possible to implement some of these applications without a complete EHR system (e.g. telemedicine), however without the EHR, digital technologies will fall short of their potential and will not fully assist PHC.

Source: World Health Organization (2018^[8]), Digital technologies: shaping the future of primary health care, <https://apps.who.int/iris/bitstream/handle/10665/326573/WHO-HIS-SDS-2018.55-eng.pdf?sequence=1&isAllowed=y>; OECD (2019^[9]), Health in the 21st Century: Putting Data to Work for Stronger Health Systems, <https://dx.doi.org/10.1787/e3b23f8e-en>.

The potential benefits of a digital transformation of PHC are also clear in the context of Brazil's National Health Plan 2020-23 (Ministério da Saúde, 2020^[6]). Among its objectives are to promote the expansion and effectiveness of PHC services in an integrated and planned fashion, to guarantee access to quality PHC services for all, and to reduce population health risks through prevention, epidemiological surveillance and health promotion.

Brazil has a long history of developing, implementing and using health information systems (Ministério da Saúde, 2017^[10]). In 2017, the government launched a public tender, the Primary Health Unit Computerisation Programme, to support the country's PHC units in using electronic health record (EHR) systems by enhancing connectivity, acquiring ICT equipment, and providing technical support and training of health professionals (OECD, 2020^[11]). Following concerns from the Federal Court of Accounts regarding the viability of a national rollout, the programme was replaced by Connect SUS. Connect SUS seeks to bring together all the health information of any one citizen under a large data network that can be used to continuously improve health care and decision making. The Connect SUS programme depends on two essential elements: the RNDS and the updated Primary Health Unit Computerisation Programme

(*Programa de Apoio à Informatização e Qualificação dos Dados da Atenção Primária à Saúde* or *Informatiza APS*).

Through the Connect SUS programme, and as part of the *Previne Brasil* programme, the *Informatiza APS* programme provides federal funding (between USD 330 or BRL 1 700 and USD 388 or BRL 2 000 per month) to family health teams that are able to collect and send data from electronic health record (EHR) systems (Harzheim et al., 2020^[12]). The programme includes the provision of training in ICT and data analysis to public employees in municipal health departments (CGI.br, 2020^[13]). The training is jointly organised by the Ministry of Health and the University of Rio Grande do Sul. It consists of 60 hours of training delivered through Telehealth Centres. In order to qualify for support from *Informatiza APS*, family health teams need to have already made investments in digital technologies, have a working EHR system and be able to share data through the PHC Health Information System (the *Sistema de Informação em Saúde para a Atenção Básica* or *SISAB*). Those teams that are eligible may use the funds from *Informatiza APS* to acquire software or hardware, pay for Internet connectivity in UBS, and hire ICT services, among others.

A key eligibility requirement of the *Informatiza APS* programme is that family health units need to already be using digital technologies. However, Connect SUS ran a pilot project in the state of Alagoas between October 2019 and June 2020 that did include an additional financial incentive for PHC units that have not yet been able to digitalise (Ministério da Saúde, 2020^[14]). The Alagoas pilot provided a lump sum of between USD 1 649 (BRL 8 500) and USD 1 940 (BRL 10 000) to each municipality in Alagoas for every PHC unit that was not yet digitalised (more than USD 1.07 million or BRL 5.5 million were disbursed to 97 municipalities). In light of the COVID-19 pandemic, the Ministry of Health instituted a time-limited incentive of USD 5 268 (BRL 27 157) to be paid to municipalities and states for each health team that has not yet been digitalised (see Box 6.3). The Ministry of Health is also collaborating with the Ministry of Science, Technology, Innovation and Communications, with support from the National Education and Research Network (*Rede Nacional de Ensino e Pesquisa* or *RNP*), to provide Internet connectivity to 16 000 PHC facilities (CGI.br, 2020^[13]).

A key element of Connect SUS focusing specifically in PHC is the e-SUS APS strategy (or *Estratégia e-SUS APS*, often also referred to as *e-SUS AB* for *Atenção Básica*) which seeks to modernise PHC through the use of health information systems, in order to better manage individual health and care, optimise data collection and improve health information (Ministério da Saúde, 2020^[6]). The strategy is based on two underlying data collection systems: the simplified data collection system (*Coleta de Dados Simplificada*) and the EHR system (*Prontuário Eletrônico do Cidadão* or *PEC*). Around 56% of family health units that have been digitalised use the *PEC*, while others use third-party EHR systems (Ministério da Saúde, 2020^[14]).

The Informatics Department of SUS – typically referred to as DATASUS – has also made various other applications available to citizens, frontline care workers and health care managers (CGI.br, 2020^[13]). These include: *MedSUS*, an application that lists all medications approved for use in SUS; *e-Gestor AB*, a platform that combines multiple PHC information systems to facilitate the planning and delivery of health services; and the *Conecte SUS Cidadão* app, which builds on a previous version called *Meu digiSUS*, which allows citizens to access their personal and clinical data collected from multiple national registries as well as access useful information, such as where are the closest health care facilities and pharmacies. There are also versions of *Conecte SUS* for frontline health workers (*Conecte SUS Profissional*) as well as managers.

Another interesting system is *DigiSUS Gestor – Módulo Planejamento (DGMP)*, a tool made available to states, the Federal District and municipalities in May of 2019. The *DGMP* allows local managers to register population health targets, annual plans, produce and send reports to the National Health Council, as well as produce annual reports. The Ministry of Health also makes available an Application Programming Interface, or API, that allows different software platforms used by health care facilities to integrate their EHR systems with the *RNDS*, as well as make use of blockchain solutions.

Municipalities are also increasingly providing digital services to citizens (CGI.br, 2020^[13]). For example, the São Paulo prefecture has developed an app called *Agenda Fácil* which allows patients to book, confirm and cancel consultations and examinations in the municipality's PHC units. According to the latest ICT Electronic Government Survey (2020^[15]), 25% of Brazilian prefectures allow patients to book appointments online and in some cases to be consulted via the Internet.

Start-ups in telemedicine are also becoming increasingly relevant. *Brasil Telemedicina*, founded in 2010, offers online medical services. It produces around 60 000 reports per month and has clients in more than 700 Brazilian cities (Brasil Telemedicina, 2020^[16]). Another startup, *Telelaudo*, provides tele radiology services 24 hours a day, seven days a week, and employs 120 radiologists. It has provided more than 4.7 million reports to more than 450 hospitals (Telelaudo, 2021^[17]).

Box 6.3. COVID-19 has been a catalyst for digital health and telehealth in Brazil

The federal government authorised on 15 April 2020 the use of telemedicine during the COVID-19 pandemic (through law number 13.989/2020). This enabled continuity of care for patients who may have feared physically visiting health care facilities, but it also protected health care professionals most at risk of serious COVID-19 (such as those with chronic conditions, those who are pregnant and those aged 60 years old and older) (see Chapters 3 and 4). To help PHC units access digital technologies, the Ministry of Health also instituted an exceptional incentive to be paid to municipalities and states for each health team that has not yet been digitalised, budgeting over USD 83.8 million (BRL 432 million) for 2020. The Brazilian Telemedicine University Network (*RUTE*) also created a special interest group on COVID-19.

Artificial intelligence, specifically machine learning algorithms, has also been employed to monitor the spread of SARS-CoV-2 infections and to actively search for individuals who may have been exposed to the virus. Algorithms have been used to predict how the virus will spread further, to assess the clinical outlook of individual patients as well as support decisions concerning the most appropriate treatments, and to manage hospital beds and other resources.

Chatbots and virtual assistants have been used by public authorities to provide accurate information on COVID-19 to populations, and also to collect data that can be used to monitor the spread of the virus. The Ministry of Health created a WhatsApp channel to answer questions from citizens as well as health care professionals, and to fight disinformation. Various states – such as Ceará, Rio Grande de Sul, and Amazonas – have also used digital tools to communicate with their populations and to aid decision making during the pandemic.

This rapid adoption of digital tools is potentially introducing some risks. For example, the emergency approval of telehealth during COVID-19 (law 13.989/2020) does not include any language relating to medical liability, data and information protection and security, medical licensing across states, nor location of medical act for tax purposes. While in some cases, it is clear that other existing laws apply (e.g. with respect to data protection and medical liability), the lack of specific text on these issues may introduce significant uncertainty and lead to lower adoption than would be desirable.

Source: CGI.br (2020^[13]), ICT in Health 2019; CGI.br (2020^[15]), ICT Electronic Government 2019; Diário Oficial da União (2020^[4]) PORTARIA Nº 3.193, DE 27 DE NOVEMBRO DE 2020; Caetano et al. (2020^[18]), Challenges and opportunities for telehealth during the COVID-19 pandemic.; <https://doi.org/10.1590/0102-311X00088920>.

6.2.2. Progress towards digital health has been uneven, especially in PHC

Out of the 40 880 PHC units (from a total of 43 200) that responded to the ICT in Health Survey (2020_[13]), it is estimated that 3 600 did not have computers and 7 200 were not connected to the Internet. Internet connection speeds in PHC facilities increased from 2018 to 2019, but still only 17% of PHC units had speeds over 10 Mbps (CGI.br, 2020_[13]). Across all health care facilities, only 20% of public facilities had speeds over 10 Mbps compared to 60% of private facilities. The United States Federal Communications Commission (FCC) recommends a minimum speed of 10 Mbps for a small practice with two to four physicians (ONC, 2019_[19]). Speeds under 4 Mbps are not recommended if practices want to make simultaneous use of EHR systems and high-quality video consultations, not to mention remote patient monitoring. Because many public PHC facilities are small, these facilities do not actually contract broadband connections themselves, but rather these are procured by municipal and state health secretaries. This could be why almost half (46%) of managers in public health facilities do not know the speed of their broadband connections (CGI.br, 2020_[13]).

While health care managers increasingly perceive their ICT equipment and Internet connections to be up-to-date and appropriate, there are marked differences between perceptions of managers in public and private health facilities (CGI.br, 2020_[13]). A majority (68%) of managers of private facilities consider their ICT equipment to be new and up-to-date, compared to only 36% of managers in public facilities, and 40% of managers in PHC units. Similarly, 67% of managers in private facilities considered their Internet connection to be appropriate given their needs, compared to only 31% of managers in public health care facilities, and 32% in PHC units. These same patterns are observed among frontline care workers, like physicians and nurses. Across both public and private facilities, managers in non-capital cities were less likely than managers in capitals to consider their ICT equipment and Internet connections were up-to-date and appropriate.

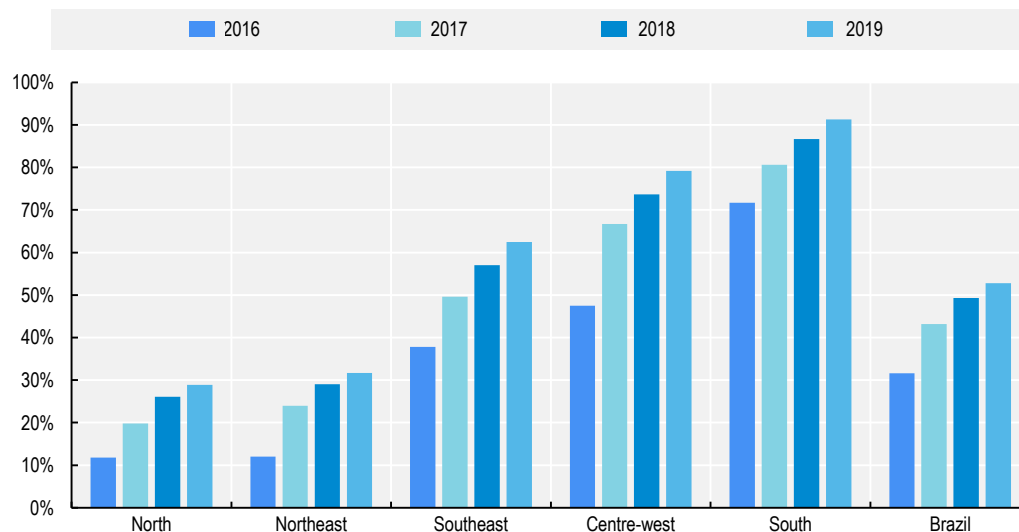
Only 14% of public health care facilities (from large institutions with over 50 beds to smaller PHC practices) had an ICT department, compared to 38% of private facilities (CGI.br, 2020_[13]). Among PHC units, 10% had in-house ICT teams. The share of public facilities (16%) that had an information security policy was significantly lower than the share of private facilities with such a policy (44%). Among PHC units, 15% had an information security policy in place. Around 14% of public facilities had adopted data loss protection and prevention tools, compared to 29% of private facilities. The use of information security tools in PHC units was especially underdeveloped, with only 40% of facilities having a firewall, only 27% with encrypted databases, and 16% having data loss prevention tools. Close to a third of PHC units did not have password-protected access to electronic systems.

In 2019, an estimated 82% of all health care facilities and 78% of PHC units had EHR systems (CGI.br, 2020_[13]). Around 18 000 health care facilities (or 18% of all facilities) did not have an EHR system, 12 000 of which were public facilities. According to the National Register of Health Establishments (CNES), 62% of FHTs were digitalised (using the same definition as that used in *Informatiza APS*), leaving almost 20 000 teams not digitalised (Ministério da Saúde, 2020_[14]).

Adoption of EHR systems was highest in the South and Southeast regions (at 90% and 83% of facilities in those regions, respectively) compared to the Northeast and North regions (77% and 74% respectively). Facilities in non-capital cities lagged behind those in capital cities in the use of EHR systems (79% of non-capital facilities compared to 92% of facilities in state capitals). Close to one in every three public facilities, and 25% of all PHC units, still maintained clinical records and patient data only in paper form, compared to one in ten private facilities. In particular, while a majority of physicians in public facilities do use electronic prescribing, 71% of physicians signed their prescriptions by hand, rather than electronically. Between 2016 and 2019, the number of PHC units using EHR systems increased from 13 375 to 21 557, a 61% increase in three years (Ministério da Saúde, 2020_[6]). Yet, the evolution has been very unequal across regions of Brazil (see Figure 6.1).

Figure 6.1. Very large inequalities in the use of EHR systems in PHC units

Share of PHC units using EHR systems in Brazilian regions, 2016-19

Source: Ministério da Saúde (2020^[6]), Plano Nacional de Saúde 2020-23.

Many of Brazil's indigenous populations are based in the North and Northeast regions where digitalisation is least advanced. Indigenous children under one year old have lower rates of utilisation of medical appointments to monitor their growth and development (Ministério da Saúde, 2020^[6]). There are also significant inequalities in the rates of vaccination against yellow fever in Brazilian states where vaccination has been recommended, ranging, in 2018, from 21.8% in Sergipe to 100% in the Federal District and Roraima (the national average for all states where vaccination is recommended is 64.1%).

Digital tools can help tackle these inequalities but the underlying infrastructure needs to be in place. Unfortunately, the rate of digitalisation of PHC units in Brazilian states seems to be inversely proportional to inequalities in health status and need. The share of hospital admissions due to conditions that are treatable in PHC is higher in regions with lower use of EHR systems in PHC units: the North and Northeast regions. Digital PHC could help reduce unnecessary admissions through, for example, better prevention, screening and monitoring of conditions that are treatable in PHC (see Box 6.2), but this cannot happen if the needed enablers – from infrastructure to software to skills – are not in place.

While the majority of health care facilities did have an EHR system in place, the information and data contained within these systems was not always complete, limiting the potential benefits. Decision support capabilities, specifically, were low, with only around a fifth of health facilities (including both large inpatient facilities with more than 50 beds as well as smaller PHC units) having EHR systems that allow, for example, drug allergy and drug dosage alerts (CGI.br, 2020^[13]). Only around a third of facilities were capable of sending and receiving discharge reports or clinical information (ibid).

Since 2018, the ICT in Health Survey collects data on how many health care facilities have electronic systems in place to send or receive data directly from electronic systems in other health care facilities (be it PHC practices or hospitals). In both 2018 and 2019, the share of health care facilities that had systems in place to exchange information was 19%, suggesting that interoperability and data exchange remain limited in Brazil (CGI.br, 2020^[13]). This is a serious barrier to effective PHC, greatly limiting care continuity and co-ordination.

Around one in five public health care facilities and PHC units offer digital services directly to patients (CGI.br, 2020^[13]). These include scheduling appointments (23% of PHC units), exams (20% of units) as well as accessing results online (22% of units). Around one in ten allow patients to access their health record online and 7% of PHC units enable communications with health professionals.

Telehealth has been in use in Brazil for quite some time now, especially in remote areas like the Amazon (Santos et al., 2016^[20]). In the rest of the country, however, its use has been very uneven. Around one-fifth of health care facilities provide remote learning (or e-learning) services (CGI.br, 2020^[13]): continued education for health care professionals delivered via videoconferencing. The share of PHC units using remote learning is higher, reaching 39%. This is the most frequent use of telehealth, with other applications, such as real-time teleconsultations and remote monitoring, having lower adoption rates. Having said that, PHC units make more use of telehealth than the average health care facility in Brazil, with 35% providing teleconsultations and 25% telediagnosis. For example, remote monitoring has been implemented in less than one in ten facilities.

Unlike with the use of EHR and the Internet, it is public health care facilities that are most active in telehealth networks, with virtually no participation among private providers (CGI.br, 2020^[13]). Among public facilities, 32% report being part of state telehealth networks, 19% report being part of the Brazil Telehealth Network Programme (*Programa Nacional Telessaúde Brasil Redes*), and 11% report participating in the Telemedicine University Network (*RUTE*). Among PHC units, the shares were 40%, 23% and 14% for state networks, the Brazil Telehealth Network Programme and *RUTE* respectively. Despite these encouraging figures, 47% of all public health care facilities are not part of any telehealth network.

The Brazil Telehealth Network Programme seeks to expand and improve health care provision, especially through PHC. The programme provides health care professionals that are part of the network both synchronous (real-time) and asynchronous teleconsultations, telediagnosics, second opinions and telelearning (Ministério da Saúde, 2020^[6]). In 2015, the SMART (*Sistema de Monitoramento e Avaliação de Resultados do Programa Telessaúde*) system was created to monitor the use of telehealth services in Brazil. From 2016 to July 2019, just under 3.5 million telehealth services (from teleconsultations to telelearning sessions) had been registered by SMART. Of these, 50% had taken place in the Southeast region and 36% in the South region. Only 1% of activities took place in the North region. Out of 26 Brazilian states, only 18 have telehealth centres participating in the network.

Few health care professionals participated in ICT training courses in 2019, according to the latest ICT in Health Survey (2020^[13]). Around 36% of nurses and 30% of physicians reported having completed capacity-building courses in health informatics, 8% of nurses and 6% of physicians participated in specialisation courses, and only 1% of nurses and 2% of physicians signed up for master's or doctorate degrees in health informatics.

Notably, around 60% of nurses and 43% of physicians did not perceive any added or reduced workload as a consequence of using computers and the Internet (CGI.br, 2020^[13]). Nurses were less likely than physicians to say that computer and Internet use had decreased their workload (17% of nurses compared to 33% of physicians). More broadly, health care professionals have overwhelmingly positive views of the use of electronic systems in their facilities. Around 90% of physicians and nurses considered the implementation of electronic systems improved efficiency in team work. Over 80% agreed that the use of electronic systems resulted in higher quality of care as well as greater efficiency in services.

6.2.3. Effective digital PHC requires a focus on the fundamental enablers

A real digital transformation of PHC in Brazil will not happen without some foundational basics. These basic enablers – individually necessary though not sufficient – include Internet connectivity, appropriate hardware (e.g. computers, tablets, servers, etc.), standards and interoperability for data collection and sharing, software (like EHR systems), ICT security, and, crucially, appropriate training of clinical and

managerial staff, including through change management. All these elements require investments and resources at different levels of government, from the federal government all the way down to the municipalities and family health teams.

Reliable energy supply and fast connectivity are indispensable

Appropriate electricity supply and Internet connectivity are vital. As discussed, a significant number of PHC units do not have an Internet connection and a majority have speeds that are likely insufficient to make simultaneous use of EHR systems, high-quality video consultations and remote patient monitoring. Among the main barriers to digitalisation reported by 12 municipalities in the context of the recent pilot project in Alagoas were unreliable electricity supply and a lack of sufficiently fast broadband (Ministério da Saúde, 2020^[14]). In some cases, there were no broadband suppliers (ibid). A 2018 study of 91 electricity suppliers serving 81 million households, found the quality of supply varies across regions, with citizens in remote localities, like the Amazonas, having significantly less stable electricity than someone living in São Paulo (Idec, 2018^[21]).

Brazil's geography (nearly eight times the size of France and Spain combined) and sparsely distributed population are certainly challenging for inclusive connectivity (OECD, 2020^[22]). The Brazilian Digital Transformation Strategy (E-Digital), along with the Connected Brazil programme, have been instrumental in expanding broadband connectivity (OECD, 2020^[11]). Yet, connectivity targets have focused exclusively on schools and have not made explicit mention of concrete connection speeds or reliability. Brazil could explore how rural co-operatives might help bring stable electricity supply and affordable broadband, as done in North Dakota, and build on private sector efforts to connect rural populations in Peru (see Box 6.4).

Box 6.4. Bringing stable electricity supply and Internet connectivity to rural areas

The Dakota Carrier Network: a consortium of small, independent rural companies and co-operatives

Local mutual and co-operative organisations, which were crucial in bringing electrical and telephone services to rural areas of the United States in the 1900s, are now attempting to do the same with Internet connectivity. With more than 200 rural co-operatives are building advanced broadband networks across the country, evidence suggests these networks may be cheaper and faster than private-sector networks (OECD, 2021^[23]). In North Dakota, a rural and sparsely populated state, in 2019, rural residents had better connection speeds than average fibre speeds in both rural and urban areas nationwide. This achievement has been credited to the Dakota Carrier Network, an umbrella organisation made up of small, independent rural companies and co-operatives. With financial support via the federal Broadband Technology Opportunities Programme, besides connecting citizens, the organisation is also deploying a dedicated 10 Gbps network to cover health care providers to enable telemedicine and data exchange.

Internet para Todos: A private sector collaboration to bring Internet connectivity to rural Peru

A collaboration between IDB Invest (the private-sector arm of the IDB Group), Telefónica, Facebook and the Development Bank of Latin America (CAF), *Internet para Todos* (Internet for All) is a wholesale operator with a new business model that aims to provide Internet connectivity in rural areas of Peru at a profit (IADB, 2020^[24]). The financial institutions provide funding with a long-term vision, while Telefónica makes its infrastructure available to rural operators. By the end of 2021, the company expects to have connected 6 million people in more than 30 000 rural areas. Expansion into other parts of Latin America is being considered, starting in Colombia.

Source: OECD (2021^[23]), Delivering Quality Education and Health Care to All, <https://dx.doi.org/10.1787/83025c02-en>; IADB (2020^[24]), <https://www.iadb.org/en/improvinglives/internet-para-todos-helping-latin-america-log>.

Remote and rural municipalities face significant barriers to dependable and fast connectivity, as well as a stable energy supply. These locations often have more limited resources yet face higher costs from broadband suppliers than better resourced large municipalities in urban areas. Partnerships between the Ministry of Health, the Ministry of Science, Technology and Innovation and the Ministry of Defence to connect PHC units, especially in remote locations, are crucial. It is important to evaluate the impact of these initiatives. For example, the already mentioned programme under the *RNP* that aims to connect 16 000 primary health units is limited to 12 months and does not seem to set concrete targets in terms of connection speeds or reliability. Without adequate Internet connectivity, the poorest and most remote regions of Brazil will be further left behind, with a digital transformation at national level compounding rather than reducing existing regional inequalities.

Purely financial incentives can be helpful but capacity building is still a major barrier

The Connect SUS pilot project in Alagoas provided municipalities with up to USD 1 940 (BRL 10 000) per unconnected PHC unit (Ministério da Saúde, 2020^[14]). Despite this, in the period under evaluation, digitalisation in Alagoas lagged behind the average in the Northeast region as well as the overall Brazilian average (*ibid*). There is no question that financial support is needed to help PHC units and municipalities acquire essential ICT products and services. Yet, the results in Alagoas show financial support alone is not enough. One of the barriers to digitalisation reported by the 12 municipalities in Alagoas was the lack of an implementation team (Ministério da Saúde, 2020^[14]).

It is not only digital skills among health care workers that matter, but also the skills, time and ownership needed to drive digitalisation of PHC units. The Alagoas pilot planned to conduct capacity-building workshops in three regions in the State, but due to COVID-19 ended up conducting workshops in only one region. That region (the *1ª Região de Saúde de Alagoas*) was responsible for half of the total increase in digitalisation during the pilot. The workshop trained professionals from 19 municipalities on the installation and use of the *e-SUS AB EHR system (PEC)* and other *e-SUS* software, as well as training those professionals attending how they themselves could train their colleagues. Another key activity in the workshop was to establish the timeline and responsibilities for implementation of the EHR system. While there are other reasons for the success of the *1ª Região de Saúde de Alagoas*, such as high degree of urbanisation, it is likely that the capacity-building workshop had a positive impact.

It is not just health care workers but also crucially managers that lead digitalisation of care services. In England, since 2017, the NHS Digital Academy has been training clinical and information technology leaders to support the digital transformation of the NHS. The Academy is delivered through a partnership between Imperial College London, the University of Edinburgh and the Harvard Medical School (Socha-Dietrich, 2021^[25]). Its flagship Digital Health Leadership programme combines content in leadership and change management, health informatics and data analytics, health systems and user centred design, as well as citizen informatics, among other subjects. In Brazil, the implementation of the *Programa Educacional em Saúde Digital* is a step in the good direction. The programme was instituted in 2020 by the *Secretaria de Gestão do Trabalho e da Educação na Saúde* and in partnership with the Federal University of Goiás. In 2021, the programme includes specialisation in digital health for both health care workers and health managers at States and Municipal levels.

As mentioned, the EHR system is at the core of digital PHC. It is impossible to send out vaccination alerts, to suggest therapies based on clinical records, and to predict clinical outcomes when patient records are not digitalised. As seen, the large majority of health care facilities, especially smaller PHC practices, do not have decision support and information exchange capabilities within their EHR systems. However, Brazil has shown it is capable of making rapid progress. In just three years (2016-19), the share of public health care facilities with EHR systems capable of (CGI.br, 2020^[13]): listing every lab result a patient has had increased from 17% to 41%; listing every patient that is using a specific pharmaceutical went from 18% to 40%; electronic drug prescribing increased from 29% to 51%. These are key functionalities in support of

PHC, especially when it is estimated that the purpose of one in five medical appointments is not clinical but to ask for prescriptions and certificates (Mendes, 2011^[26]; Marshall, Shah and Stokes-Lampard, 2018^[27]). Tying financing of digital PHC to the use of specific functionalities can be a powerful way to increase adoption (more on this in section 6.4.2).

Box 6.5. Key recommendations to increase digitalisation of PHC units

Ensure reliable energy supply and fast connectivity in all PHC units

- Set ambitious and clear targets for both reliability and speed of connections as part of existing and future collaborations between the Ministry of Health and other ministries and the *RNP* to increase Internet connectivity in PHC units. Improve co-operation among governmental entities and across the different levels of government (federal, state and municipal) for implementation of broadband connectivity initiatives.

Expand digitalisation of PHC units through financial support and capacity building

- Expand financial payments included in the Connect SUS pilot project of Alagoas to the entire Brazilian territory. Condition receipt of financial support to participation of municipal and state officials in capacity-building workshops, and develop training specifically for senior executives. Set ambitious and clear quantitative targets for adoption and use of EHR systems (such as *e-SUS AB*), including effective use of electronic drug prescribing and exchange of clinical and administrative data across units. Emphasise importance of cybersecurity as an inevitable consequence of PHC units' operational dependence on digital systems.

Note: see also recommendations in the *OECD Telecommunications and Broadcasting Review of Brazil 2020* (OECD, 2020^[22]) and the *OECD Reviews of Digital Transformation: Going Digital in Brazil* (OECD, 2020^[11]), as well as the *Recommendation of the Council on Broadband Connectivity* (OECD, 2021^[28]).

6.3. Digital divides pose barriers to an effective digital PHC

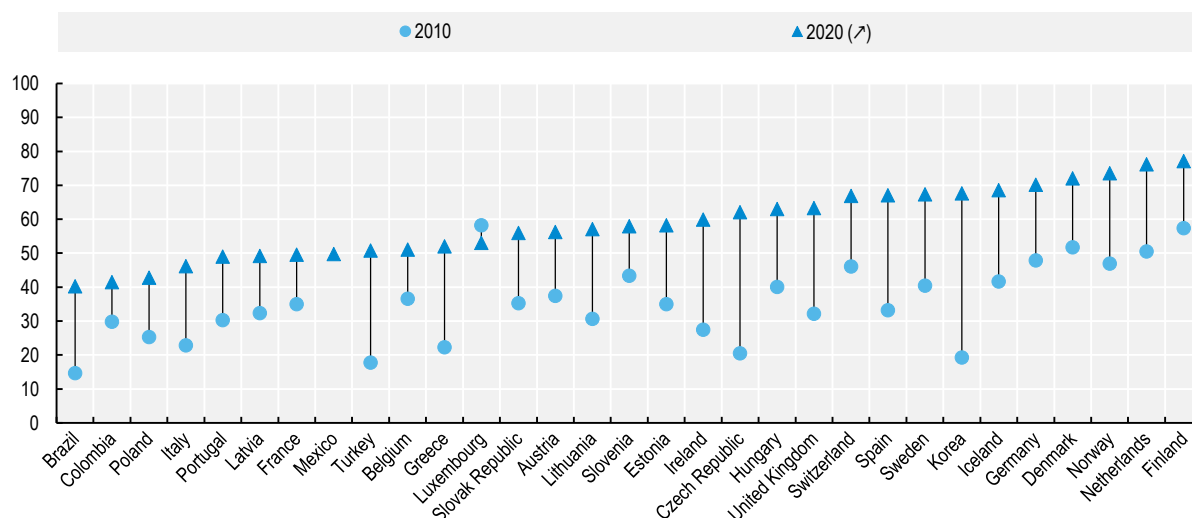
Even if Brazil were to break down all barriers to connectivity and digitalisation of PHC units, there would still persist significant inequalities in the use of digital technologies and tools among its citizens. While the country has made enormous progress in recent years, millions of Brazilians are still digitally excluded, and what is worse there is a clear association between digital and social inequalities in the country. Evidently, digital PHC cannot be effective if populations – especially those that most need it – do not have access to digital services.

6.3.1. Brazilian citizens have never been as connected but clear digital divides persist

According to the most recent ICT Households Survey, in 2019, 50.7 million Brazilian households (or 134 million individuals) had Internet access (corresponding to 71% of all households), an increase of 5.2 million connected households in relation to 2018 (CGI.br, 2020^[29]). Brazilians are more connected but also more digitally engaged. More than two in three Internet users aged 16 or over used electronic government services in 2019 (ibid). Between 2010 and 2019, the share of individuals using the Internet to look for health information more than doubled from around 15% in 2010 to 40% in 2019 (see Figure 6.2). Despite these clear improvements, the country ranks below OECD countries in indicators of digital uptake and use, and digital divides tend to reflect “analogue” divides (OECD, 2020^[11]; BNDES, 2017^[30]).

Figure 6.2. More Brazilians are seeking health information online

Share of individuals using the Internet for seeking health information in the last three months, in 2010 and 2020



Note: Differences in methodology in Colombia in 2010 and in Korea in 2010 and 2019. Most recent data point for Colombia is for 2018 and for France, Switzerland and Mexico it is 2019 (not 2020). For Brazil specifically, the data are for 2010 and 2020 and are collected by CETIC.

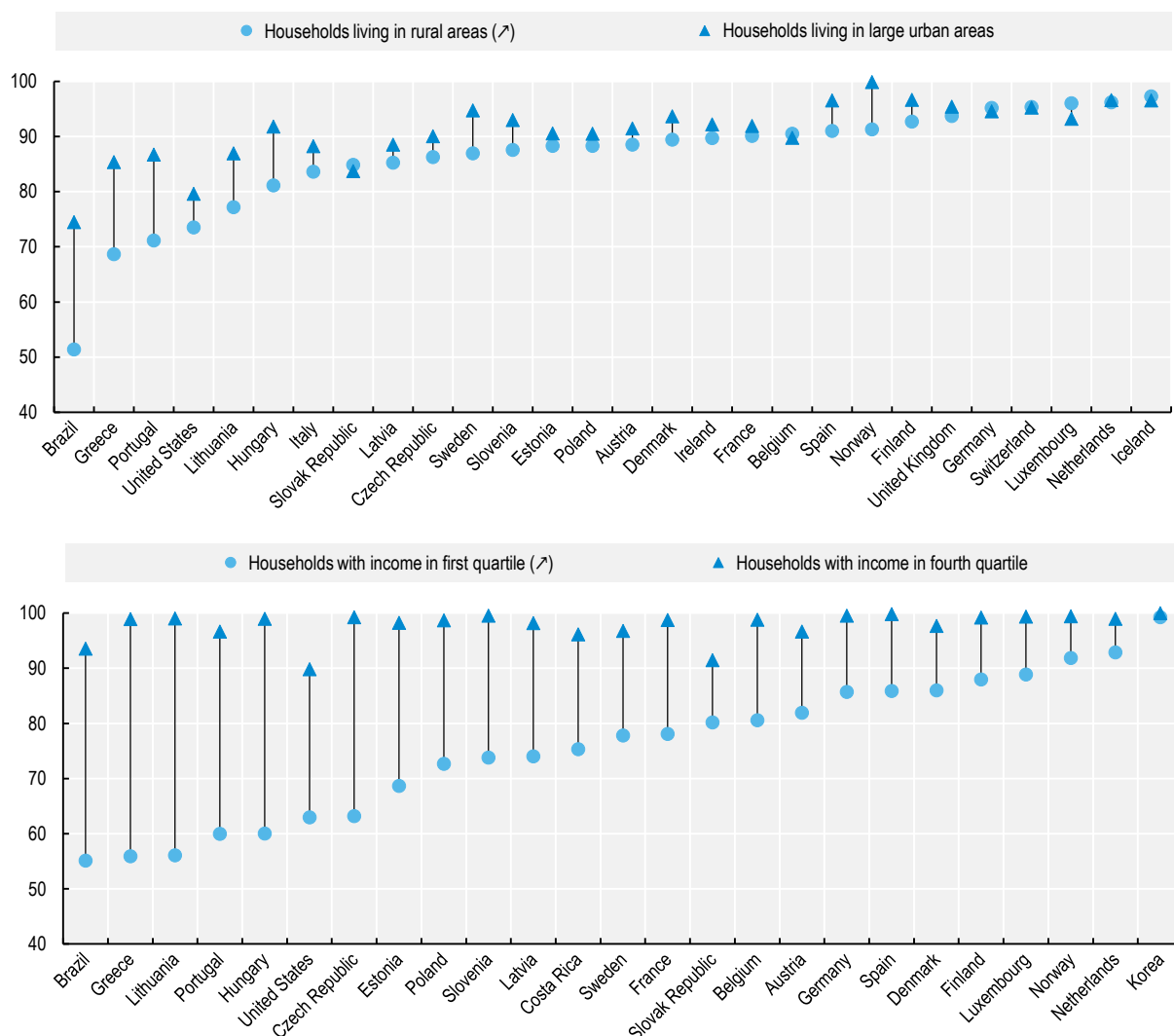
Source: OECD ICT Access and Usage by Households and Individuals (database), http://stats.oecd.org/Index.aspx?DataSetCode=ICT_HH2.

Digital divides mirror social inequalities limiting effective digital PHC provision

There is a clear association between digital and social inequalities in Brazil (see Figure 6.3). Despite important increases in connectivity, nearly 47 million Brazilians did not use the Internet in 2019 (CGI.br, 2020^[29]). Among non-users, the large majority had lower education (85%) and lower socio-economic status or SES (95% belonged to classes C and DE²). Furthermore, regional disparities in use persist. The difference between the proportion of connected households in the Southeast (75%) and the Northeast (65%) regions is of 10 percentage points (ibid). According to the most recent ICT Households Survey (2020^[29]), close to 20 million Brazilian households did not have Internet access in 2019, of which 6.4 million households were from the Northeast region. Households with the lowest SES made up the majority of households without Internet access (13 million households in classes DE did not have access in 2019). In 2019, nearly 35 million were disconnected in urban areas (23%) and 12 million in rural areas (47%).

Figure 6.3. There are important digital divides in Internet access at home

Share of households with broadband Internet access at home, 2020



Note: Differences in methodology for income in Korea, Brazil and the United States, and for rural/urban in the United States only. Data on income for Brazil, the United States, Costa Rica and France are for 2019; data on rurality for Brazil, the United States, France, the United Kingdom and Switzerland are for 2019. For Brazil specifically, the data are collected by CETIC. Income data are collected using ranges in minimum wages. Reported data were aggregated in a way that best fit the distribution into four balanced categories. Income non-response are not included. Areas are defined as urban or rural according to local legislation, as compiled by the NSO. Reported data refer to urban (densely populated) and rural (thinly populated).

Source: OECD ICT Access and Usage by Households and Individuals (database), http://stats.oecd.org/Index.aspx?DataSetCode=ICT_HH2.

As mentioned, in 2019, 40% of Internet users in Brazil searched for information on health or health care services online (CGI.br, 2020^[29]). The share of individuals seeking health information on the Internet was lower among those aged 60 years and older, as well as in groups with lower SES (classes DE). Only 23% of Internet users searched for or used public health services (CGI.br, 2020^[29]).

While Brazil has made significant progress in implementing digital government services, certain groups of individuals are not taking full advantage. According to the 2019 ICT Household Survey (2020^[29]), those less likely to use e-government services include people aged 60 years and older (46% use e-government services compared to 75% of 25 to 34-year-olds), those with lower education levels (46% of those with

elementary education compared to 87% of those with tertiary education), and those from groups with lower SES (48% in classes DE compared to 88% in class A).

6.3.2. Equitable digital PHC starts with ensuring digital uptake and use

It is clear that a digital transformation of PHC can have multiple benefits for populations and individuals, but there is also a risk that digital divides may add to social divides, compounding existing inequalities and deepening social exclusion. With digital divides so clearly aligned with social inequalities in Brazil, there is a serious risk that a digital transformation of PHC will compound existing social divides. An effective digital PHC that reaches all Brazilians is only possible if every citizen is not only connected but can have a say in the design of digital PHC services.

Digital inclusiveness as a foundation for digital PHC for all

Brazil's policies for digital inclusion have focused on bringing the Internet to remote areas and isolated communities (OECD, 2020^[11]). This is the case with the Electronic Government – Citizen Assistance Service (*Governo Eletrônico – Serviço de Atendimento ao Cidadão* or *GESAC*) programme, created in 2002 to provide free broadband Internet connections to telecentres, schools, health care units as well as indigenous villages and socially vulnerable communities (OECD, 2018^[11]). Other programmes³ include the Digital Inclusion Programme, the Connected Citizen Project, the One Computer Per Student Project and the Telecentros.BR Programme (OECD, 2018^[11]). The Internet for All (*Internet para Todos*) programme is a more recent expansion of the *GESAC*, focusing on subsidising broadband connections for the more than 30 000 localities and districts in Brazil without a connection.

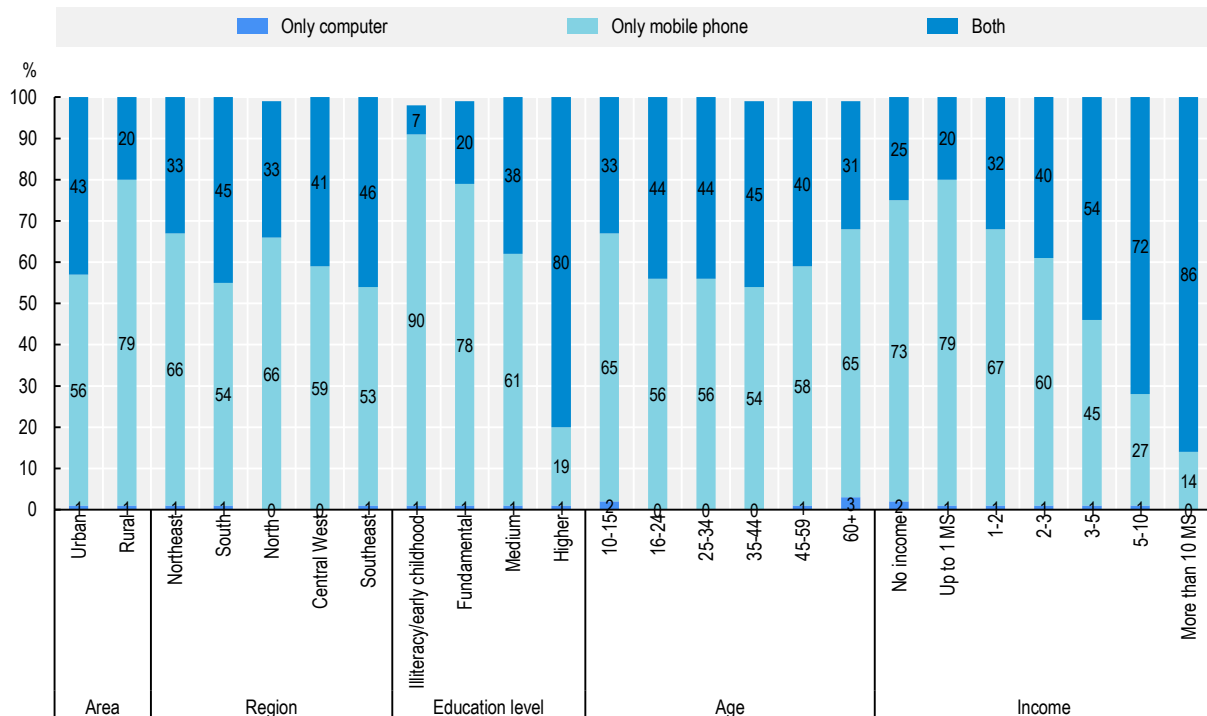
The creation of telecentres throughout Brazil in the 2000s was one of the first initiatives at the federal level to promote digital inclusion (CGI.br, 2020^[31]). Telecentres are public spaces that provide computers with an Internet connection free for anyone to access, as well as workshops, courses and training. According to the most recent ICT Public Access Centres 2019 Survey, which gathers information on telecentres, 45% of telecentres were not operational in the three months prior to the data collection, the most cited reason being a lack of computers and Internet connection, followed closely by lack of maintenance and financial resources (CGI.br, 2020^[31]). The number of telecentres in use decreased from the previous Survey in 2013, one possible reason being that the original telecentre programme stipulated that after an initial five-year period of support by the federal government, the equipment would be donated to the municipalities (OECD, 2020^[11]).

In 2015, the Federal Audit Office (*Tribunal de Contas da União*) reviewed public policies related to digital inclusion, finding difficulties in the co-ordination of the different levels of government, limited previous studies to inform and frame new policies and actions, and no systematic monitoring and evaluation of programmes (Tribunal de Contas da União, 2015^[32]). In 2018, the Federal Audit Office again drew attention to the lack of co-ordination across the federal, state and municipal governments, and pointed out that many municipalities do not have any telecentres (Tribunal de Contas da União, 2018^[33]).

Many policies and programmes to promote digital literacy and ICT use focus on younger people and on schools, which are undeniably important and will lead to improvements in digital inclusion in the coming decades (OECD, 2020^[11]). Yet, it is older people who most need and access PHC (Mendes et al., 2019^[34]), and more must be done today to include older Brazilians in the digital transformation. Besides strengthening the role of telecentres as providers of training, especially in rural and remote areas, it is important to develop a multi-channel approach to the inclusion of vulnerable groups who are most likely to need and access PHC. Such an approach cannot ignore the crucial role that mobile phones play as a means to access the Internet for older individuals as well as those from rural areas, those on lower incomes, and those with lower levels of education (see Figure 6.4).

Figure 6.4. Mobile phones are the only means to access the Internet for more vulnerable groups

Older, rural, lower income and lower education groups are more likely to use only mobile phones to access Internet



Note: MS = minimum salary.

Source: ICT Households 2019: Survey on the Use of Information and Communication Technologies in Brazilian Households (database), <https://cetic.br/en/pesquisa/domicilios/indicadores/> (accessed in March 2021).

Access to mobile broadband is behind the rapid increase in Internet access among the Brazilian population (OECD, 2020^[11]). Reliance on mobile connectivity exclusively may prevent some users from accessing more sophisticated services that require higher speeds (e.g. real-time teleconsultations), either because those speeds are not available or data plans are prohibitively expensive. The fact that exclusive use of mobile phones for Internet access is associated with more vulnerable groups could mean that those citizens who most stand to benefit from digital PHC may also be less likely to be able to access it. This is why it is important to adopt a dual strategy of developing new channels to reach these vulnerable groups on mobile platforms, coupled with improvements in high-speed broadband connectivity and digital skills, for example through an expansion of telecentres.

The use of short message service (SMS) and mobile applications to deliver government services is already a reality in many countries, especially in Korea, a country that topped the OECD 2019 Digital Government Index, and where citizens can access government services on their mobile phones. In the United States, the government has set up a mobile first strategy to ensure government services are mobile friendly (United Nations, 2020^[35]). In Finland, the government has created a centralised multi-channel one-stop-shop portal that is linked with other specialised portals, thus promoting equity in citizen access to digital government services (ibid).

Adopting a user-driven approach to digital PHC

A *user-driven* approach describes government actions that allow citizens and businesses to indicate and communicate their own needs and, thereby, drive the design of government policies and public services

(OECD, 2020^[36]). This requires a shift from a government-centred focus to a user-centred approach that places the needs of users at the core of digital transformation processes and services. A common pitfall of many digital health initiatives is that they fail to adequately engage with health professionals and citizens, leading to products and services that are not relevant to users and add no value (WHO and ITU, 2012^[37]).

User needs should be the basis of a digital transformation of PHC. From service design and development through to implementation and monitoring, the user should be present during the entire lifecycle (OECD, 2020^[36]). Throughout the process there should be opportunities for feedback and even co-creation. What is key, especially in the Brazilian context, is that a user-driven approach that is accessible and inclusive is also an approach that produces more accessible and inclusive services. Coupled with a strategy to increase digital literacy among citizens, a user-driven approach can engage those citizens that are both most vulnerable and most digitally excluded. Indeed, the now widely used participatory budgeting approach is a Brazilian creation, first implemented in the city of Porto Alegre (OECD, 2019^[38]). Participatory budgeting allows citizens a say in how (usually local) budgets are formed and how public moneys are spent. In Porto Alegre, widely seen as a success story, thousands of people have voiced their preferences over the years, affecting how millions of dollars were spent, and importantly – because minorities and vulnerable groups were sometimes overrepresented – shifting public money to where it is most needed (Local Government Association, 2016^[39]).

Box 6.6. Key recommendations to promote citizen uptake and use of digital PHC

Focus on eliminating digital divides as a foundation for effective digital PHC

- Strengthen the role of telecentres as providers of training, especially in rural and remote areas, and ensure appropriate funding and technical assistance at all levels of government. In parallel, develop a multi-channel approach that focuses on mobile access to reach vulnerable groups who are most likely to need and access PHC.

Users must be at the centre, if not drivers, of a true digital transformation of PHC

- Develop content, services and applications that meet the needs of both health professionals but especially citizens, particularly those with low digital uptake, e.g. low-educated, low-income and elderly people. Continue to expand patient access to their personal health information.

Note: see also recommendations in the *OECD Reviews of Digital Transformation: Going Digital in Brazil* (OECD, 2020^[11]) and the *OECD Digital Government Review of Brazil: Towards the Digital Transformation of the Public Sector* (OECD, 2018^[11]), as well as the *Recommendation of the Council on Broadband Connectivity* (OECD, 2021^[28]).

6.4. A governance structure that is fit to deliver a digital PHC

Brazil's government is divided into 26 states and the federal district, and well over 5 000 municipalities. Municipalities are granted the status of federal entities, at the same level as the states. While there is great heterogeneity in the geographical area and population, as well as social and economic indicators, among the subnational jurisdictions, all Brazilian municipalities enjoy the same legal status (OECD/UCLG, 2019^[40]). In 2015, 17 municipalities had more than 1 million inhabitants (22% of population). About 44% of municipalities had less than 10 000 inhabitants. Brazil also has 438 health regions, which are made up of municipalities, and are also very heterogeneous in their demographics, socio-economic indicators and governance structures (Ministério da Saúde, 2020^[6]). Collaboration and co-ordination across these multiple structures is very challenging, yet it is crucial for a successful digital transformation of PHC.

6.4.1. Brazil's decentralised government poses challenges to digital PHC

Decentralisation can have economic and administrative benefits, from services that are better aligned with local needs, to more innovative local experimentation, to economic growth driven by competition and accountability in public services (OECD, 2019^[38]). However, there are challenges too. Some of the most relevant drawbacks in the context of a digital transformation of PHC in Brazil include overlapping responsibilities, diseconomies of scale, and lack of human and technical capacities.

A digital transformation of PHC cannot take place without adequate governance arrangements, from leadership and co-ordination to institutional models and resources (OECD, 2020^[2]). The *OECD Digital Government Index* sets out four transversal facets that can be used to frame progress towards digital public services, including in PHC (see Box 6.7): a *strategic approach* with a clear vision, objectives, goals and actions; the specific *policy levers* and tools to connect strategies to policy actions; *implementation* capacity that transforms goals and strategies into effective and concrete programmes; and finally *monitoring* activities to take stock of progress and feedback into strategies.

Box 6.7. The OECD Digital Government Index

The OECD Digital Government Index (DGI) is a cornerstone of the OECD work on digital government and public sector data. Built on the OECD Recommendation on Digital Government Strategies, the Index responds to the need for a measurement instrument to track the adoption of digital government policies across OECD member and partner countries. The Digital Government Index draws upon the OECD Digital Government Policy Framework, which characterises a mature digital government as:

- being **digital by design** when governing and leveraging digital technologies to rethink and re-engineer public processes, simplify procedures, and create new channels of communication and engagement with stakeholders;
- being **data-driven** when valuing data as a strategic asset and establishing the governance, access, sharing and re-use mechanisms for improved decision-making and service delivery;
- **acting as platform** when deploying platforms, standards and services to help teams focus on user needs in public service design and delivery;
- being **open by default** when making government data and policy making processes available to the public, within the limits of legislation and in balance with national and public interest;
- being **user-driven** when according a central role to people's needs and convenience in the shaping of processes, services and policies; and by adopting inclusive enabling mechanisms;
- being **proactive** when anticipating people's needs and respond to them rapidly, avoiding the need for cumbersome data and service delivery processes.

Source: OECD (2020^[2]), Digital Government Index: 2019 results, <https://doi.org/10.1787/4de9f5bb-en>.

Brazil's strategic approach is exemplary, embodied in the excellent and comprehensive ESD28. However, like other OECD countries, Brazil has been better at adopting strategies than at promoting and monitoring their implementation. The ESD28 sets out an ambitious vision for digital health in Brazil, but it will be difficult to realise. This is because there are significant risks, as identified in the WHO/ITU *National eHealth Strategy Toolkit* (2012^[37]), namely: insufficient and geographically imbalanced resource capacity, scarce funding, too broad of an implementation focus, and unavailability of required infrastructure (this last one has already been discussed in previous sections). To these barriers, one can add a lack of clarity in the assignment of responsibilities across levels of government.

In Brazil, the division of responsibilities is unclear in a number of areas, including in health care (OECD, 2019^[38]). This is frequently the case in policy areas that are “shared” across levels of government, as is the case with health care in Brazil. This can result in duplication, cost shifting and scale inefficiencies. The government’s 2017 eHealth Strategy noted that initiatives to create a national eHealth architecture had lacked alignment and continuity, leading to frustration, scepticism that such a programme could work and demotivation in tackling the inevitable barriers (Ministério da Saúde, 2017^[10]). It was also mentioned that strategies and investments had been traditionally defined and executed in isolation by the federal, state and municipal governments. Despite the existence of high-quality initiatives in digitalising health care, these remained “islands of excellence” due to a lack of integration across levels of government (Ministério da Saúde, 2017^[10]).

Small municipalities, especially, face significant barriers to digitalisation of PHC units and teams. Because responsibilities are “shared”, the three levels of government are often dependent on each other to implement the actions set out in the ESD28. This can lead to delays in adoption due to holdups in different levels of government (Andrade et al., 2019^[41]). Municipalities are responsible for many key steps in digitalising PHC, from purchasing ICT equipment and software to hiring ICT specialists. Smaller municipalities may have limited capacity to manage these processes. One example is the requirement that the contracting of essential services be renewed every five years, which has been flagged as an impediment to the expansion and upkeep of ICT infrastructure (Ministério da Saúde, 2017^[10]). It is not uncommon for smaller municipalities, and even larger ones, to not have any long-term ICT planning (Veloso et al., 2011^[42]).

Financial commitments have also perhaps not been as ambitious as the agendas set out in previous strategies and in the ESD28. This has led to a situation in which some municipalities are responsible for digitalising PHC services but lack the necessary resources to do so. In other words, these are unfunded mandates (OECD, 2019^[38]). Since 2014, the Ministry of Health has provided a monthly payment of USD 117 (BRL 603.38) for PHC units who seek to improve their connectivity (Ministério da Saúde, n.d.^[43]). The target was to connect 12 000 units, however only 8 000 had been connected and only 4 800 were sending data back to the Ministry of Health.

More recently, the *Informatiza APS* programme provides monthly support of USD 330 (BRL 1 700) to USD 388 (BRL 2 000) to family health teams that are able to collect and send data to central repositories. A key eligibility requirement of the programme is that family health units need to already be using digital technologies. The pilot project in the state of Alagoas has expanded support to include an additional financial incentive for PHC units that have not yet been able to digitalise (a lump sum of between USD 1 649 (BRL 8 500) and USD 1 940 (BRL 10 000) to each municipality in Alagoas for every PHC unit that was not yet digitalised). As this is still a pilot, this form of pre-digitalisation support is not yet available for all municipalities. In light of the COVID-19 pandemic, the Ministry of Health did institute a time-limited incentive of USD 5 268 (BRL 27 157) to be paid to municipalities and states for each health team that has not yet been digitalised, but it is still too early to know what impact this has had on the numbers of PHC units without computers and Internet access.

6.4.2. Moving from digital strategies to implementation and monitoring

Determining which level of government should take over which responsibility related to digitalisation is not without challenge, but it is also of the utmost importance. Clear assignment of responsibilities is crucial for accountability, monitoring and effectiveness of investment (OECD, 2019^[38]). Particularly with shared responsibilities, as is the case with health care in Brazil, clarity is all the more important, so that all parties mutually understand what their individual responsibilities are. It is important to avoid multiple co-ordination mechanisms with no clear role in decision-making processes (OECD, 2019^[38]). Finally, it is vital to build the necessary capacity to deliver on mandates. This means aligning funding with functional responsibilities

to avoid unfunded mandates, but also encouraging sharing of best practices and specialised knowledge across jurisdictions, making expertise available to all levels independent of size (OECD, 2019^[38]).

Setting clear well-funded mandates for digitalisation at the right levels of government

A major barrier to digital PHC in Brazil is human and technical capacity in municipalities. Having more than 5 500 municipalities procure ICT products and services for over 42 000 PHC units every five years is not good practice. Around one in five Brazilian municipalities have less than 5 000 inhabitants, and yet they are responsible for hiring ICT specialists and purchasing ICT products and services, from hardware to software, to connectivity and cybersecurity. Often, municipalities either have no providers to choose from, or face prohibitively high costs. As the pilot project in Alagoas has demonstrated, municipal managers lack the capacity to digitalise, even when financing is provided by federal and state governments (Ministério da Saúde, 2020^[14]). Setting responsibilities for digitalisation of PHC at municipal level for all of Brazil is likely leading to diseconomies of scale, and creating challenges for small municipalities that do not have the human and technical capacities.

In OECD countries with small subnational government units, such as the Nordic countries, France, the Slovak Republic and Spain, subnational partnerships, joint authorities and outsourcing are frequently used to create scale economies (OECD, 2019^[38]). In the context of digital health, specifically, regional health authorities that bring together smaller jurisdictions and provide shared services to all members are a way to achieve both economies of scale and valuable expertise. For example, in New Zealand, primary health organisations or PHOs, especially larger PHOs like ProCare (which serves 800 000 patients in greater Auckland), offer its member practices a centralised human resource, practice management and procurement service (Varsamis, 2019^[44]).

Established in 2005, the English NHS Shared Business Services (SBS) provide a mix of finance and accounting, procurement, digital innovation services, employment services, innovation and technology services, and change management to hundreds of NHS organisations (NHS Shared Business Services, 2020^[45]). Through a partnership with the Royal Papworth Hospital NHS Foundation Trust, NHS SBS has developed a digital patient booking system that can be rolled out to all NHS organisations (ibid). Also in the United Kingdom, the NHS GP IT Futures programme will create a new environment for the supply of clinical ICT systems to general practice, including a new online NHS Digital Buying Catalogue to find PHC technology systems and services (NHS Digital, 2020^[46]). In Portugal, the Shared Services of the Ministry of Health (SPMS) is the Ministry's central purchasing and ICT authority, providing shared services in the areas of purchasing and logistics, financial services, human resources and ICT. Recently, SPMS added new electronic dynamic purchasing systems for NHS organisations to procure consultancy services in blockchain, artificial intelligence, telemedicine and mHealth from providers that have been certified (SPMS, 2020^[47]).

Brazil could look to these examples from New Zealand, England and Portugal for a new model of shared services for digital PHC. Given the geography, population and population density of Brazil, shared services and responsibilities could be aggregated at the level of the federal government, state governments or even health macro-regions and regions, depending on the specific service being considered. For example, dynamic purchasing systems can be set up by the federal government, building on the experiences of the *Co-ordenação-Geral de Tecnologia da Informação e Comunicações* (CGTIC), which already plans, co-ordinates, supervises and guides ICT investments for the federal government. For smaller states (like Roraima, Amapá and Acre), health macro-regions and bigger health regions could then set up agencies for shared services, providing specialised human resources, purchasing and management services for digital PHC to municipalities.

There are three important policy levers to carry out this regionalisation of competencies for digital PHC:

- First, it is crucial that standards and interoperability remain a central federal responsibility. This does not mean that these should be developed by only the federal government, but it does mean

that states and municipalities should participate in this process rather than lead it. This would ensure that whatever systems are procured by states and municipalities, the ability to share data and patient mobility remain possible throughout Brazil.

- Second, these new mandates need to be adequately funded and responsibilities be clearly set. This would guarantee that the new shared services agencies are both able to execute their responsibilities and accountable for the results.
- Third, local leadership, community participation and local ownership remain essential (OECD, 2019^[38]). A new model of shared services needs to have the buy-in and ownership of the municipalities and PHC units it is supposed to serve.

Larger and smarter investments are needed to deliver on ambitious digital health strategies

Digitalisation requires investments in infrastructure and connectivity (e.g. high speed broadband), in hardware (e.g. computers), software (e.g. EHR systems), specialised human resources (ICT specialists), user training (e.g. digital literacy of professionals and patients), and importantly, though often neglected, change management and implementation. Brazil has invested significantly in connectivity (e.g. *GESAC*) and software (e.g. *e-SUS AB*), as well as providing support for hardware and training (e.g. *Informatiza APS*). However, as seen in previous sections, substantial numbers of PHC units and teams, as well as citizens, remain disconnected, and thus unable to benefit from a digital transformation of PHC. The vision set out in ESD28 and previous digital health strategies requires both larger and smarter investments, tying financial support to effective and equitable use.

The International Telecommunication Union has recently estimated that connecting every Brazilian to the Internet would cost USD 11.1 billion (ITU, 2020^[48]). Brazil's Connect SUS programme is likely to involve total investments of USD 0.77-1.16 billion (BRL 4-6 billion) over five years (OECD, 2020^[11]). In 2019, according to the World Bank Group, Brazil had a population of 211 million and a gross domestic product (GDP) of USD 1.84 trillion. By comparison, in Sweden, a country that in 2019 had just over 10.2 million inhabitants and a GDP of USD 530 billion, the country's 21 regions invest around USD 1.2 billion annually in health care ICT (U.S. Department of Commerce, 2020^[49]). Only 1% of Swedish drug prescriptions are issued in paper format (ibid).

As previously discussed, Brazil instituted an exceptional incentive for municipalities and states digitalise PHC units, budgeting over USD 83.8 million (BRL 432 million) for 2020 (Diário Oficial da União, 2020^[50]). In the United States, the FCC's Emergency Broadband Benefit is a USD 3.2 billion programme to support households struggling to pay for internet service during the pandemic, providing monthly discounts towards broadband service, as well as one-time discounts to purchase equipment like laptops or tablets, with higher discounts on Tribal lands (FCC, 2021^[51]). The FCC's COVID-19 Telehealth Program provides USD 200 million in immediate support to nonprofit and public eligible health care providers responding to the COVID-19 pandemic by fully funding their telecommunications services, information services, and devices necessary to provide critical connected care services (FCC, 2021^[52]).

While it is challenging to compare investments across different geographies, populations and economies, it is helpful to place financial commitments across countries in the contexts of their digital health strategies. The Brazilian ESD28 sets out an ambitious agenda, including milestones for the achievement of certain capabilities (e.g. ePrescribing by December 2022). Looking to the order of magnitude of investments in countries that have already achieved these milestones is a practical way to assess whether current investments are sufficient. As an example, in the United Kingdom, a USD 33.3 million (GBP 26 million) national fund has been set up to have all NHS doctors, nurses and other clinical staff on e-rostering systems by 2021 (GOV.UK, 2021^[53]). E-rostering, which allows clinicians to select shifts digitally, providing them with more flexibility and more efficient work schedules, is just one capability.

Investments should also be comprehensive in their scope. It is often the case that one of the most important steps in implementing new technologies – change management and support for implementation – is not

appropriately funded (Varsamis, 2019^[44]). As the pilot project in Alagoas illustrates, it is possible to have buy-in from municipalities, provide funding, and still have limited adoption. While support in the form of capacity building is undoubtedly important, it might need to be combined with financial support for lost productivity and service disruption during periods of change and adaptation (Bakhai et al., 2019^[54]).

It is not just the magnitude of investments that matters. Conditions for accessing these funds by the beneficiaries can promote, or hinder, the adoption and everyday use of digital technologies. Tying financial support to the use of specific functionalities or programmes can be a way to target investments and increase their effectiveness. The *Informatiza APS* programme already does this by conditioning eligibility for financial support to the reporting of data to the *SISAB*, but it can potentially go further. The Australian Practice Incentives Program (PIP) eHealth Incentive encourages practices to adopt key digital services, from patient records to secure messaging capabilities (to communicate with both patients and other practices), to ePrescribing, among others (Australian Government, 2016^[55]).

Finally, while a digital transformation of PHC has enormous potential (see Box 6.2), there is little high-quality evidence regarding recent digital-first PHC models, such as the one being implemented in the United Kingdom (Rodgers et al., 2019^[56]). It is important that funds be made available for continuous monitoring and evaluation of digitalisation in PHC in Brazil.

Box 6.8. Key recommendations for a co-ordinated well-funded digital PHC

Set clear well-funded mandates for digitalisation at the right levels of government

- Consider aggregating shared services and responsibilities related to digitalisation of PHC at less decentralised levels. Build on experiences of the *CGTIC* to explore setting up dynamic purchasing systems at federal level, with appropriately funded agencies at state or regional level providing specialised human resources, purchasing and management services for digital PHC to municipalities.

Make larger and smarter investments to promote baseline digital PHC services for all

- Review magnitude of investments needed to deliver on vision of ESD28, including funding for change management, capacity building, and monitoring and evaluation. Tie financial support to specific milestones in the adoption and use of digital PHC functionalities, starting with ePrescribing, referral management, appointment scheduling and patient access to EHR.

Note: see also recommendations in *Making decentralisation work: A handbook for policy makers* (OECD, 2019^[38]) and in the *OECD Digital Government Review of Brazil: Towards the Digital Transformation of the Public Sector* (OECD, 2018^[11]).

6.5. Conclusion

Over the last decades, Brazil has been making impressive progress towards a digital transformation of its public sector, and is aiming to offer all of its more than 3 000 public services via digital channels by the end of 2022 as part of its Digital Government Strategy 2020-22. The country has made significant inroads also in building a foundation for digital PHC, with successive ambitious and comprehensive digital health strategies, coupled with vital investments in hardware, interoperability and skills. However, progress towards effective use of digital technologies in PHC has been slow, and fundamentally unequal. Thousands of PHC units have no computers or Internet access. Only 17% of PHC units have speeds over 10 Mbps. Around 18 000 health care facilities (or 18% of all facilities) do not have an EHR system.

The COVID-19 pandemic has likely accelerated the adoption of digital technologies but its true impact remains unclear for now.

At the same time, and despite significant progress in recent years, millions of Brazilians are still digitally excluded. More than two in three Internet users aged 16 or over used electronic government services in 2019, and the share of individuals using the Internet to look for health information more than doubled between 2010 and 2019. However, nearly 47 million Brazilians did not use the Internet in 2019. Digital divides are also clearly associated with social inequalities, with older Brazilians, those from rural areas, those with lower incomes and lower education levels all less likely to use the Internet, and more likely to use mobile phones only. There is thus a serious risk that a digital transformation of PHC will only compound existing social divides.

Against this backdrop, Brazil's decentralised government creates challenges to digitalising PHC. The country's 26 states, one federal district, well over 5 000 municipalities (around 44% of which had less than 10 000 inhabitants), and 438 health regions are very heterogeneous in their demographics, socio-economic indicators and governance structures. The municipality of Serra da Saudade, with around 800 inhabitants, has the same responsibilities to manage the digitalisation of PHC as the municipality of São Paulo, with its over 12 million inhabitants. There are significant risks for the country's ambitious strategy for digital health, including insufficient and geographically imbalanced resource capacity, scarce funding, too broad of an implementation focus, and unavailability of required infrastructure.

Brazil has established a solid basis on which to build an effective, equitable and efficient digital PHC. But, to do so, it must tackle three sets of interconnected challenges: digitalisation of all PHC units and teams; digital inclusion for all Brazilian citizens, especially those that are most vulnerable; and a governance structure that is fit to deliver a digital transformation of PHC, by setting clear well-funded mandates at the right levels of government.

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Notes

¹ Here and throughout this chapter, an exchange rate of BRL 5.155/USD for the year 2020 from <https://stats.oecd.org>.

² The 2020 Brazilian Criteria for Economic Classification (*Critério de Classificação Econômica Brasil*) defines six classes (A, B1, B2, C1, C2, DE) of socio-economic stratification based on a system of points that aggregates survey data on wealth, living conditions, education and income. Category A represents a higher socio-economic status.

³ See also Table 3.1. in (OECD, 2020_[11]).

OECD Reviews of Health Systems

Primary Health Care in Brazil

Primary health care in Brazil is well-organised, the result of sustained commitment to providing high quality primary health care for the whole population. Brazil has implemented a set of reforms over the past decades to improve the distribution of doctors, develop new forms of service organisation, introduce new financing models, and implement a range of quality improvement initiatives. This review uses internationally recognised indicators and policy frameworks to examine the performance of primary health care in Brazil. While the review points to notable successes, Brazil continues to face challenges as its population ages, risk factors such as obesity are on the rise, and emerging pandemic threats require resilience and adaptability. The report points to key actions that Brazil should consider in the coming years to strengthen performance of primary health care, especially screening and prevention for major non-communicable diseases, improve quality of primary health care provision, address workforce shortages and pursue a digital transformation. A companion publication with a health system review of Brazil examines the main challenges and approaches needed to improve the performance of the Brazilian health system.



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1. Executive Summary

Approval is sought for a **two-year £523,432** (£261,716 per annum) **Community Health & Wellbeing Worker (CHWW) pathfinder** in West Wales, representing a targeted system intervention to reduce avoidable demand on health and care services by addressing the underlying social, economic and environmental drivers of poor health and wellbeing.

The pathfinder will establish an initial CHWW team linked to the Ty Elli Practice Group as a first implementation site, forming part of a wider West Wales model across Carmarthenshire and Pembrokeshire.

The programme is expected to contribute to measurable system and population benefits, including:

- Reduction in non-medical GP consultations
- Reduction in unscheduled care interactions
- Improved wellbeing and reduced loneliness
- Improved self-management of long-term conditions
- Optimal access to benefits
- Increased childhood and adult immunisation

Unlike traditional service expansion, this proposal does not add capacity into existing reactive pathways. Instead, it introduces a proactive, community-based mechanism that reduces demand at source by identifying and addressing unmet need earlier.

Health and wellbeing outcomes are shaped not only by healthcare access but by a complex interaction of social, economic and environmental factors, including poverty, social isolation, unemployment, education, housing and community cohesion.

Traditional health services often struggle to address these wider determinants because they operate primarily through clinical pathways and referral-based models. In Hywel Dda, an estimated 460,000 GP consultations per year relate to non-medical need. Even a modest reduction in this demand would release significant clinical capacity and reduce pressure across primary and urgent care services.

This proposal developed by the Centre for Social Innovation at the request of Hywel Dda Public Health directly addresses this pressure by shifting the model of care upstream, introducing proactive, household-based engagement that identifies need earlier, builds trust within communities, and supports sustained interaction with a coordinated coalition of services.

The Community Health & Wellbeing Worker (CHWW) model offers a proven, preventative and community-based approach: a community-embedded workforce focused on proactive engagement, early identification of need and connection to existing services.

In West Wales, this model is further strengthened through partnership with the Centre for Social Innovation (CfSI), by applying a whole-household approach that addresses interconnected issues including housing, education, employment and social isolation, with the aim of building long-term resilience and reducing reliance on reactive public services.

The model has strong international precedent through the Brazilian Family Health Strategy, with over 300,000 workers supporting approximately 70% of the population and has been successfully adapted within the UK through programmes such as those in Westminster and Cornwall.

Evidence from these programmes demonstrates:

- Improved engagement with preventative services,
- Increased uptake of vaccination and screening,
- Reductions in demand on primary care services.

The proposed West Wales model will deploy four CHWWs in Llanelli through established voluntary sector partners, working as part of a coordinated, multi-agency coalition spanning primary care, public health, local authorities, education provision, voluntary sector partners and community assets, reflecting the reality that the drivers of demand do not sit within a single service boundary.

Importantly, the model introduces a proactive neighbourhood-based outreach function that complements and strengthens existing services rather than duplicating them, improving engagement, continuity and outcomes for individuals with complex or unmet needs.

The programme is supported by a nationally established infrastructure through the National Association of Primary Care (NAPC), providing training, digital tools and evaluation capability, reducing implementation risk and enabling rapid mobilisation.

This pathfinder will test how the model can be effectively delivered within the West Wales context and generate robust evaluation evidence to inform future commissioning decisions.

Approval is therefore sought to implement a two-year, £523,432 pathfinder that will test a scalable, evidence-based model capable of reducing system demand, improving population outcomes and strengthening the long-term sustainability of health and care services across West Wales.

The scale and nature of these challenges within West Wales are illustrated below.

CHALLENGES facing Health & Wellbeing in West Wales

Demographic change, health inequalities and rural access challenges are increasing pressure on health, social and community care services across West Wales.

LIVING LONGER WITH COMPLEX NEEDS



1 in 4
residents
aged **65+**

85+ population projected to
increase **28%** by **2030**



RURAL COMMUNITIES

Large rural geography
with **dispersed**
communities creates
challenges for **service**
access and delivery

9+ years



Up to difference in life
expectancy between the
most and least deprived
communities

HEALTH INEQUALITIES



DEMAND
~460,000

GP consultations
per year relate
to non-medical need

Rising demand for health,
care and community services
driven by **ageing population**
and increasing prevalence of
long term conditions



WIDER DETERMINANTS OF HEALTH

Key drivers:

- **poverty** and financial **insecurity**
- **social isolation**
- housing **quality**
- **unemployment** and economic inactivity

POPULATION

388,000 residents



Carmarthenshire ~ 49%
Pembrokeshire ~ 33%
Ceredigion ~ 18%

These challenges highlight the need for **community-based** approaches that build **trust**, strengthen **prevention**, improve **engagement** with services and **support** earlier intervention

Data Sources: Office for National Statistics (ONS) Mid-Year Population Estimates; ONS Subnational Population Projections (2020-based); StatsWales; Public Health Wales Observatory; Welsh Index of Multiple Deprivation (Welsh Government); Hywel Dda Population Needs Assessment

2. Strategic Case

2.1. The Case for Change

Across Wales there is growing recognition that the current health and social care system is under increasing pressure from rising demand, workforce constraints and widening health inequalities. Many services are currently configured to respond to crisis or deterioration, rather than supporting individuals and families earlier in their health and wellbeing journeys.

Key challenges in Wales include:

- High levels of deprivation in many communities
- Rurality and transport barriers
- Social isolation and loneliness
- Mental health pressures and long waiting lists
- An ageing population with increasing multimorbidity
- Cultural and societal risk factors

The World Health Organisation has identified that factors such as poverty, poor housing, social isolation, low health literacy and chronic stress exert a far greater influence on population health, driving long term conditions, multimorbidity and unscheduled care. Addressing these upstream drivers is therefore critical to improving outcomes, reducing inequalities and relieving pressure on health services.

2.1.1. Determinants of Health

Fixed

1. *Biological Factors* (we are largely born with). Examples include:

- Age
- Sex
- Genetic inheritance

Modifiable

2. *Individual Lifestyle Factors* (How people live day to day) - often the focus of traditional health interventions but are strongly shaped by wider conditions. Examples include:

- Smoking and alcohol use
- Diet and nutrition
- Physical activity
- Sleep patterns
- Medication adherence

Such lifestyle “choices” are constrained by income, housing, employment, education, and stress.

3. *Social and Community Networks* (The relationships around people). Social connection is a powerful protective factor for health, examples include:

- Family and household support
- Social isolation or loneliness
- Community cohesion
- Informal carers and peer support
- Trust in services and institutions

The health impact of this factor includes links to mental health outcomes, influencing self-care and health-seeking behaviours.

4. *Living and Working Conditions* (The core drivers of health inequalities). Often has the largest impact on population health. Examples include:

- Housing quality (damp, cold, overcrowding)
- Employment status and job security
- Income and financial stress
- Education and skills
- Transport access
- Digital access and literacy
- Access to green space and safe environments

These aspects typically explain why health inequalities persist even when healthcare access is universal.

5. *Wider Socio Economic, Cultural, and Environmental Conditions* (The outer system context) and shape the other determinants. These include:

- Poverty and inequality
- Economic policy and welfare systems
- Education systems
- Urban planning and transport policy
- Climate and environmental quality
- Cultural norms and discrimination
- National and local policy decisions

This is where prevention has the highest leverage, but often the longest time horizon.

These drivers sit largely outside traditional clinical pathways yet account for a significant proportion of repeat demand, placing sustained pressure on primary care, urgent care and wider services. GPs often act as the default point of access when wider support systems are unavailable, resulting in avoidable demand and pressure within primary care.

Evidence consistently shows that clinical services alone cannot resolve these challenges. A survey of GPs¹ (Citizens Advice 2015) suggested that significant proportion of demand presented to GPs relates to social, psychological, and practical issues that fall outside traditional medical responses, yet strongly influence health outcomes:

¹ [CitizensAdvice_AVeryGeneralPractice_May2015.pdf](https://www.citizensadvice.org.uk/cach3.com/Global/CitizensAdvice/Public%20services%20publications/CitizensAdvice_AVeryGeneralPractice_May2015.pdf)
https://www.citizensadvice.org.uk/cach3.com/Global/CitizensAdvice/Public%20services%20publications/CitizensAdvice_AVeryGeneralPractice_May2015.pdf

Around 20% of GP consultation time is spent on non-medical issues² (Hood et al, 2015). These issues most commonly include:

- Housing problems
- Benefits and financial stress
- Employment issues
- Relationship and family stress
- Loneliness and social isolation

The estimated total population for Hywel Dda University Health Board is 385,600. The NHS Strategy Unit states that the average consultation rate in UK General practice is 5.5-6.5 consultations per person per annum, with rural and older populations (like HDUHB) tend to be at the higher end of this range.

Illustration for Hywel Dda University Health Board

386,000 population x6 consultations = 2,316,000 total consultations of which 20% (463,000) are likely being utilised for non-medical issues.

GPs reported these issues reduce time available for clinical care and frequently re-present because they are not medically resolvable.

Welsh Government and Public Health Wales policy recognise that most health outcomes are driven by social, economic and environmental factors, not healthcare alone. A non-medical approach, “Social Prescribing” has been developing since 2016, with the aim to support holistic health, wellbeing and social needs. For example, it includes connecting people to groups and services within a community to reduce loneliness, improve mental health and address long-term conditions.

Key timeline for social prescribing in Wales:

- Government commitment (2016)
- Wales Social Prescribing Research Network (2018)
- National Framework for Social Prescribing (2023)

2.1.2 From determinants to delivery: Social Prescribing and CHWWs

The determinants of health outlined above demonstrate that a significant proportion of demand within health services is driven by non-medical need. These factors – including housing, financial insecurity, social isolation and low confidence – cannot be resolved through clinical intervention alone, yet frequently present within primary care and urgent care settings.

In Wales, social prescribing has emerged as an important mechanism to address these wider determinants. It provides a structured pathway to identify non-medical needs and connect individuals

² Hood, C. M., K. P. Gennuso, G. R. Swain, and B. B. Catlin. 2016. County health rankings: Relationships between determinant factors and health outcomes. *American Journal of Preventive Medicine* 50(2):129-135. <https://doi.org/10.1016/j.amepre.2015.08.024>

to community-based support. This includes services such as welfare advice, community groups, employment support and mental wellbeing services.

However, social prescribing is primarily a referral-based model. It typically relies on individuals presenting to services, after which they are signposted or referred onwards. While effective, this approach can be limited where:

- needs are complex or multiple
- individuals lack confidence or trust to engage
- people disengage after referral
- programmes are time or use limited
- issues fluctuate or persist over time

The Community Health & Wellbeing Worker (CHWW) model strengthens this approach by introducing a proactive, relationship-based workforce. Rather than waiting for individuals to present, CHWWs engage directly with households and communities, building trust, identifying need earlier and supporting sustained engagement with a range of services.

CHWWs therefore provide a complementary function to social prescribing, addressing its key limitations by:

- proactively identifying unmet need before escalation
- building longitudinal, trust-based relationships with residents
- supporting individuals to navigate services, rather than simply referring
- maintaining continuity of support for people with complex or changing needs
- encouraging individual connections and community cohesion

This distinction is critical. Social prescribing identifies support and answers the question: “**What services are available?**” CHWWs enable the change and answers: “**Who will walk alongside this person to achieve change?**”. By combining both approaches, the model moves from signposting to sustained behaviour change and improved outcomes.

By combining the strengths of social prescribing with proactive community outreach, the CHWW model creates a more complete and effective approach to addressing the social determinants of health and reducing avoidable demand on health and care services. Therefore, it is not an alternative to existing services, but a critical enhancement that strengthens their effectiveness and potentially reduces the long-term pressures on those services.

Taken together, these factors create a strong case for a proactive, relationship-based model in West Wales. CHWWs provide that function by building trusted relationships with households, identifying unmet need earlier and connecting people to the wider coalition of services that can support health, wellbeing and longer-term resilience. This is particularly important in West Wales, where rurality, isolation, deprivation and fragmented access to services can make referral-based models alone insufficient.

This integrated approach reflects growing evidence that relationship-based community outreach is one of the most effective ways of reaching individuals who are least likely to engage with services until needs escalate. Many existing services operate through referral-based pathways, meaning individuals

typically receive support only once needs have already been identified, often during periods of crisis or deterioration.

CHWWs address this gap by providing proactive engagement within neighbourhoods, identifying unmet needs earlier and strengthening connections between residents and existing services. The model also aligns strongly with the principles of *A Healthier Wales* and the *Wellbeing of Future Generations Act*, supporting prevention, integration and community-based care.

Social Innovation Approach (CfSI Contribution)

The model is further enhanced through collaboration with the Centre for Social Innovation (CfSI), which brings a distinct approach focused on addressing systemic barriers and enabling long-term behavioural and societal change.

While the CHWW model is already proven in improving engagement and reducing demand, the CfSI approach strengthens this by:

- Taking a whole-household perspective rather than focusing solely on individual needs,
- Addressing interconnected issues such as housing, education, employment and social isolation alongside health,
- Supporting behaviour change through sustained, trust-based relationships within communities,
- Enabling communities to become more connected and resilient, reducing long-term reliance on public services.

This approach represents a shift from traditional service models, where individuals are expected to identify problems and seek help, towards a proactive model where support is brought into the home and tailored to the realities of people's lives. By combining proven CHWW delivery with social innovation principles, the programme aims not only to improve individual outcomes but to test how community-based models can contribute to wider system transformation and long-term population health improvement.

The proposed model has been intentionally designed as a coalition-based approach rather than a standalone service. It brings together primary care, public health, local authorities, voluntary and community sector partners, academic evaluation, and wider community assets such as housing, education, training and employment support. This coalition model reflects the reality that the barriers affecting households do not sit neatly within one service boundary and therefore require a coordinated, relationship-based response across the wider system.

2.2. Evidence and Precedent

The CHWW model is supported by substantial international precedent and a growing body of evidence from UK implementation demonstrating its effectiveness in strengthening prevention and community engagement. There are currently 24 live CHWW sites across England, employing approximately 192 workers, supported through the NAPC national programme. This demonstrates that the model is not experimental, but a proven, scalable intervention adapted successfully across multiple contexts.

2.2.1. International Origins – Brazil

The CHWW model originates from Brazil's Family Health Strategy, introduced in the early 1990s as part of major national health system reform designed to strengthen prevention and community-based care.

The programme embeds community health workers within neighbourhoods to provide proactive household engagement, health promotion and connection to primary care services.

The programme now includes:

- over 300,000 community health workers
- coverage of approximately 70% of the Brazilian population

This represents one of the largest community-based public health workforces in the world.

Evidence from Brazil demonstrates significant improvements in population health outcomes, including:

- increased vaccination uptake,
- improvements in maternal and child health,
- improvements in chronic disease management,
- reductions in preventable hospital admissions.

The Brazilian experience demonstrates that community-based health workers can play a transformative role in strengthening prevention and improving access to care at scale.

2.2.2. UK Implementation – Westminster

The first UK adaptation of the model was introduced in Westminster in 2021 with support from the National Association of Primary Care. This programme applied the core principles of the Brazilian model within a UK primary care context, focusing on proactive household engagement in areas experiencing high levels of health inequality.

Peer-reviewed research (Junghans, 2023) evaluating the Westminster programme demonstrated significant improvements in engagement with preventative health services.

Households receiving visits from Community Health & Wellbeing Workers experienced:

- 40% higher uptake of prevention opportunities,
- 47% higher vaccination uptake,
- 82% higher participation in screening and NHS health checks,
- 7.4% reduction in GP consultations.

These outcomes were achieved in communities with historically low engagement with preventative health services. The findings demonstrate that proactive, relationship-based outreach can significantly improve preventative health engagement while reducing demand on primary care services.

Importantly, the model works because it combines health promotion with broader social support, recognising that barriers to health engagement often arise from wider social circumstances rather than lack of clinical services.

2.2.3. Cornwall Programme

The most relevant UK comparator for West Wales is the programme delivered in Cornwall through Volunteer Cornwall.

The programme initially launched with 10 Community Health & Wellbeing Workers and has since expanded significantly. It now includes approximately 55 workers operating across the county, working through voluntary sector organisations and integrated with primary care and community services. Each

worker typically supports around 120 households, building long-term relationships with residents and helping them access health and wellbeing services.

Cornwall shares many characteristics with West Wales, including:

- rural and dispersed communities,
- pockets of deprivation and health inequality,
- strong voluntary sector infrastructure,
- close working between health, local government and community organisations.

These similarities make Cornwall a particularly relevant example of how the CHWW model can be successfully implemented in rural and mixed population settings. Learning from the Cornwall programme also demonstrates the effectiveness of delivering the workforce through voluntary sector organisations, which are often best placed to build trust and engage with communities.

2.2.4. Relevance to West Wales

Taken together, the evidence from Brazil, Westminster and Cornwall demonstrates that the Community Health & Wellbeing Worker model can:

- strengthen prevention and early intervention,
- improve engagement with health and wellbeing services,
- reach individuals who may not otherwise access support,
- reduce demand on reactive health services.

The success of the model across both urban and rural contexts provides a strong foundation for testing the approach within West Wales. The proposed deployment will therefore build on international evidence and UK learning to assess how the model can be effectively adapted and delivered within the Welsh health and social care context.

2.2.5. Workforce Sustainability and Local Economic Impact

A key strength of the Community Health & Wellbeing Worker (CHWW) model is its ability to establish a sustainable, locally rooted workforce that contributes not only to health and wellbeing outcomes, but also to skills development and economic participation within communities.

The proposed West Wales model will utilise nationally recognised training pathways through the National Association of Primary Care (NAPC) to ensure rapid mobilisation and consistent workforce capability from the outset.

Building on this, the model is designed to support long-term workforce sustainability through regional capability and national alignment. This includes partnership with the University of Wales Trinity Saint David (UWTSD) to develop locally delivered training provision, alongside early engagement with Health Education and Improvement Wales (HEIW) to explore alignment with emerging national workforce pathways, including the potential development of a pan-Wales apprenticeship model informed by the Therapy Assistant Practitioner (TAP) approach.

This integrated approach positions the CHWW model as both a delivery intervention and a workforce development mechanism, enabling:

- accessible entry routes into health, social care and community-based careers
- workforce development aligned to local population need and system demand

- structured progression pathways into health, social care and specialist community roles
- improved recruitment and retention through locally rooted employment opportunities

Importantly, the model recognises that the CHWW role itself represents a skilled and sustainable career, supporting both progression where desired and long-term role stability where appropriate.

This approach directly aligns with Welsh Government priorities relating to community-based workforce models, economic participation and regional skills development, particularly in rural and underserved areas, and strengthens the case for scalability across Wales.

2.2.6. Strategic Fit

The proposed pathfinder aligns strongly with Welsh policy direction, including:

A Healthier Wales

- prevention and early intervention
- care closer to home
- integrated services across sectors.

Public Health Priorities

- improving access to preventative support
- tackling health inequalities
- strengthening community resilience.

Neighbourhood and Place-Based Care

- supporting collaboration between health, local government and voluntary sector partners.

2.2.6 Innovation and Future Development

The programme will also explore opportunities to enhance the CHWW model through technology and innovation, including the potential future use of digital tools such as remote monitoring, wearables or home-based sensors where appropriate.

These elements will not form part of the core pathfinder delivery but will be considered as part of the evaluation and future development of the CfSI model, ensuring that innovation is introduced in a controlled and evidence-based way.

3. Benefits

3.1. Benefit of Community Health and Wellbeing programmes

While the most immediate measurable impacts are likely to be seen in primary care demand, wellbeing and urgent care avoidance, the full benefit of the model is wider. The programme is designed to support a coalition of services, with impacts anticipated across health, social care, community resilience, access to entitlements, education, training and employment support.

Primary Care Capacity & Demand (Short–Medium Term)

A. Reduction in GP consultations for non-medical need (Workforce liberation)

- Fewer repeat GP appointments driven by:
 - loneliness
 - stress/anxiety
 - housing/financial problems
 - low-confidence self-management

B. GP time released (Workforce liberation)

- Fewer short, non-clinical consultations
- Improved focus on:
 - complex clinical need
 - multimorbidity
 - continuity

2. Urgent & Emergency Care (Medium Term)

A. Reduced escalation into urgent care

- Earlier intervention prevents:
 - anxiety-driven ED attendances
 - deterioration of LTCs
 - crisis presentations

B. Avoided hospital admissions

- Reduced admissions for ambulatory care sensitive conditions (ACSCs)
- Better adherence to care plans

3. Mental Wellbeing & Psychosocial Outcomes (Short Term)

A. Improved wellbeing scores

What changes

- Improved wellbeing, reduced loneliness and strengthened community connection

B. Reduced reliance on crisis mental health routes

- Fewer GP visits for stress-related symptoms
- Better coping skills and support networks

4. Long-Term Condition Management (Medium Term)

A. Improved engagement & adherence

- Better attendance at:
 - reviews
 - diagnostics

- monitoring
- Improved self-management behaviour

B. Slower deterioration trajectories

- Reduced volatility in conditions with strong social drivers

5. Prevention & Early Intervention (Medium–Long Term)

A. Earlier risk detection

- Issues identified **before they become clinical**
- Safeguarding and welfare risks surfaced early

B. Falls and frailty prevention

- Improved confidence and mobility
- Reduced fear-driven inactivity

6. Workforce Experience & Sustainability (Short Term)

A. GP and practice team experience

- Reduced moral distress from “unfixable” problems
- Improved sense of system support

B. Reduced “failure demand”

- Fewer repeated contacts for same unresolved issues

7. Population Insight & System Intelligence

A. Better understanding of unmet need

What changes

- Rich, real-world insight into:
 - housing issues
 - poverty markers
 - isolation patterns
 - societal shifts

3.2. Alignment of benefit to Hywel Dda University Health Board priorities

HDUHB Priority	CHWW Contribution	Measurable Alignment
Move from a sickness service to a wellness	• Early identification of social and wellbeing risks	• Reduced crisis presentations • Improved wellbeing scores

and prevention-focused system	• Prevention of escalation into clinical crises • Behaviour change support in real-world context	• Earlier support before service thresholds
Sustainable primary care and reduced unplanned demand	• Diverts non-medical demand from GPs • Reduces repeat, unresolved attendances • Releases GP capacity for complex clinical need	• Reduction in GP attendances for CHWW cohort • GP hours released (capacity equivalent) • Reduced ED/urgent contacts over time
Tackling inequalities across rurality and deprivation	• Proactive outreach into hard-to-reach households • Support for housing, poverty, isolation, digital exclusion • Relationship-based engagement in deprived areas	• Improved engagement in previously under-engaged groups • Reduced inequality in access to support • Population insight to target investment
Better support for frailty, multimorbidity and Long-Term Conditions	• Supports adherence and attendance • Reduces volatility and deterioration • Addresses social drivers of LTC outcomes	• Improved LTC review attendance • Reduced exacerbation-driven contacts • Fewer preventable admissions
Whole-system integration with local authorities and voluntary sector	• Acts as connective tissue between sectors • Strengthens social prescribing effectiveness • Provides local intelligence to clusters	• Increased joint working activity • Improved MDT flow • Better use of community assets

3.3. Benefit Realisation

Outcome measurement will utilise validated tools such as MYCaW (Measure Yourself Concerns and Wellbeing), a widely used patient-reported outcome measure developed in primary care and community settings (Paterson et al., 2008; Polley et al., 2017), supported through the National Association of Primary Care (NAPC) national offer. This ensures a consistent, standardised approach to capturing patient-reported outcomes alongside system-level metrics, strengthening the robustness and comparability of evaluation findings. In addition to health service impacts, the evaluation will assess wider coalition outcomes including household resilience, community connection, access to entitlements, engagement with training or employment support, and the effectiveness of whole-household approaches in sustaining change.

Dimension	Benefit	KPI	Measure	Data Source	Timeframe
Primary Care Capacity & Demand	Reduced non-medical GP consultations	% reduction in GP attendances for CHWW cohort	GP appointments per patient (pre vs post)	GP practice systems (EMIS/Vision)	3–6 months
	Reduced repeat attendance	% of frequent attenders (top decile)	Number of contacts per frequent attender	Practice BI / audit	6–12 months
	GP capacity released	GP hours released (capacity equivalent)	Appointments avoided × 10 mins	Derived metric	6–12 months
Mental Wellbeing & Psychosocial Outcomes	Improved wellbeing	Change in MYCaW / ONS-4 score	Pre- / post- engagement score	CHWW outcome collection	6–12 weeks
	Reduced loneliness	% reporting reduced loneliness	Loneliness scale (e.g. UCLA-3)	CHWW assessment	3–6 months
	Reduced stress-related demand	Reduction in GP anxiety/stress codes	Rate per 1,000	GP systems	6–12 months
Urgent & Emergency Care Avoidance	Reduced crises use	ED attendances per CHWW cohort	Rate per 1,000	ED dataset	6–18 months
	Reduced conveyances	Ambulance conveyances avoided	Call-out/conveyance rate	WAST data	6–18 months
	ACSC admission prevention	Emergency admissions for ACSCs	COPD, HF, falls	PEDW	12–24 months
Long-Term Condition Management	Improved engagement	DNA rate for LTC reviews	% DNAs	GP systems	6–12 months
	Improved self-management	Attendance at monitoring	HbA1c / BP reviews attended	Audit	6–12 months
	Fewer exacerbations	Unplanned contacts	Contacts per patient	GP / ED	12–24 months
Prevention, Frailty & Falls	Reduced falls risk	Falls per CHWW older cohort	Rate per 1,000	Community datasets	12–24 months
	Early frailty identification	% identified pre-crisis	Frailty flags raised	MDT records	6–12 months
	Delayed escalation	Time to crisis	Months to first urgent contact	Longitudinal review	12–24 months
Workforce Experience & System Sustainability	Improved GP experience	Qualitative feedback	Survey / narrative	GP cluster feedback	3–6 months
	Reduced failure demand	Repeat attendance rate	Same-issue recontacts	Practice audit	6–12 months

	Better MDT functioning	Perceived value of CHWW	MDT feedback	MDT survey	6 months
Population Insight & Place-Based Intelligence	Better insight into unmet need	Thematic issue reporting	Housing, loneliness, poverty themes	CHWW reporting	Ongoing
	Targeted prevention	Actions informed by insight	Changes to local interventions	Programme review	12 months
	Improved cross-sector working	Joint actions with LA/VCSE	Number of joint interventions	Programme log	Ongoing
Other	Increased access to entitlements	% of eligible households with a completed benefits check	Benefit claims submitted monthly	CHWW assessment	6–12 months
	Increased childhood coverage	% of children with MMR dose 1 by 2nd birthday and % with 2 doses by age 5	MMR1 coverage at age 2 and MMR2 coverage at age 5	PHW / WIS / Practice baseline	12-24 months
	Adult Flu uptake	% vaccinated among target population e.g. over 65s and those with LTCs	Flu vaccinations	PHW / WIS / Practice baseline	12-24 months

4. Economic Case

4.1. Overview

The purpose of the Economic Case is to demonstrate that the proposed Community Health & Wellbeing Worker (CHWW) deployment represents the most effective and efficient use of public resources to address the identified gap in proactive community engagement and prevention support.

The analysis considers how best to strengthen early identification of need and improve engagement with health, social care and community services across West Wales.

The options appraisal therefore examines alternative approaches to improving preventative engagement and reducing demand on reactive services, before identifying the preferred option.

4.2. Options Considered

Three delivery options were considered.

Option 1 – Do Nothing

Under this option, no new intervention would be introduced, and existing services would continue to operate within their current models. This option would avoid additional expenditure in the short term but would fail to address the underlying system challenges identified in the Strategic Case.

Existing services primarily operate through referral-based pathways, meaning support is typically provided only once individuals have already presented to services or experienced deterioration in their circumstances. As a result, opportunities for earlier intervention and prevention are often missed, contributing to avoidable pressure on primary care, urgent care and social care services.

Maintaining the status quo would therefore perpetuate:

- reactive service models,
- increasing demand on healthcare services,
- persistent health and social inequalities,
- missed opportunities for preventative engagement.
- disengagement and dissatisfaction.

This option would not contribute to system transformation or prevention objectives and is therefore not considered a viable long-term approach.

Option 2 – Expand Existing Referral-Based Services

This option would involve increasing capacity within existing services such as community connectors or social prescribing roles. These services play an important role in supporting individuals to access community-based wellbeing support once a need has been identified. However, expanding these services alone would not address the fundamental gap identified in the Strategic Case.

These roles typically rely on referrals from primary care or other professionals, meaning they engage individuals only after their engagement or when needs have already been recognised. While expanding these roles may improve access to support for individuals already within the system, it would not introduce the proactive neighbourhood-based outreach function that is central to the CHWW model.

As a result, this option would have limited ability to:

- identify unmet need earlier,
- engage residents who are not currently accessing services,
- build trusted relationships within communities.

Although beneficial, this option would primarily strengthen reactive pathways rather than delivering a preventative community engagement model.

Option 3 – Community Health & Wellbeing Worker Model (Preferred Option)

This option introduces a dedicated workforce focused on proactive community engagement, early identification of need and connection to existing services.

The CHWW model differs fundamentally from traditional service models. Rather than waiting for individuals to present to services, Community Health & Wellbeing Workers actively engage with residents within neighbourhoods, building trusted relationships and identifying a wide spectrum of emerging needs before they escalate.

This proactive approach enables the workforce to:

- reach individuals who may not otherwise engage with services,
- support earlier intervention across health, social care and community support,

- strengthen connections between residents and existing services,
- address wider determinants of health including employment, education and social isolation.
- encourage and strengthen community cohesion.

International evidence from the Brazilian Family Health Strategy, together with UK experience such as Westminster and Cornwall, demonstrates that this model can increase engagement with preventative health services while reducing demand on reactive healthcare provision.

4.3. Value for Money Considerations

The proposed CHWW deployment represents a high-value, preventative investment when compared with the costs associated with reactive healthcare interventions.

For example:

- a GP consultation typically costs approximately £40³,
- an emergency department attendance costs approximately £110 - £565,
- an emergency hospital admission may cost between £1,500 and £3,500⁴.

If 5% of the 480 households (24 households) experience one fewer GP consultation per year, this equates to ~£960 in direct cost avoidance (24 × £40)

If 10% of households (48 households) avoid one GP consultation, this equates to ~£1,920 annually

These illustrative figures represent only a small proportion of potential impact and exclude wider system effects such as reduced repeat attendance, avoided escalation, and improved workforce productivity.

More significant value is expected to arise from reduced repeat demand, improved care navigation, and avoided escalation into urgent and emergency care, which will be evaluated through the programme's monitoring framework. This aligns with national learning from NAPC-supported sites, where impact is demonstrated through multi-domain system value rather than a single cost-saving metric.

This approach presents a prudent, evidence-aligned view of value rather than relying on high-impact but less predictable assumptions such as avoided admissions.

4.4. Wider Social and Economic Benefits

In addition to potential health system impacts, the CHWW model supports wider social outcomes by addressing the social determinants of health. These include:

- improved engagement with employment and training opportunities,
- increased participation in community activities,
- reduced social isolation,

³ Source: Curtis & Burns (2023) *Unit Costs of Health and Social Care; NHS England National Cost Collection 2022/23*

⁴ NHS England (2023) *National Cost Collection for the NHS 2022/23*

- improved access to education and youth services.

These wider benefits contribute to broader public sector objectives relating to economic participation, community resilience and social inclusion.

4.5. Preferred Option

Based on the analysis above, Option 3 – the Community Health & Wellbeing Worker model, represents the preferred option.

This option offers the strongest alignment with strategic priorities for prevention, community engagement and early intervention while providing a practical and evidence-informed approach to addressing the identified gap in proactive outreach.

The proposed deployment will allow the model to be tested within the West Wales context and generate robust evidence to inform future commissioning decisions and potential wider adoption.

Option 3 is the only option that introduces a proactive, preventative mechanism into the system, addressing root causes rather than increasing capacity within existing reactive pathways.

5. Commercial Case

5.1. Overview

The Community Health & Wellbeing Worker pathfinder will be delivered through a partnership-based commissioning model involving voluntary sector delivery partners and central programme coordination. This approach reflects the importance of community trust, local intelligence and established voluntary sector networks in successfully implementing the CHWW model. This coalition-based delivery model is a defining feature of the proposal, enabling coordinated, whole-system support that no single organisation could deliver in isolation.

The proposed commercial model has been informed by learning from existing programmes, particularly the Volunteer Cornwall Community Health & Wellbeing Worker programme, which has demonstrated the effectiveness of delivering the workforce through voluntary sector organisations embedded within local communities. This model enables the programme to combine the credibility and reach of community-based organisations with the oversight and consistency provided by central programme coordination.

5.2. Delivery Model

The delivery model is designed to operate across a coalition of partners, combining the reach of voluntary sector organisations, the legitimacy and pathways of primary care, the strategic coordination of public health, the wider support offer of local authorities, and the research and development capability of CfSI and UWTSD.

The programme will adopt a hybrid delivery model, comprising:

- local delivery through third sector organisations,
- central coordination of programme principles, standards and evaluation.

This model has been selected because voluntary sector organisations are uniquely positioned to:

- recruit workers with strong community connections,
- build trusted relationships with residents,
- engage effectively with local community groups and networks,
- navigate the local service landscape.

Community-based delivery is a core feature of successful Community Health Worker models internationally and has been identified as a key factor in building trust and engagement within communities.

Local delivery partners will be identified and appointed based on their ability to deliver the CHWW model effectively within their communities, rather than being limited to a predefined set of organisations. This ensures the model remains flexible, locally responsive and aligned to existing community assets.

Potential delivery partners may include established voluntary sector organisations, housing associations, community anchor organisations or other place-based providers with strong local presence, trusted relationships and the operational capability to support workforce delivery.

Organisations such as Carmarthenshire Association of Voluntary Services (CAVS) and local housing associations represent examples of the type of partner well positioned to deliver this model. However, partner selection will be based on best fit against defined criteria, including:

- demonstrated community reach and trust within target populations,
- experience of delivering community-based or outreach services,
- ability to support safeguarding, supervision and lone working,
- existing relationships across health, social care and community networks,
- capacity to host and operationally manage the workforce.

This approach ensures that delivery is rooted in the strongest available local infrastructure, supports collaboration with existing partners, and enables the model to adapt to local context and opportunities as the programme develops. These organisations will host the Community Health & Wellbeing Workers and provide operational management and supervision.

Additionally, by leveraging existing national training, digital and evaluation support provided through NAPC, this model avoids the need to develop new infrastructure locally.

5.3. Market Capability

The voluntary sector in West Wales has a well-established infrastructure supporting community engagement and partnership working across the health and social care system.

Organisations such as CAVS and PAVS already play a central role in coordinating voluntary and community sector activity across their respective counties.

These organisations provide:

- established relationships with community organisations and residents,

- experience of delivering community-based programmes,
- local knowledge of community needs and assets,
- established governance and safeguarding arrangements.

Utilising these organisations therefore reduces delivery risk by building on existing capability and trusted community networks.

5.4. Commissioning Approach

The programme will utilise grant or service-level agreement (SLA) arrangements with voluntary sector partners to support delivery of the workforce. This approach reflects the collaborative nature of the programme and aligns with existing partnership arrangements between statutory organisations and the voluntary sector.

Service agreements will define:

- workforce numbers and deployment areas,
- responsibilities for recruitment and workforce management,
- safeguarding and governance requirements,
- reporting and monitoring arrangements,
- participation in programme evaluation.

These agreements will ensure clear accountability for delivery while maintaining the flexibility required to respond to local community needs.

5.5. Workforce Development Model

The proposed delivery model includes a phased approach to workforce development, balancing national consistency with long-term local sustainability.

In the initial phase, training and accreditation will be delivered through the National Association of Primary Care (NAPC), providing access to established Level 3 qualifications, structured learning pathways and national best practice. This enables rapid mobilisation of the pathfinder using proven training infrastructure and supports consistency with existing UK delivery models.

In parallel, the programme will work in partnership with the University of Wales Trinity Saint David (UWTSD), to explore the development of a locally delivered training model, building regional capability and reducing reliance on external provision over time. This includes early engagement with Health Education and Improvement Wales (HEIW) to explore alignment with national workforce development priorities and the potential development of a pan-Wales apprenticeship pathway for Community Health & Wellbeing Workers, informed by the Therapy Assistant Practitioner (TAP) model developed by UWTSD and Hywel Dda University Health Board.

This phased approach ensures:

- rapid mobilisation of the pathfinder through existing national infrastructure,
- development of a sustainable local training pipeline,
- alignment with regional workforce and skills strategies.

Over time, this creates the potential for West Wales to act as a regional hub for CHWW training and development, supporting future scale-up and standardisation of the model across Wales.

5.6. Central Programme Coordination

While delivery will be local, the principles and standards of the CHWW model will be coordinated centrally to ensure consistency across delivery areas. Central coordination responsibilities will include:

- defining the core CHWW role and operating model,
- supporting the recruitment process,
- establishing training and workforce development standards,
- coordinating programme evaluation and reporting,
- facilitating shared learning between delivery partners.

This structure ensures that the programme maintains a consistent model of delivery while allowing local partners the flexibility to adapt engagement approaches based on community context.

5.7. Value for Money

Delivering the programme through voluntary sector partners offers several potential efficiencies, including:

- lower organisational overheads compared with statutory delivery,
- access to existing community networks and infrastructure,
- established local recruitment pipelines,
- flexibility in workforce deployment.

These factors contribute to a cost-effective delivery model while ensuring strong community engagement and reducing implementation risk.

5.8. Contract Management

Delivery partners will operate under agreed service-level arrangements that define:

- programme objectives,
- workforce deployment expectations,
- reporting requirements,
- safeguarding standards,
- participation in evaluation activity.

Programme management and oversight will be provided through the governance structure described in the Management Case.

5.9. Future Commercial Considerations

The pathfinder phase will provide an opportunity to test the delivery model and assess its scalability. If successful, future commissioning options may include:

- expansion of the workforce or through additional voluntary sector partners,
- integration within place-based health and wellbeing programmes,
- wider regional or national implementation.
- additional innovations and services targeting community building.

The deployment will therefore provide valuable learning regarding the most effective commissioning and delivery model for Community Health & Wellbeing Workers within Wales.

5.10. National Support Offer

The programme will initially utilise support from the National Association of Primary Care (NAPC), which provides a nationally developed infrastructure to support Community Health & Wellbeing Worker programmes.

The NAPC offer represents a bundled package of support that would otherwise require local development and includes:

- Accredited training pathways (Level 3 Certificate and Apprenticeship), supporting workforce capability and professionalisation – *full details are included in 6.4.1 National Training and Accreditation Offer*,
- A bespoke case management system (Octolink) to enable structured recording of activity, caseload management and reporting,
- Access to validated evaluation tools (e.g. MYCaW) to support consistent measurement of patient-reported outcomes,
- Participation in a national peer network enabling shared learning across CHWW programmes,
- Quarterly learning workshops and an annual national conference, supporting continuous development and dissemination of best practice.

The cost associated with this national support offer is approximately £15,000 per annum.

This represents good value for money, as it:

- avoids the need to develop training, digital infrastructure and evaluation tools locally,
- ensures alignment with established national models and best practice,
- strengthens the robustness of evaluation and evidence generation,
- supports consistency and scalability of the model across Wales.

The inclusion of the NAPC offer therefore reduces delivery risk, accelerates implementation and enhances the ability of the pathfinder to generate meaningful and transferable learning. Without this national infrastructure, equivalent capability would need to be developed locally at significantly higher cost and risk.

6. Financial Case – Indicative Cost Model

6.1. Community Health & Wellbeing Worker Pathfinder (West Wales) Overview

The purpose of the Financial Case is to demonstrate that the proposed Community Health & Wellbeing Worker (CHWW) deployment represents a realistic, proportionate and affordable investment aligned with the scale and objectives of the proposed pathfinder programme.

The pathfinder will deploy four Community Health & Wellbeing Workers (Band 4 equivalent) across Carmarthenshire, delivered through voluntary sector partners and a central programme coordinator.

The total estimated annual programme cost, for 2026/27, is £268,173. All financial modelling, including cost-per-household and value-for-money calculations, is based on this consistent annual figure.

This cost includes workforce salaries, programme management, training, infrastructure and programme coordination. All financial assumptions are aligned to national benchmarks (NAPC modelling and NHS Agenda for Change pay scales) and have been standardised across the document to ensure internal consistency and transparency.

6.2. Workforce Costs

Community Health & Wellbeing Workers are proposed at Band 4 equivalent level, reflecting the level of responsibility required to work independently within communities while navigating across health, social care and community services. The Welsh model reflects a higher-skilled workforce aligned to complexity, safeguarding and multi-agency working.

Typical Band 4 salary range (Agenda for Change): £27,898 - £30,615

For financial planning purposes a midpoint salary has been used to estimate workforce costs.

Salary Assumptions

Item	Cost per annum	Total cost per annum
Salary (Band 4 midpoint)	£27,898	
Employer on costs (NI, pension approx. 25-30%)	£ 7,650	
Total annual employment cost	£38,265	
Total Community Health & Wellbeing Worker x4		£153,024

Total workforce cost: approximately £153,024 per year

6.2.2 Banding rationale

Banding for Community Health & Wellbeing Workers has been informed by national learning and NHS Wales comparators. Earlier pilots in England utilised Band 2 roles; however, these were largely driven by affordability rather than role requirements.

Evidence from NAPC and existing sites indicates that lower banding limits recruitment, retention and the ability to deliver the full model.

The CHWW role requires:

- independent working within communities,
- safeguarding awareness and lone working,
- behaviour change and motivational support,
- navigation across health, social care and community services,
- management of a defined household caseload.

Benchmarking against NHS Wales roles indicates Band 3 represents the minimum appropriate banding, with Band 4 justified where the role includes caseload responsibility, structured assessment and MDT contribution.

The proposed model therefore adopts a Band 4 equivalent to ensure workforce capability, delivery quality and sustainability.

6.3. Team Management Co-ordination

Operational delivery of the Community Health & Wellbeing Worker (CHWW) model will be managed locally through third sector partner organisations, who will retain responsibility for day-to-day line management, safeguarding arrangements and employment responsibilities for the workforce.

Given the community-based nature of the role, Community Health & Wellbeing Workers will regularly operate independently within neighbourhoods and households. Experience from the Cornwall programme and similar community health worker models demonstrates that this type of outreach role requires strong local leadership, safeguarding support and professional supervision to ensure safe delivery and maintain workforce wellbeing.

A 1.0 FTE Team Lead role is therefore included within the programme structure to provide leadership, coordination and practical support to the CHWW workforce. The Team Lead role has been sized proportionately to the initial deployment (4 CHWWs), with scope to scale as the model expands.

This role will work closely with voluntary sector partners and local health services and will support:

- leadership and professional support for Community Health & Wellbeing Workers,
- safeguarding oversight and escalation of complex situations,
- support for lone working arrangements and workforce wellbeing,
- coordination with GP practices and local health services,
- establishing honorary arrangements such as with GP practices where required to enable appropriate integration with primary care teams,
- ensuring workers maintain compliance with key governance requirements such as GDPR and information governance protocols
- supporting community engagement activities, including facilitating community-based initiatives such as coffee mornings and outreach sessions,
- maintaining morale and professional development within the CHWW workforce through regular supervision and peer support.

The Community Health & Wellbeing Worker role involves engaging with individuals and families experiencing complex social and health challenges. Strong leadership and support structures are therefore essential to ensure that workers are well supported, confident in their role and able to sustain effective community engagement over time. This Team Lead role will help ensure the workforce operates safely, remains connected to local services and maintains the strong relational approach that is central to the CHWW model.

Typical Band 6 salary range (Agenda for Change): £39,263 - £47,280

For financial planning purposes a midpoint salary has been used to estimate workforce costs.

Management Cost

Role	Cost per annum	Total cost per annum
Team Lead Manager (Band 6 equivalent)	£41,437	
Employer on costs (NI, pension approx. ~27.5%)	£11,425	
Total Management cost (1.0 FTE)		£52,862

Total Management cost: approximately £52,862 per year

6.4. Training and Workforce Development

Community Health & Wellbeing Workers require an initial training and induction programme covering:

- Safeguarding
- community engagement
- health promotion
- system navigation
- motivational interviewing / health coaching
- lone working and safety.

Where possible, existing training provision within partner organisations and national programmes will be utilised to minimise additional costs. Based on comparable community workforce training programmes within the NHS and voluntary sector, an estimated training cost of approximately £1,000–£1,500 per worker has been assumed.

Estimated Training Costs

Item	Cost	Total cost per annum
Initial training and induction (per worker)	£1,500	
Training cost for 4 workers		£6,000

Total Training cost: approximately £6000

6.4.1. National Training and Accreditation Offer

To ensure consistency, quality and sustainability of the Community Health & Wellbeing Worker (CHWW) workforce, nationally recognised training pathways are available through the National Association of Primary Care (NAPC).

Two structured education routes are available:

- **Level 3 Certificate in Community Health and Wellbeing (Royal Society for Public Health)**
A structured 120-hour programme providing core competencies in community engagement, health improvement, safeguarding and system navigation.

- **Level 3 Apprenticeship in Community Health and Wellbeing (360 hours)**
A nationally recognised apprenticeship programme delivered across the four UK nations, incorporating the Level 3 certificate alongside broader workforce development.

Both routes:

- Are nationally recognised qualifications
- Provide consistency of workforce capability
- Support professionalisation of the role
- Enable scalability across Wales

The availability of these established training pathways reduces the need for local programme development and ensures alignment with best practice emerging from UK programmes.

6.5. Programme Infrastructure

Additional operational costs are required to support delivery of the programme and enable Community Health & Wellbeing Workers to operate safely and effectively within communities. These costs reflect typical requirements for community-based workforce programmes and include equipment, travel, monitoring infrastructure, and the systems required to support safe lone working and safeguarding practice.

Detailed infrastructure costs will be refined during implementation with delivery partners, but the above estimates provide a prudent allowance for safe and effective delivery of the initial deployment.

These include:

- **IT and Equipment** - This estimate is based on typical equipment costs for community-based workers requiring mobile working capability, including laptops or tablets, mobile phones and basic software access.
- **Travel expenses** - Travel costs are based on typical mileage reimbursement rates for community-based roles, recognising the rural geography of West Wales and the requirement for workers to travel between communities and households.
- **Monitoring and evaluation tools** - Monitoring and evaluation tools will support structured collection of activity data, outcome measures and evaluation outputs required to assess the effectiveness of the pathfinder.
- **Safeguarding and lone-worker systems** - As Community Health & Wellbeing Workers will be operating within community settings and, at times, undertaking home visits, provision is required for safeguarding support and lone-worker arrangements. This may include lone-worker devices or apps, check-in systems, incident reporting arrangements, DBS processing, and safeguarding process support within host organisations.

Infrastructure Estimate

Item	Total Annual Cost
IT and equipment	£4,000
Travel and community engagement	£8,000
Monitoring and evaluation tools	£5,000

Safeguarding and lone-worker systems	£3,000
Total infrastructure cost	£20,000

Total infrastructure cost: £20,000

6.6. Central Programme Coordination

Central programme coordination will be required to ensure that the Community Health & Wellbeing Worker (CHWW) model is implemented consistently across delivery areas and that learning from the pathfinder is captured effectively to inform future commissioning decisions.

This function will focus on strategic programme support rather than operational workforce management, as day-to-day line management and supervision of Community Health & Wellbeing Workers will be undertaken locally by the voluntary sector delivery partners.

Central programme coordination will support:

- programme oversight and coordination across partner organisations,
- development and implementation of the monitoring and evaluation framework,
- collation and analysis of programme activity and outcome data,
- generation of evidence to inform future commissioning decisions,
- coordination of reporting to programme governance structures,
- dissemination of learning and best practice emerging from the pathfinder.

This coordination will be delivered through a proportionate allocation of time from a Band 7 Project Manager, reflecting the level of programme management expertise required to support a multi-partner initiative involving health, voluntary sector and academic partners.

For the purposes of financial planning, an allocation equivalent to approximately 0.25 Full Time Equivalent (FTE) has been assumed to support programme coordination, monitoring and evaluation activities.

This approach ensures that appropriate programme oversight and evidence generation capability is in place while maintaining a proportionate and cost-effective level of programme infrastructure for a pathfinder of this scale.

Role	Cost per annum	Total cost per annum
Project Manager (Band 7 equivalent)	£51,027	
Employer on costs (NI, pension approx. 16.25%)	£8,292	
Total Project Co-ordination cost (0.25 FTE)		£14,830

Total Project Co-ordination cost: £14,830

6.7. Total Estimated Deployment Cost

Cost Category	Estimated Cost
Workforce (4 CHWWs)	£153,024
Team management	£52,862

Training	£6,000
NAPC	£15,000
Infrastructure	£20,000
Central coordination	£14,830
Total cost	£261,716

Total Estimated Annual Project Cost: £261,716

6.8. Cost Per Household Coverage

Based on comparable programmes (Cornwall and Westminster):

- each CHWW typically supports approximately 120 households
- four workers would therefore support approximately 480 households

Estimated cost per household: £261,716 ÷ 480 households ≈ **£545 per household per year**

This represents a relatively modest investment in preventative community engagement compared with the costs associated with reactive healthcare services.

6.9. Relative Cost Comparison

Community Health & Wellbeing Workers represent a cost-effective preventative workforce.

Approximate NHS activity costs include:

Activity	Approx Cost
GP consultation	£40
A&E attendance	£110 - £565
Emergency hospital admission	£1,600–£3,500

As stated in Section 2, it is estimated that 20% of GP consultation time is spent on non-medical issues and that for Hywel Dda University Health Board, this equates to over 460,000 GP consultations.

Applying this methodology to Ty Elli Practice (over 14,000 patients registered), this estimates nearly 17,000 non-medical consultations, equal to nearly £700,000.

6.10. Wider System Value

In addition to potential reductions in healthcare utilisation, the programme may also generate wider social value through:

- improved employment engagement,
- reduced social isolation,
- improved mental wellbeing,
- stronger community resilience.

These benefits align closely with the objectives of place-based health and wellbeing systems and the principles of the Wellbeing of Future Generations Act.

6.11. Benchmarking Against UK Delivery Models

The proposed cost model aligns with national delivery experience from existing Community Health & Wellbeing Worker programmes.

NAPC modelling based on UK pathfinder sites indicates a typical annual cost of approximately £215,000 for a team comprising:

- 4 Community Health & Wellbeing Workers,
- 1 Team Manager,
- Associated infrastructure, training and support.

The proposed West Wales model (£261,716) reflects:

- Use of Band 4 workforce (higher than Band 2 in England models),
- Inclusion of enhanced governance, safeguarding and evaluation requirements,
- Rural delivery considerations (travel and infrastructure).

This demonstrates that the proposed model is:

- Consistent with national benchmarks,
- Realistic in cost assumptions,
- Appropriately adjusted for the Welsh context.

6.12. Financial Summary

The proposed Community Health & Wellbeing Worker deployment represents a modest and proportionate investment relative to wider system costs while offering the potential to:

- strengthen prevention,
- reduce inequalities,
- improve engagement with health and wellbeing services,
- reduce demand on reactive services.

The estimated annual programme cost of approximately £261,716 therefore represents a realistic and proportionate investment to test the model within the West Wales context. Over two years, this equates to a total investment of £523,432

7. Management Case

7.1. Governance and Programme Management

Effective governance and programme management are critical to ensuring the successful implementation and evaluation of the Community Health & Wellbeing Worker (CHWW) deployment.

Given the breadth of services this model is intended to support, governance will need to provide assurance not only over delivery of the CHWW workforce itself, but also over coordination across the wider coalition of partners involved in prevention, community support and household resilience.

The programme will operate within a structured governance and delivery framework designed to provide clear accountability, effective oversight and strong partnership working across health, local government, academia and voluntary sector organisations. It builds on established delivery models and existing partnerships, reducing implementation risk and enabling rapid mobilisation.

This governance model ensures that the programme benefits from strategic leadership, operational coordination and community-based delivery capability.

7.2. Strategic Partnerships

The Centre for Social Innovation (CfSI) will act as a strategic partner supporting model design, innovation and evaluation, rather than as a contracted delivery provider of CHWW services. Operational delivery of the workforce will be commissioned through voluntary sector organisations via grant or SLA arrangements.

The CfSI will play a key role in shaping the model beyond traditional service delivery, supporting:

- development of innovative, community-led approaches to health and wellbeing,
- identification of systemic barriers affecting communities,
- design of approaches that support long-term behavioural and societal change,
- evaluation of wider impacts beyond traditional health metrics, including community resilience and social outcomes.

This ensures that the programme not only delivers immediate service benefits but also generates learning on how public services can evolve to better support communities in the long term. Where CfSI provides support to the programme (e.g. model development, evaluation design or innovation), this will be delivered through a clearly defined partnership arrangement aligned to governance and procurement requirements.

This ensures a clear distinction between strategic design, delivery, and commissioning responsibilities.

7.3. Senior Responsible Owner

The programme will operate under a Senior Responsible Owner at Executive Director level will be appointed. The SRO will hold overall accountability for the programme and will be responsible for:

- strategic leadership of the programme
- ensuring alignment with national public health priorities
- oversight of programme performance and delivery
- management of strategic risks
- reporting progress to senior stakeholders and funding bodies.

The SRO will provide clear executive accountability, ensuring delivery remains aligned to strategic priorities and that risks, performance and outcomes are actively managed throughout the pathfinder lifecycle.

7.4. Programme Board

Strategic oversight of the programme will be provided through a Programme Board, which will provide assurance that the pathfinder is being delivered effectively and in accordance with agreed objectives.

The Programme Board will monitor:

- programme delivery and milestones
- financial performance
- risk management
- evaluation progress.

The Programme Board will also ensure that the programme remains aligned with regional and national strategic priorities.

Membership of the Programme Board will include representatives from:

- Public Health Wales
- Hywel Dda University Health Board
- Centre for Social Innovation (University of Wales Trinity Saint David)
- Voluntary Sector
- local authority partners.

The Programme Board will meet quarterly to review progress, risks and programme performance.

7.5. Project Management

Day-to-day coordination of the programme will be undertaken by a Project Manager, who will be responsible for ensuring that implementation progresses effectively across all partner organisations.

Key responsibilities of the Project Manager include:

- coordinating recruitment and workforce development,
- supporting integration with health, social care and community services,
- maintaining the programme risk register,
- monitoring programme performance,
- coordinating monitoring, reporting and evaluation.

The Project Manager will report to the Senior Responsible Owner and Programme Board, ensuring that delivery issues can be escalated and addressed promptly.

7.6. Operational Delivery

Operational delivery of the programme will be undertaken through third sector partner organisations, which will host and manage the Community Health & Wellbeing Workers.

Local delivery partners could include:

- Carmarthenshire Association of Voluntary Services,

- Housing Association

These organisations will be responsible for:

- recruitment of Community Health & Wellbeing Workers,
- day-to-day line management and supervision,
- safeguarding oversight,
- supporting integration with local services and community organisations.

The voluntary sector is particularly well placed to deliver this workforce due to its strong community presence, established networks and local intelligence regarding community needs.

7.7 Workforce Development and Career Pathways

A structured workforce development approach will be implemented to support recruitment, retention and progression of Community Health & Wellbeing Workers.

This will include:

- access to nationally recognised Level 3 qualifications through NAPC (certificate or apprenticeship routes),
- ongoing professional development supported through peer networks and supervision structures,
- exploration of accredited training delivery through UWTSD to support long-term sustainability.

The programme will also work with partners to define clear career pathways, enabling CHWWs to progress into wider roles across health, social care and community services where appropriate, while also recognising and supporting the CHWW role as a valuable standalone career.

This approach supports both workforce retention and system integration, ensuring that skills developed through the programme contribute to the wider health and care workforce over time.

The model also provides a sustainable workforce solution, creating local employment opportunities and structured pathways into health and social care careers.

7.8 Risk Management

The following risks have been informed by learning from existing Community Health & Wellbeing Worker programmes (including Cornwall and Westminster), as well as wider experience from social prescribing and community-based delivery models. This ensures that risks reflect real-world implementation challenges rather than theoretical assumptions.

Risk	Likelihood	Impact	Mitigation
Workforce recruitment and retention challenges	Medium–High	High	Early engagement with Voluntary Care Sector partners; targeted local recruitment; flexible role design; clear career pathways (NAPC → UWTSD); strong supervision model (Team Lead); learning from Cornwall shows retention improves with peer support and local identity

Role ambiguity and duplication with existing services (e.g. social prescribing, community connectors)	Medium	High	Clear role definition and comms; onboarding with GP clusters and MDTs; pathway mapping; positioning CHWW as <i>complementary (proactive + relational)</i> not duplicative; learning from Westminster highlights need for clarity early
Limited engagement or buy-in from primary care	Medium	High	Early GP engagement via Ty Elli cluster; identify clinical champion; demonstrate workload benefit (non-medical demand reduction); regular feedback loops; align to cluster priorities
Data sharing and access barriers (Digital systems, IG constraints)	Medium	High	Early IG engagement; DPIA and data sharing agreements; use anonymised/aggregate data where required; phased access model;
Difficulty demonstrating measurable impact (attribution challenge)	Medium	High	Built-in evaluation design with comparator group; pre/post analysis; use of validated tools (MYCaW); academic partner support (UWTSD/CfSI); mixed-methods approach to capture both quantitative and qualitative impact
Over-reliance on individual CHWW relationships (model fragility)	Medium	Medium–High	Standardised training and operating model (NAPC); shared case management system (Octolink); team-based supervision
Safeguarding risk in community-based lone working	Low–Medium	High	Lone worker systems; safeguarding training; clear escalation pathways; Team Lead oversight; alignment with existing safeguarding frameworks
Inconsistent delivery across partners / variation in quality	Medium	Medium	Standardised training and induction; central coordination; peer network (NAPC); regular review and learning sessions; use of common tools (Octolink, MYCaW)
Demand exceeds capacity (early success risk)	Medium	Medium–High	Clear cohort definition (Ty Elli focus); phased onboarding; prioritisation criteria; regular caseload review; learning from Cornwall shows demand grows quickly once model is visible
Community mistrust or low engagement in target populations	Low–Medium	High	Recruit locally embedded workforce; partner with trusted organisations; relationship-first approach; no referral threshold barrier; learning from Westminster shows trust-building is critical in first 3–6 months
Insufficient integration with wider system (LA, housing, benefits)	Medium	High	Coalition model; formal partner engagement; Programme Board representation; pathway mapping; CHWW role explicitly includes “joining the dots” across services
Financial sustainability beyond pathfinder unclear	Medium	High	Clear evaluation framework; cost-per-household model; early engagement with commissioners; demonstrate value through

			multiple benefit domains (health + social + economic)
Digital / data tool adoption challenges (Octolink, MYCaW)	Low–Medium	Medium	Training via NAPC; simple user workflows; ongoing support; phased implementation; avoid overburdening workforce with data collection
Programme perceived as “pilot that doesn’t scale”	Medium	Medium–High	Design with scalability in mind; standardised model; national alignment (NAPC); local workforce pipeline (UWTSD); early documentation of model and learning
Role drift into statutory or backlog functions	Medium	Medium	Clear role definition; maintain prevention focus; regular scope review through Programme governance

Risks will be actively managed through the programme governance structure. A risk register will be maintained by the Project Manager and reviewed regularly by the Programme Board.

Where risks cannot be resolved at programme level they will be escalated to the Senior Responsible Owner for resolution.

7.9 Monitoring and Evaluation

A structured monitoring and evaluation framework will be implemented to ensure that the pathfinder generates robust evidence to inform future commissioning decisions.

Performance monitoring will include:

- workforce activity metrics,
- community engagement levels,
- service referral and connection rates,
- preventative health engagement,
- qualitative feedback from residents and partners.

These measures will enable the programme to track early outcomes, identify learning and refine the delivery model during the pathfinder phase.

To strengthen attribution of impact, the evaluation will incorporate a comparator approach. This will include:

- Comparison with a matched population within the same GP cluster not receiving CHWW support (e.g. similar demographic and deprivation profile),
- Pre- and post-intervention analysis for the CHWW cohort,
- Trend comparison against baseline practice-level activity data.

This approach will enable differentiation between programme impact and natural variation in demand over time, strengthening the robustness of findings and supporting future commissioning decisions.

In addition to standard health and service utilisation metrics, the evaluation will also consider wider social and community outcomes, including:

- changes in household resilience and self-sufficiency,
- levels of social connection and community participation,
- barriers to accessing services and how these are overcome,
- the effectiveness of relationship-based approaches in driving behaviour change.

This broader evaluation approach, supported by CfSI and academic partners from the University of Wales Trinity Saint David, will provide deeper insight into how community-based models can contribute to long-term system transformation while strengthening the overall evidence base generated by the programme.

7.10 Programme Assurance

Programme assurance will be provided through regular performance reporting to the Programme Board and Senior Responsible Owner.

This will ensure that:

- delivery remains aligned with programme objectives,
- risks are identified and managed early,
- learning from the pathfinder is captured effectively.

This assurance framework will support informed decision making regarding the future commissioning and potential scaling of the CHWW model.

8 Conclusion

This business case presents a clear opportunity to implement a proven, evidence-based model that addresses one of the most significant and persistent challenges facing the health and care system: avoidable demand driven by unmet social and community need.

International evidence from Brazil and emerging UK experience in areas such as Westminster and Cornwall demonstrate the potential for this workforce to significantly increase engagement with preventative health services while reducing demand on reactive healthcare provision.

The proposed West Wales pathfinder provides a practical and proportionate opportunity to test how this model can be implemented within the Welsh context while strengthening collaboration across health, social care and community services.

By combining locally embedded voluntary sector delivery with central programme coordination and academic evaluation, the pathfinder will generate valuable learning to inform potential future commissioning decisions across Wales.

This represents a strategic opportunity to test a scalable model for prevention, community resilience and system sustainability. Approval will enable West Wales to lead the development of a new, integrated approach to health and wellbeing that can inform future policy and commissioning across Wales.

8.7 Recommendation

It is recommended that approval is granted to proceed with the Community Health & Wellbeing Worker pathfinder in West Wales. This represents a low-cost, high-impact intervention that directly addresses one of the most significant drivers of demand within the health and care system.

This will enable:

- recruitment of four Community Health & Wellbeing Workers (Band 4 equivalent),
- recruitment of a Team Leader (Band 6 equivalent)
- initial deployment through the Ty Elli Practice Group across Carmarthenshire,
- delivery through third sector partners,
- implementation of a structured monitoring and evaluation framework.

The pathfinder will generate evidence to inform future commissioning decisions regarding the potential expansion of Community Health & Wellbeing Workers across Wales.

8.8 Decision Required

Approval is sought for:

1. Implementation of the Community Health & Wellbeing Worker pathfinder in West Wales.
2. Deployment of four Community Health & Wellbeing Workers through voluntary sector delivery partners.
3. Allocation of the proposed pathfinder budget of approximately £261,716 for the first year of delivery, within an overall two-year investment of £523,432
4. Establishment of the governance and evaluation framework outlined in this business case.

8.9 Next Steps

Subject to approval, the following next steps will be undertaken:

- establishment of programme governance arrangements
- finalisation of service-level agreements with delivery partners
- recruitment and onboarding of Community Health & Wellbeing Workers
- implementation of training and induction
- launch of pathfinder delivery within target communities.

The programme will commence within six months of approval, with evaluation findings informing future commissioning decisions.

Addendum A: Clarifications on Workforce Banding, Return of Investment and Scale

Purpose

This addendum provides clarification on workforce banding, return on investment (ROI) and model scale to support decision-making and ensure alignment with national evidence and delivery experience.

Workforce Banding (Rationale and Benchmarking)

Community Health & Wellbeing Workers (CHWWs)

The proposed banding for Community Health & Wellbeing Workers has been informed by:

- Learning from existing UK CHWW programmes supported by the National Association of Primary Care (NAPC)
- Benchmarking against NHS Wales community-facing roles
- The functional requirements of the role within the West Wales model

Early CHWW pilots in England utilised Band 2 roles, largely driven by affordability constraints rather than the requirements of the model. Evidence from these programmes indicates that lower banding:

- restricts recruitment to candidates with limited experience
- impacts confidence and autonomy in community settings
- reduces effectiveness in navigating complex systems
- contributes to higher turnover and reduced continuity

As the model has matured, Band 3 has emerged as the minimum recommended level, with many programmes adopting higher banding where the role includes greater responsibility.

The CHWW role in West Wales requires:

- independent community-based working
- proactive household engagement
- safeguarding awareness
- navigation across health, social care and community services
- longitudinal relationship management

Benchmarking against NHS Wales roles indicates that Band 3 aligns with entry-level community support roles, while Band 4 is appropriate where the role includes caseload management, structured assessment and multi-agency coordination.

The proposed Band 4 equivalent therefore reflects the level of autonomy, responsibility and system navigation required to deliver the model effectively and sustainably. This approach also supports longer-term workforce sustainability through alignment with emerging national workforce pathways, including early discussions with Health Education and Improvement Wales (HEIW) regarding potential apprenticeship development.

Team Lead

The original proposal included a Band 7 Team Lead role. Following further review and benchmarking, this has been refined to ensure proportionality to the scale of the initial deployment.

The team role will:

- provide leadership and professional support to CHWWs
- oversee safeguarding and lone working arrangements
- act as a key interface with GP practices and local services
- support integration across partner organisations
- maintain workforce wellbeing and performance
- coordinate local delivery alongside central programme support

A Band 6 equivalent is considered proportionate for the scale of the initial deployment, providing sufficient seniority to manage safeguarding, workforce supervision and multi-agency coordination, without over-specifying the role relative to team size.

Return on Investment (Measurement Approach)

The CHWW model delivers value across multiple parts of the system and should not be assessed against a single cost-saving metric.

Key value domains include:

1. Acute Demand Reduction

- Reduced A&E attendances
- Reduced non-elective admissions
- Potential reductions in length of stay

2. Primary Care Impact

- Reduction in avoidable GP consultations
- Improved targeting of clinical time towards higher-need patients
- Reduction in repeat attendance (“failure demand”)

3. Prevention and Early Intervention

- Increased vaccination uptake
- Increased screening participation
- Earlier identification of health and social needs

4. System Efficiency

- Reduced missed appointments (DNA rates)
- Increased uptake of community and voluntary sector services
- Improved service navigation and utilisation

5. Patient Outcomes and Experience

- Improved wellbeing (e.g. MYCaW outcomes)
- Increased patient activation and confidence
- Strengthened trust and engagement with services

6. Wider Determinants and Social Value

- Improved access to housing, benefits and employment support
- Reduced social isolation
- Increased community resilience

ROI will be presented as:

- cost per household supported,
- cost relative to avoided or reduced demand,
- system-wide value across health, social care and community services.

This approach avoids reliance on a single high-impact metric (e.g. avoided hospital admissions) and reflects national experience that most value is realised through reduced repeat demand, improved care navigation and avoided escalation, rather than isolated high-cost events.

Illustrative scenario modelling, included within this Business Case, demonstrates that even small reductions in demand (e.g. reduced GP consultations across a proportion of households) begin to offset programme costs, while wider system benefits provide additional value.

Scale and Evidence Base (National Context)

UK Delivery

The CHWW model is now an established and scaling approach within the UK.

- 24 live sites across England delivery includes Westminster, Cornwall and multiple London and regional sites,
- Approximately 192 CHWWs currently deployed,
- Supported through the NAPC national programme.

Across these sites, consistent patterns are emerging, including:

- improved engagement with preventative services,
- earlier identification of need,
- increased uptake of vaccination and screening,
- reduced pressure on primary and urgent care services.

This level of adoption demonstrates that the model is established and scalable, rather than experimental, reducing implementation risk for West Wales.

International evidence (Brazil):

The strongest international evidence comes from Brazil's Family Health Strategy, one of the largest community-based health programmes globally.

- Over 300,000 community health workers,
- Coverage of approximately 70% of the population.

Published evaluations have demonstrated:

- sustained reductions in hospitalisation rates,
- improved chronic disease management,
- increased vaccination uptake,
- improvements in maternal and child health outcomes.

Reported outcomes also include:

- 34% reduction in cardiovascular mortality,
- 31% reduction in stroke mortality.

Implications for West Wales

This evidence demonstrates that:

- the CHWW model is proven, not experimental,
- the model is adaptable across urban and rural settings,
- impact is consistent across multiple domains of health and wellbeing,
- effectiveness increases with scale, integration and maturity.

The proposed West Wales pathfinder therefore represents:

- a low-risk, evidence-informed deployment,
- aligned with established national and international models,
- capable of generating robust, locally relevant evidence to inform future commissioning decisions.

Conclusion

This addendum reinforces that:

- Band 4 CHWW roles and Band 6 Team Lead roles are proportionate and evidence-based within NHS Wales context,
- Return on investment should be understood as multi-domain system value rather than a single financial metric,
- The CHWW model is already established at scale across the UK and internationally, with a strong and growing evidence base.

Taken together, this confirms that the proposed West Wales model is appropriately costed, proportionately staffed, and represents a credible, low-risk and evidence-informed investment aligned to prevention, system sustainability and community-based care.

Demonstrator project report of Measure Yourself Concerns and Wellbeing[®], demographic data and outcome measures analysis for the CHWW project - Year One



**Dr Marie Polley
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June 2024

Disclaimer: The views expressed in this report are those of the authors and do not represent those of the Cornwall and Isles of Scilly Integrated Care Board or Volunteer Cornwall.

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Meaningful Measures Ltd provides innovative person-centred evaluation, audit and research solutions. Our mission is to enable people's own voices to shape the understanding and improvement of health and wellbeing services around the world.

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Executive summary

In the UK, the Office of National Statistics calculates that 4.7 million people live in poverty. Deprivation has severe impacts on physical and mental health, and this puts a strain on public services which are already overstretched. Some areas of Cornwall and Isle of Scilly are in the top 10% most deprived areas in England.

In February 2023 Cornwall and Isles of Scilly Integrated Care System introduced teams of “Community Health and Wellbeing Workers” (CHWW) to support underserved communities with the aim of building relationships and identifying people’s needs. The initial pilot has now been extended to run for a minimum five years, working in the Kerrier area of Cornwall, and extending to Penwith, Bodmin and Liskeard. There is an aim to have a total of 50-60 CHWWs in post by Autumn 2024.

CHWW’s encourage householders to initially identify their biggest concern and then work with them to reduce or remove that concern. Since Autumn 2023 Measure Yourself Concerns and Wellbeing® (MYCaW) data has been collected to show the impact of the CHWW. MYCaW is a validated individualised questionnaire designed for evaluating holistic and personalised approaches to supporting people and is accredited by NHS England for supported self-management services (such as social prescribing). MYCaW provides qualitative (written concerns) and quantitative (score change on 0 to 6 scale) data. MYCaW data analysis for this project is reported to demonstrate the initial impact of the CHWW project.

Aim:

To analyse all data on the initial and follow-up MYCaW forms from the CHWW project, as well as any relevant available demographic data about the project recipients.

To carry out a comparison of commonly used outcome measures with concerns documented from MYCaW data.

Method:

MYCaW data was gathered by CHWW teams during their home visits. The first MYCaW was carried out after a CHWW had got to know the resident and was applied after at least the third meeting. Follow-up MYCaW data was captured 3 months later.

Results:

Demographics:

N=211, the modal age was 35-44 years old, 60% of the sample was female. Nearly 80% of residents lived in either St Austell or Newquay.

Measure Yourself Concerns and Wellbeing® score changes:

Score changes for all people with baseline and paired follow-up scores were analysed. The lower the score the better the rating of the severity of a concern or wellbeing status. Overall, the data suggest that most people who were supported by the service, their concerns and wellbeing improved as a result.

Average MYCaW change scores for Concern 1 (-2.6, n=147; $p \leq 0.001$) Concern 2 (-2.4, n=130; $p \leq 0.001$) and Wellbeing (-1.8, n=147; $p \leq 0.001$) showed a large improvement at follow-up and were statistically significant.

91% and 88% of Concern 1 and 2 score changes (respectively) and 90% of the wellbeing score changes met the threshold for minimal important difference. This result is extremely high for percentage improvements compared to other MYCaW datasets.

What do the MYCaW concerns tell us?

412 concerns were reported when combining Concern 1 and Concern 2. Concerns were coded using an existing established framework by Meaningful Measures Ltd based on other existing national data sets.

The top 5 individual concern categories reported were Finances, Housing, Mental health, loneliness and isolation and access to food.

Nearly 50% of all concerns were practical ones, (e.g. problems with finance, housing, managing the household admin, managing to look after ones-self, social justice and safety issues, work and training, school and education and carer burden). Around 25% of concerns related to a broad range of mental and emotional issues.

What else is going on in your life?

MYCaW also asks people if there is anything else going on in their life that is important, besides the concerns they have nominated. 130 people recorded a total of 139 other things going on in their life. 42% of events were positive in nature most frequently an improvement in their finances or work situation (it was implied that the CHWWs had helped get benefit payments sorted or enable job finding). 49% of events were negative in nature. The most frequent negative issue was ongoing issues with physical and mental health (20%) and this often related to issues with long term conditions and pain.

What is important about the support you have received? 142/212 people responded to this question, providing 202 things that were important about the support they received from the CHWWs. The most frequent response (38%) related to the support, understanding and provided by the CHWWs as well as the encouragement to make changes or improvements.

Conclusion:

The first aim of the demonstrator report was to investigate the MYCaW score changes and understand the types of concerns people experience, as well as what aspects of the CHWW support are important to them.

Average changes in MYCaW scores after receiving support from the CHWWs were very large and highly statistically significant. A comparison group is needed for this to ascertain how much change is attributable to the CHWW intervention. Qualitative data patterns showed that the top concerns were: Finances, housing, mental health, loneliness & isolation and access to food. The demonstrator report shows that the CHWW project is a highly successful and valuable service supporting people in Cornwall and IoS to access what are considered their basic needs and rights to appropriate finance, housing, food and support with their physical and mental health. We conclude that this service should be continued, and more data collected to establish the longer-term impacts on the current residents.



The second aim of the evaluation was to carry out a comparison of commonly used outcome measures informed by the types of concerns documented from MYCaW data. The most often picked MYCaW super-category was “practical concerns” and most of these concerns related to the social determinants of health. Other outcome measures for health and wellbeing may not be able to reach all of the areas that MYCaW reaches, especially in terms of social determinants of health. We provide a summary review of the most useful measures, those which are short, free, sensitive and provide economic value data which could partner with MYCaW (suggestions include: SF-36, ONS4, EQ5D, and Dialogue+).

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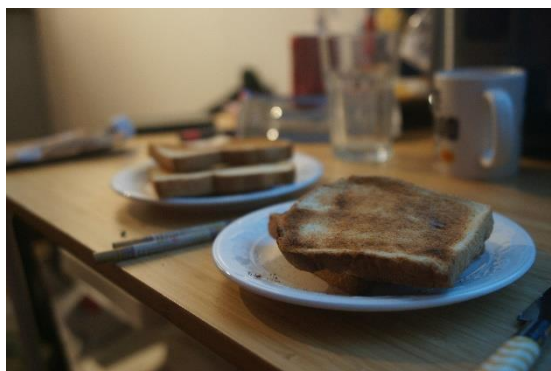
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1.0 Introduction

1.1 Background

The number of people experiencing deprivation (people struggling to afford to meet their most basic physical needs to stay warm, dry, clean and fed), has increased by 148% from 2017 to 2022. 3.8 million people experienced destitution in 2022, including around one million children¹. Deprivation has severe impacts on physical and mental health and this puts a strain on public services which are already overstretched². Improvements in life expectancy have been shown to have stalled, and for women in the most deprived 10% of areas life expectancy has declined. The health gap has grown between wealthy and deprived areas³.

In 2024 the County of Cornwall and IoS faces significant challenges with deprivation for its population. Salaries in Cornwall and IoS are about 80% of the national average, contributing to poverty among families and children⁴. Some areas in Cornwall and IoS are in the top 10% most deprived areas in England⁵. NHS England are keen to support projects that tackle deprivation and health inequalities.



In 2023 an innovative project to help tackle inequalities was launched within the NHS Cornwall and Isles of Scilly Integrated Care System. This project was the introduction of teams of “Community Health and Wellbeing Workers” (CHWW). The CHWW

¹ Firzpatrick, S. et al. (2023) Destitution in the UK 2023, Rowntree Foundation, <https://www.jrf.org.uk/deep-poverty-and-destitution/destitution-in-the-uk-2023>

² Bestwick, M. and Schmuecker, K. (2024) The impact of hardship on primary schools and primary healthcare. York: Joseph Rowntree Foundation.

³ Marmot M, Allen J, Boyce T, Goldblatt P, Morrison J. Health Equity in England: The Marmot Review 10 Years On. Institute of Health Equity; 2020 (health.org.uk/publications/reports/the-marmot-review-10-years-on).

⁴ <https://www.local.gov.uk/case-studies/responding-cost-living-challenges-cornwall-council>

⁵ <https://transformation-cornwall.org.uk/resources/cornwall-council-index-of-multiple-deprivation-imd-2019-and-2015-cornwall-reports>

initiative is based on the successful “Brazilian Family Health Strategy”^{6, 7} developed in the 1990s. The CHWW role involves proactively going door-to-door in underserved communities with the aim of building relationships and identifying people’s needs. This model will also identify people who are not engaging in healthcare services and often excluded from them. The teams will actively encourage residents to register with a GP to enable them to take part in vaccinations, screening and other healthy behaviours.

Since 2019, the National Association of Primary Care alongside Imperial College London have been working together to support the implementation and spread of this in many areas in the UK⁸. This model was firstly tested in the London borough of Westminster⁹ and spread to Bridgewater, Calderdale, Wandsworth, Oxford, and North Wales⁶, amongst other areas.



Figure 1. Brazilian Community Health Worker model.

⁶ Johnson, C.D., Noyes, J., Haines, A. et al. Learning from the Brazilian Community Health Worker Model in North Wales. *Global Health* 9, 25 (2013). <https://doi.org/10.1186/1744-8603-9-25>

⁷ Junghans, C., Antonacci, G., Williams, A. et al. Learning from the universal, proactive outreach of the Brazilian Community Health Worker model: impact of a Community Health and Wellbeing Worker initiative on vaccination, cancer screening and NHS health check uptake in a deprived community in the UK. *BMC Health Serv Res* 23, 1092 (2023). <https://doi.org/10.1186/s12913-023-10084-8>

⁸ <https://napc.co.uk/community-health-and-wellbeing-workers/>

⁹ Dallera, G., Antonacci, G., Junghans C, Harris M. (2022). Measuring the impact of the Community Health and Well-being Worker initiative in Westminster on immunisation, screening, NHS Health Check uptake, and GP consultation rates. Report commissioned by Westminster City Council. Oct 2022

Within Cornwall and IoS there was an initial starter pilot in February 2023 that recruited 10 CHWW in four locations in the Central Cornwall Integrated Care Area (ICA). This involved working with the voluntary sector in three areas and a Primary Care Network (PCN) in the fourth. Teams of CHWWs are now being rolled out in the 10% most



deprived areas within Cornwall and IoS. Initial work has been reported on in a qualitative evaluation¹⁰ investigating how the CHWW project got off the ground, how CHWWs were recruited and trained. This work did not investigate any of the impact of the CHWW work on residents. The initial report found many positive aspects of the project and also identified some areas of learning. One of these areas was the need to have a clear understanding between the CHWW role and the social prescribing link worker roles, and that elements of best practice in social prescribing could be echoed in the CHWW role.

The initial pilot that started in February 2023 has been extended to run for a minimum five years with a larger workforce of CHWW¹¹. As part of the extension, in January 2024, a team of 22 CHWWs were appointed in the Kerrier area of Cornwall. With the further investment from Cornwall and Isles of Scilly Integrated Care Board, CHWW teams are now developing in Penwith, Bodmin and Liskeard. There is an aim to have a total of 50-60 CHWWs in post by the Autumn of 2024¹².

1.2 Logistical details of the current CHWW project

CHWWs work within the ICS, so while they are part of the primary care team, they are also integrated into the wider health and social care system and local communities allowing them to quickly connect with all relevant services for their householders.

¹⁰ John Tredinnick-Rowe, Richard Byng, Tamsyn Brown et al. Piloting a Community Health and Well-being Worker Model in Cornwall: a guide for implementation and spread, 29 March 2024, PREPRINT (Version 1) available at Research Square [<https://doi.org/10.21203/rs.3.rs-4144720/v1>]

¹¹ <https://www.volunteercornwall.org.uk/how-we-help/in-the-community/chww#CHWWNews>

¹² <https://cios.icb.nhs.uk/2024/03/06/cornwalls-success-with-community-health-and-wellbeing-workers/>

Depending on caseloads and working hours (full or part time), the CHWWs are each appointed around 120 households to support and build long term relationships with whole families following the principles of comprehensive intergenerational, hyperlocal, universal and integrated .

CHWWs are recruited locally based on the knowledge of the local community. Recruitment is not qualifications based, but instead focused on a candidates' ability to build relationships and to understand a community¹³. CHWW staff also work alongside those in social prescribing link worker roles.

In Cornwall and IoS, CHWWs typically meet people within their own homes, or within the local community (e.g. a café or community hub). Part of the CHWW's role is to encourage householders to initially identify their biggest concern and then work with them to reduce or remove that concern. Since Autumn 2023 the CHWW project has been collecting Measure Yourself Concerns and Wellbeing® (MYCaW) data to show the impact of the CHWW.

1.3 What is the Measure Yourself Concerns and Wellbeing® (MYCaW) tool?

The MYCaW questionnaire is a validated individualised questionnaire designed for evaluating holistic and personalised approaches to supporting people and is accredited by NHS England for supported self-management services (such as social prescribing). MYCaW only takes a few minutes to complete and can routinely be incorporated into a consultation to understand and prioritise what a person most wants support with. MYCaW allows a more rigorous approach to capturing the voice of residents beyond the anecdotal.

First form - Each person writes down the one or two things that concerns them the most, that they want help with, in a box. These concerns can relate to any area of a person's life and are rated for severity using a simple numerical Likert scale and a person's wellbeing is also scored.

¹³ See video: <https://www.youtube.com/watch?v=HMXOnbCEfi0>

At follow-up - the original concerns are rescored, and wellbeing is rescored. Two further questions ask if anything else important is happening in a person's life and what has been most important about the service received. These questions are useful for identifying confounding variables which may affect concerns and wellbeing score changes, as well as capturing what the residents value about the support they are receiving.

MYCaW will capture concerns that are medically related, but more often people report concerns that affect their biopsychosocial wellbeing. People also report practical concerns around finance, housing, job status, concerns about their future, their life purpose, relationships and carer burden. All of these concerns may impact a person's overall wellbeing, yet, many of these topics are not routinely recorded.

There are several coding frameworks for MYCaW which sit behind this tool allowing researchers the ability to easily categorise the qualitative data captured by the tool. Quantitative data can be analysed to track score changes and work out the percentage of people making a clinically relevant change to their concerns and wellbeing and assess the statistical relevance of the mean score changes.

MYCaW can be accessed as a paper or digital version and a licence is required from www.meaningfulmeasures.co.uk to use the tool.

1.4 Aim of report

- To analyse all data on the initial and follow-up MYCaW forms from the CHWW project, as well as any relevant available demographic data about the project recipients.
- To carry out a comparison of commonly used outcome measures with concerns documented from MYCaW data. (Please see Appendix 1 at the end of the report for this).



2.0 Methods

Data was gathered by the CHWW teams during their home visits. The first MYCaW was presented to an individual once a CHWW had got to know the resident, it was not necessarily administered at the first meeting, but was applied after at least the third meeting. Follow up MYCaW data was captured 3 months later (however, this time frame was not rigid, and it may vary by several weeks per participant). If someone wished to leave the project then a follow-up MYCaW was administered.

MYCaW data was held in a centralised excel sheet which CHWW added their information to.

Anonymised service data including demographics and MYCaW data was sent to Meaningful Measures Ltd in a password protected excel file. The data file was checked and cleaned then converted into a data file for IBM SPSS Statistics version 29.0.1.0 (171).

2.1 Analysis of Demographic data

All demographics were calculated as percentage frequencies.

2.2 Analysis of Measure Yourself concerns and Wellbeing® concern types

The MYCaW concerns were analysed for the whole dataset. To do this, all the first and second concerns were combined. They were initially coded, using the personalised care coding framework devised by Meaningful Measures Ltd. This was reviewed by a second researcher and a final version agreed. The proportion of concerns in each theme and category was then calculated to enable visual representation in the results section of this report.

2.3 Analysis of score changes

The mean score changes and accompanying standard deviation for MYCaW Concern 1, Concern 2 and Wellbeing were calculated. To determine if the overall change in

Concerns and wellbeing scores was statistically significant, a paired samples 2-way T-Test was carried out on all concerns scores that had a baseline and follow-up score. This meant that all Concern or Wellbeing scores that did not have a follow-up score were omitted from this calculation.

Finally, the percentage of scores that met the minimal threshold of change to signify an important difference were calculated. This includes scores that met the improvement threshold of 1 point and scores that met the deterioration threshold of 1 point. This level of changes (minimal important difference) is the level of change needed for a demonstrable difference to be experienced by the participant.

3.0 Results

3.1 Demographics

211 people's data was available to analyse. Demographic data on age, gender and area of Cornwall were all analysed. Figure 2 shows the range of ages that engaged in the CHWW service. Approximately a quarter of all residents were between 35-44 years old. 18% of residents were between 55-64 years old and 15% were between 70-89 years old. There was a very large range between the youngest and oldest residents e.g.15-19 years old through to 95 -99 years old.



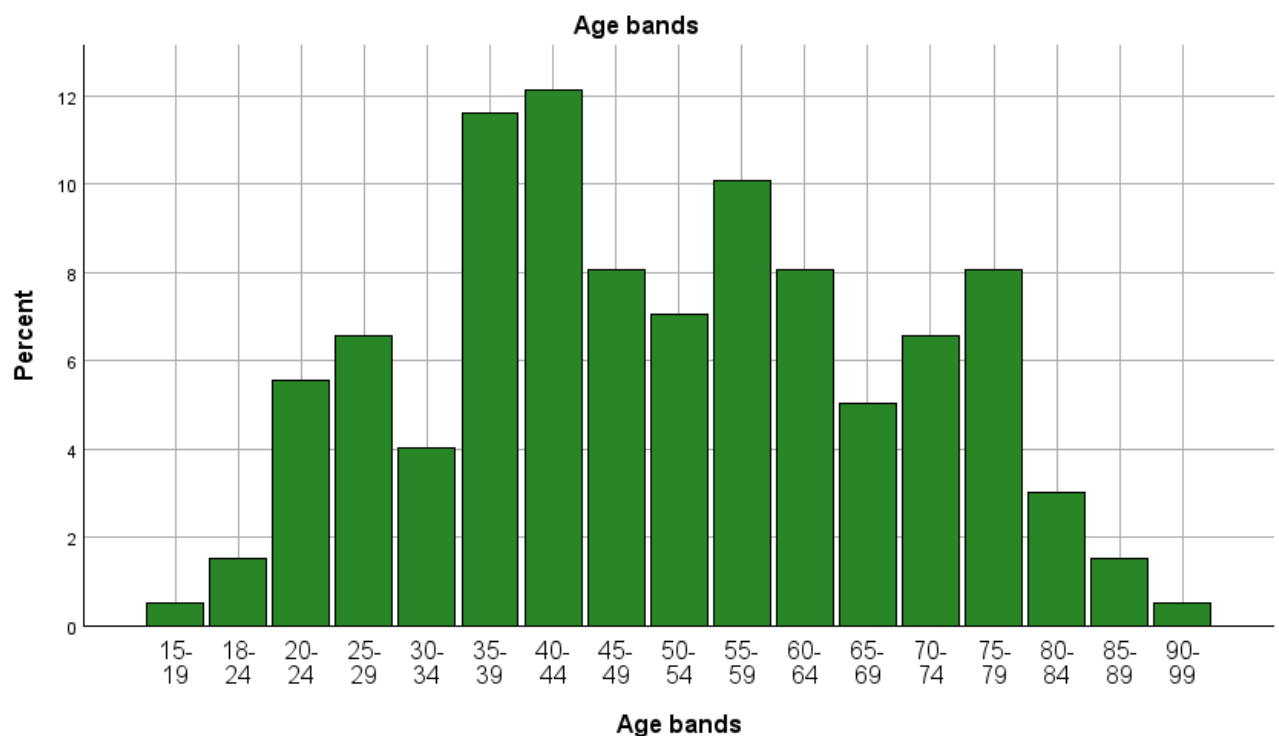


Figure 2. Range of ages of CHWW residents.

As shown in Table 1 below, the CHWW service engaged with a significantly higher proportion of females (60%) compared to males (29%). Nearly 80% of residents lived in either St Austell or Newquay, which is expected as 60% of the overall CHWW resource is located in these two locations.

Category		(%) whole dataset N=211
Gender (n=190)	Male	29%
	Female	60%
	Binary or Trans	1%
	Not stated	10%
Area of Cornwall	Newquay	42%
	St Austell	36%
	Falmouth	13%
	Truro	10%

Who completes the MYCaW form?	<i>CHWW supporting a resident</i>	66%
	<i>Resident on their own</i>	33%
	<i>Carer about themselves</i>	0.5%
	<i>Carer supporting a Resident</i>	0%
Type of consultation	<i>Face-to-face</i>	76%
	<i>Via phone or video</i>	7%
	<i>Not stated</i>	18%

Table 1. Summary of demographic and consultation information (note percentages may not add to 100% due to rounding).

3.2 How was the MYCaW form complete and by whom?

We recommend that MYCaW is completed by the residents themselves, but that is not always possible for a variety of reasons. We therefore also recommend that a professional can complete the form for the resident ensuring that they write down verbatim how a resident describes their concerns. Sometimes the person completing the form is a carer supporting the resident and occasionally they are a carer completing it form themselves.

In Table 1 it shows that two thirds of the MYCaW forms were completed by the CHWWs supporting the residents and one third of MYCaW forms were completed by the residents themselves.

It can be useful to know what type of consultation was carried out, e.g., was it face-to-face, via phone, online etc. Face-to-face was by far the most frequent consultation method (76%).

3.3 Measure Yourself Concerns and Wellbeing® score changes

All evaluation entries that had baseline scores and matching follow up scores were deemed to have ‘paired’ data and were used in the analysis of MYCaW® Concerns and Wellbeing score changes. For the analysis of Concern 1 score changes there were 147/211 (70%) paired data entries to analyse. Concern 2 is an optional part of the questionnaire and there are always fewer entries for Concern 2 compared to Concern 1. There were 130/211 sets of paired data for Concern 2. 148/211 people provided a baseline and follow-up wellbeing score.

Table 2 outlines the score changes, statistical significance levels and the relevance of these score changes. Both MYCaW Concerns scores and Wellbeing scores showed an average reduction in the severity at follow-up. To determine if these score changes were down to change or were likely to have been influenced by the support from the CHWWs, a two-tailed, paired T-Test was carried out. All score changes were highly statistically significant, meaning that there was a less than 5% probability that the score changes were due to chance.

MYCaW©	Baseline score (±SD)	Follow-up score (±SD)	Score change (±SD)	P value	% minimal positive important difference	% minimal negative important difference
Concern 1 (n=147)	5.4 (±0.8)	2.8 (±1.5)	-2.6 (±1.6)	≤0.001	91%	1%
Concern 2 (n=130)	5.3 (±0.8)	2.9 (±1.5)	-2.4 (±1.5)	≤0.001	88%	3%
Wellbeing (n=148)	4.8 (±1.1)	3.0 (±1.2)	-1.8 (±1.3)	≤0.001	90%	4%

Table 2. Summary of the MYCaW scores and changes over time. A negative score change denotes an improvement.

To further understand the relevance of the score changes we calculated what proportion of score changes met the minimum threshold level of change. This level of changes (minimal important difference) is the level of change needed for a demonstrable difference to be experienced by the participant. It is similar to asking the question “*so what does the score change really mean?*”

We can see that 91% and 88% of the Concern 1 and 2 score changes (respectively) met this positive threshold level, and 90% of the wellbeing score changes. These are extremely high percentage improvements compared to other MYCaW datasets. Overall, the average scores indicate that for the vast majority of people who were supported by the service, their concerns and wellbeing improved as a result.

To ensure that these results are fully representative of the people who complete baseline MYCaW questionnaires it is important to collect as much follow up data as possible. This score data presented here represents 70% of everyone who had taken part in this demonstrator project at the time data was submitted for analysis to Meaningful Measures Ltd. As the service is ongoing, there will be a proportion of people who hadn't had chance to have their follow up data collected before it was analysed as an arbitrary cut-off point was agreed upon. This largely explains why the data is not 100% representative of the residents using the service. Some people may not want to be followed up, and often people may move, return to work or just be unavailable for a further consultation, when the follow-up data needs to be collected.

3.4 Analysis of Measure Yourself Concerns and Wellbeing® concerns

Data for 211 people was received by Meaningful Measures Ltd to analyse. For the qualitative analysis of the concerns, there were a total of 412 concerns provided when combining Concern 1 and Concern 2. All of the concerns stated were coded using an existing framework of concerns developed by Meaningful Measures Ltd. This framework was based on over 2,500 concerns from people around the UK using personalised care services.

Initially all concerns are designated a theme and the proportion of concerns in each theme is shown in the pie chart below - Figure 3.

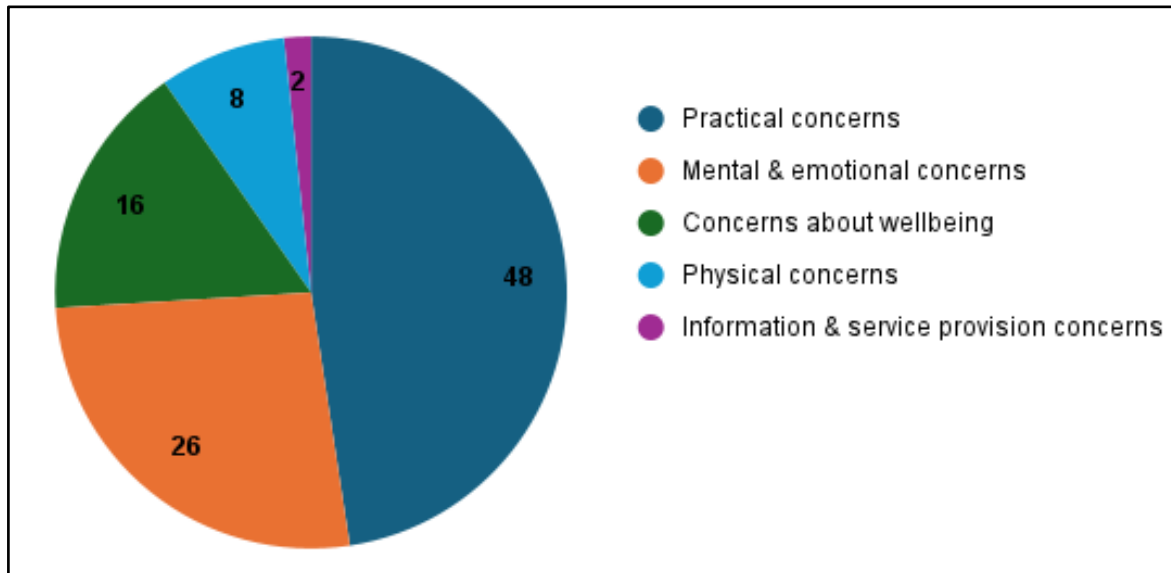


Figure 3. The percentage frequency of concerns falling in each main theme.

Almost half of the concerns were practical ones, which included problems with finance, housing, managing the household admin, managing to look after ones-self, social justice and safety issues, work & training, school and education and carer burden. A quarter of concerns related to a broad range of mental and emotional issues.

Across the whole dataset we identified the top 5 individual concern categories reported - shown below in Table 3. It is worth noting that most of these concerns are social determinants of health and it is, therefore, not surprising that people's mental health is being affected.



Category	n (%)
Finances	68 (17%)
Housing	52 (13%)
Mental health	40 (10%)
Loneliness and isolation	28 (7%)
Access to food	22 (5%)

Table 3. The top 5 concerns most frequently reported.

3.4.1 Practical Concerns

The theme about practical concerns was split into 8 specific categories. Figure 4 shows the breakdown of the frequency of concerns in each category. Below is some description and examples of concerns in each of the categories.

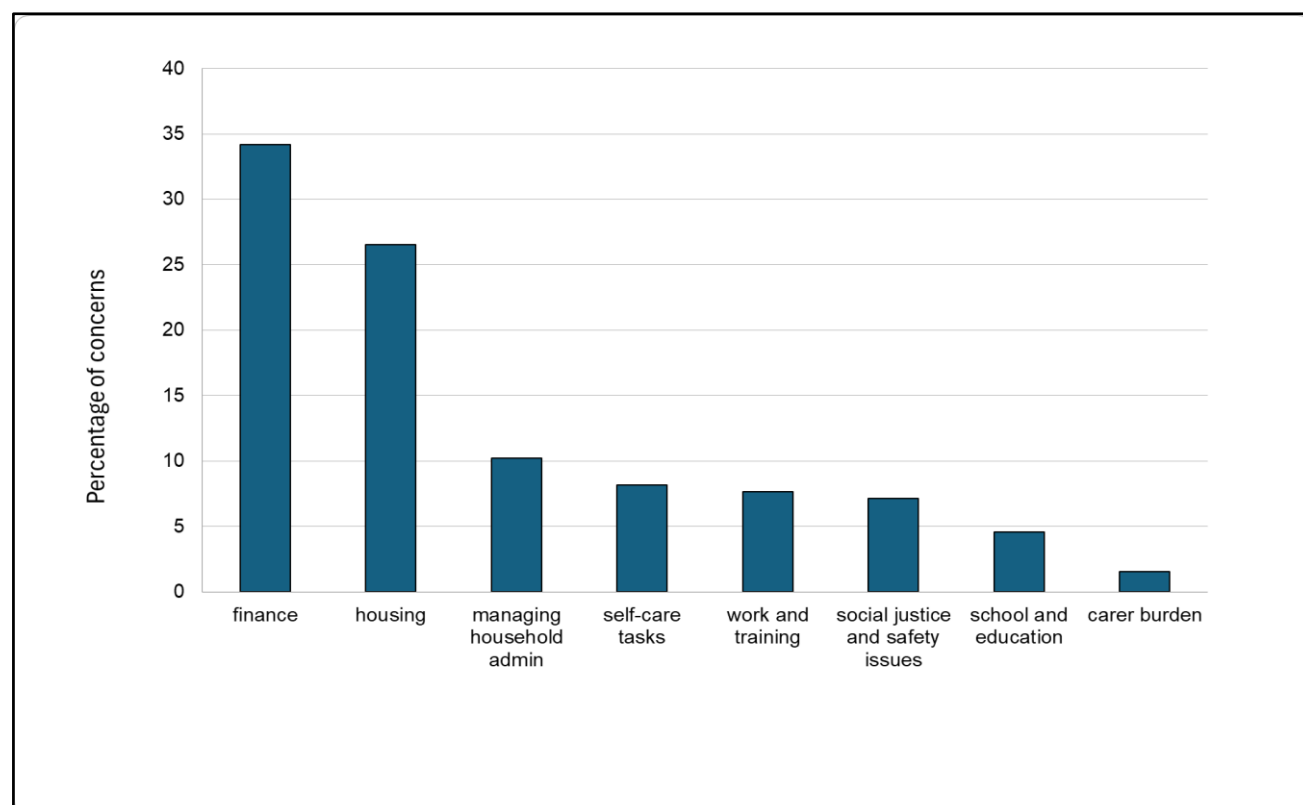


Figure 4. Breakdown of practical concerns.

3.4.1.1 Financial concerns

The most frequent practical concerns (n=67, 34%), were finance based. Approximately 45% of all finance concerns did not specify what the actual finance concerns related to but stated 'concerned about finances' or 'finances' / 'finance support' / 'financial struggles'.

25% of people with financial concerns were in debt, with a range of organisations including the housing association, energy companies, DWP, council tax, or water company e.g.

"My Council Tax bill - keep getting threatened with the bailiffs, but I just can't afford it"

"Bedroom Tax, [housing association] and DWP all stressing me out. They all reckon I owe them money!"

"Electric bill, they changed how I paid it & I have ended up in debt with it now. I want to go back to how it was before, I knew where I was"

Some people weren't in debt but were finding it hard to make ends meet

"Am struggling to afford food for my family."

About 20% of people with financial concerns needed help accessing benefits and welfare payments, such as attendance allowance, carers allowance, personal independent payments or support for their children e.g.,

"I feel like a really bad Mum, I am trying to get my youngest the support he needs. He needs Nursery and his portage, they won't fund both"

3.4.1.2 Housing concerns

26% (n=52) of practical concerns related to housing. Approximately 20% of these concerns just stated 'housing' so it was not clear what the specific issue was.

27% of the housing concerns related to the state of housing, where repairs were needed, people had no flooring, mould growing on the walls and often issues with their heating.

"The damp and mould in my flat is making me ill. It effects my asthma. Can't open the windows, might let rats in again".

"I'm so cold. Just can't get warm. Heating boiler has been condemned. I had a heart attack every February for past 6 years due to the cold".

"Having problems with my flat, landlord turns the heating off when he leaves for months".

A further 27% of the housing concerns related to wanting to change where a person lived - some people were living in overcrowded conditions, and others wanted a smaller house or a house to suit their additional access needs. Some people wanted to move closer to family, others wanted to move away from the negative impact of their family. A small number of people were homeless, in emergency accommodation or had been served with Section 21 eviction notices.

"Overcrowding in housing, want a house near to mum to help support with childcare and health"

"Moving to a bungalow as my partner has to sleep downstairs"

"Housing - need new home that's accessible for just me. Restricted on what I can eat, when I can use bathroom/wash clothes etc by [family member]".

Just over 15% of people had housing concerns relating to furniture. Most people needed items of furniture for themselves or their children or babies. A few people needed help to clear out and declutter.

3.4.1.3 Concerns with managing the household admin

10% of people with practical concerns were struggling to manage their household admin. Some concerns were quite general, stating 'form filling' whereas other concerns gave a more detailed picture of some people were overwhelmed with the paperwork that had built up, or where they needed support to apply for particular things, e.g. a blue badge or PIP payments.

"I need help with lots of bills and paperwork, it's all overwhelming."

"I need help accessing my ID like birth cert or bank statements."

"I need to apply for PIP but keep putting it off."

3.4.1.4 Concerns about self-care task

7% of the practical concerns were about the ability of people to care for themselves. Some concerns stated that more support at home was needed, other concerns explained that people needed help with cooking, cleaning or accessing their medication, because of a disability or due to the impact of having treatment for a health condition. As well as looking after the house, some people needed support with their gardens which were overgrown.

"I need help to cook as I have a disability and can't use kitchen equipment."

"I need help with cleaning, since my prostate cancer I have remained unwell and can no longer bend to pick things up and clean things."

"I cannot get to the pharmacy and need my medication delivered, I cannot be without it. Can you help? I get anxious making calls."

3.4.1.5 Concerns about Work and Training

7% of practical concerns were about getting a job. Most people just wanted to find a job to have some income. A few people mentioned wanting a job that they could enjoy or would align with their personal values. Some people were also concerned about their level of confidence, feeling that they wanted to improve their confidence to get back to work.

“Looking for apprenticeship.”

“Not enjoying my current work environment. Affecting my mental and emotional health.”

“Confidence building to re-enter employment.”

3.4.1.6 Concerns relating to social justice and safety issues

There were a discrete set of concerns in this theme about social justice and safety. These concerns included concerns about neighbours selling drugs, being antisocial and crime in their buildings, - to the point that people were scared to leave their house or put their bins out. Other people mentioned issues with abuse from family and care workers and sexual harassment.

“My last support worker owes me money and has my house and car keys”.

“Problems with neighbours keeping us housebound”.

“Recent abuse and related court case”.

3.4.1.7 Concerns about school and education

5% of people’s practical concerns related to education - several people wanted advice about furthering their education, one person wanted to complete their studies.

Several people had concerns about their children not being settled at school or skipping school.

“Adult education advice.”

“Children struggling with behaviour and attendance at school.”

3.4.1.8 Carer burden

This was only mentioned by a few people, who wanted to be healthier to manage their caring role better or who were concerned how to protect themselves from the person they were caring for.

Overall, the theme of practical concerns has demonstrated a very diverse range of concerns that the CHWWs have recorded during their consultation. Whilst finance and housing were by far the most frequently mentioned concerns, the concerns ranged from ones about crime and safety, education, employment, carer burden, the ability to look after oneself or the home. In the next section we explore the mental and emotional concerns. It is highly likely that some of the practical concerns are causing mental and emotional concerns.

3.4.2 Mental and Emotional Concerns

These types of concerns were expressed by 26% of all the residents and are detailed in Figure 5. The categories will be explained in more detail below.

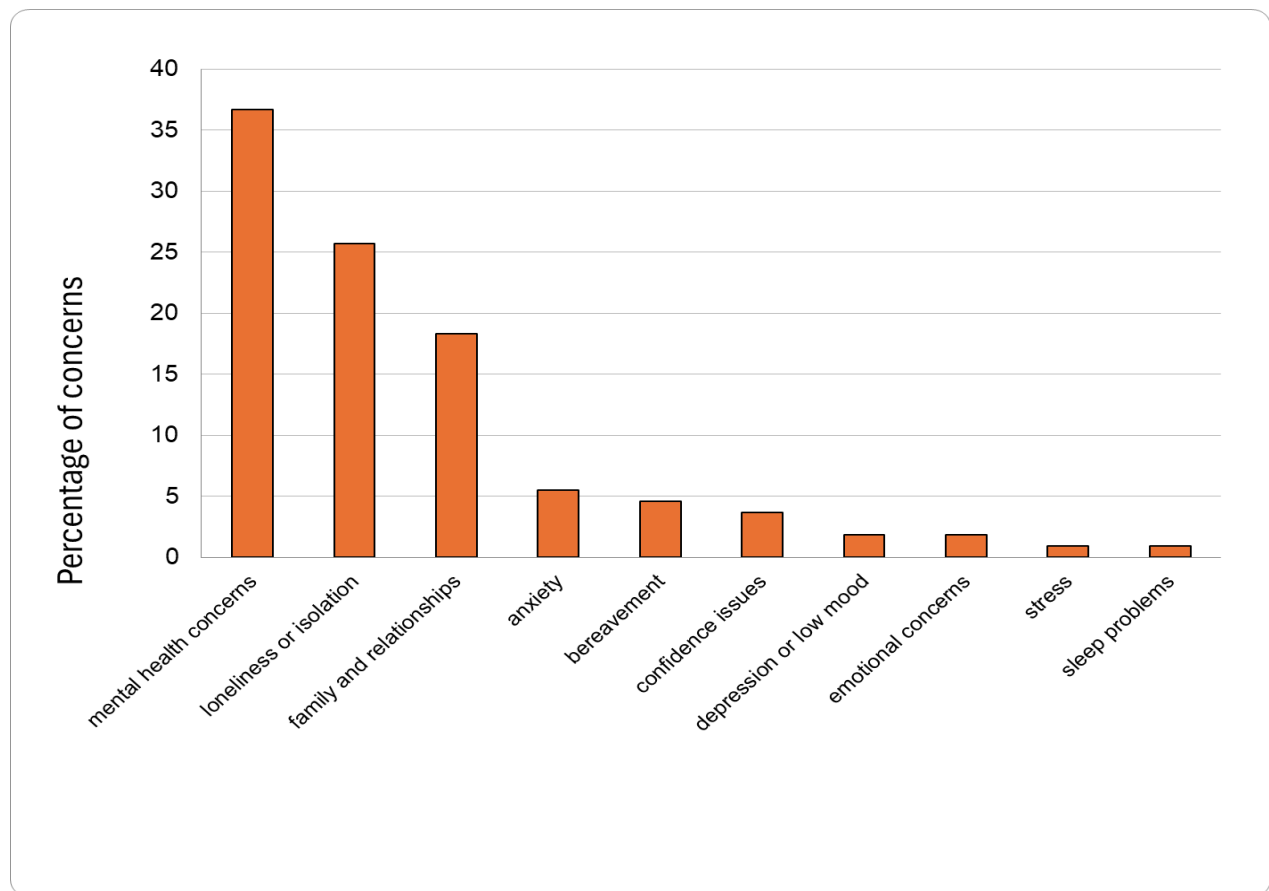


Figure 5. Breakdown of Mental and Emotional Concerns.

3.4.2.1 Mental Health (In general)

The most common category (36%) related to general mental health concerns. Here nearly two thirds of concerns just stated: ‘mental health’ or ‘mental health support’. Where there was more detail, some people explained what was causing their mental health issues e.g.

“Mental wellbeing due to a difficult couple of months.”

“Mental health due to housing and family.”

“Mental health surrounding finances.”

Some people had experienced childhood or past trauma which was still affecting their mental health, others stated conditions such as agoraphobia were impacting their mental health.

A few people also detailed the impact that their poor mental health was having on them, such as starting to self-harm or attempt suicide.

“Things get too much for me and I cut myself.”

“My mental health is bad and I'm finding it hard to do things at home.”

“I feel really useless & helpless, wellbeing is up and down”

3.4.2.2 Loneliness or Isolation

About one quarter of issues in the mental and emotional concerns theme were due to loneliness or isolation. About one third of this group reported they were ‘lonely’ and two thirds mentioned isolation.

“I feel really alone. My daughter is at Uni and I feel alone.”

“I am lonely and isolated, I worry no one knows I exist.”

“Feeling isolated and trapped in flat.”

3.4.2.3 Family and Relationships

This was the third largest reported category of concerns for the mental and emotional theme. Some people were experiencing conflict within the family, and others were concerned for their partners.

“Worried about [name], if something every happens to her I'd be lost.”

Some people were worried about their school aged children whilst other people were worried about their adult children who were in unhealthy relationships, or experiencing homelessness.

“I'm worried about my eldest. It's hard to get him out and about. Can be a struggle getting him to school.”

“I’m really worried about my daughter and baby grandson. She is in rehab at the moment. Worried about her ex getting back in contact with her.”

3.4.2.3 Anxiety

A discrete and small group of people mentioned their main concerns was anxiety - sometimes stating what type of anxiety it was e.g.,

“Bad social anxiety and I have anxiety going places.”

3.4.2.4 Bereavement, depression, confidence issues, stress, emotional concerns and concerns about sleep

All of these remaining concern categories were all only mentioned three or four times and are self-explanatory.

3.4.3 Concerns about wellbeing

This theme of concerns was expressed by 16% of residents and relates to concerns that directly affect a person's wellbeing. The two most frequently report concerns related to access to food and to social interaction. The theme will be explained in more detail below, the frequency of concerns in this section are shown in Figure 6 below.

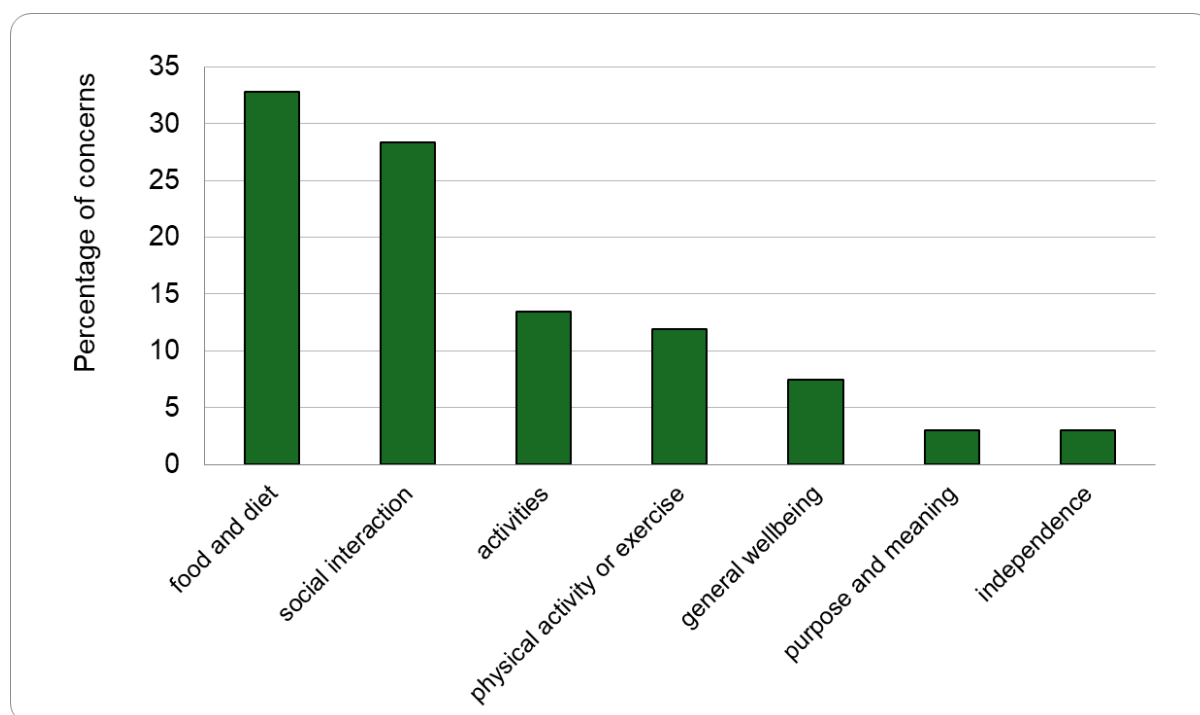


Figure 6. Breakdown of concerns about wellbeing.

3.4.3.1 Food and diet

For many people who nominated food as their concern, they needed support to access food and get enough food for everyone in their household to eat. A couple of people needed a cooker or were unmotivated to cook whilst living on their own.

“Am struggling to afford food for my family.”

“Haven’t got enough food for my family.”

“I need a cooker. I need to be able to cook for and with my daughter. It’s awful not having a cooker.”

3.4.3.2 Social Interaction

Most people in this category wanted to increase the amount of social interaction they were having, which included needing support to find appropriate groups to join and or to have people to talk to regularly, which may not necessarily be through a community group but could be via a befriender

“I want to be more social and do something for myself.”

“Find some more social connections.”

3.4.3.3 Activities

In a similar vein to the social interaction category, another group of people specifically wanted to join a group to do an activity either for themselves or for their child.

“Help accessing toddler groups.”

3.4.3.4 Physical activities or exercise

The small group of people in this category wanted to start physical activities such as going to the gym, so they could improve their strength and get fitter.

“Need to go to the gym. It's good for my RA. Need to get stronger.”

3.4.3.5 General Wellbeing, Purpose and meaning, Independence

There were only a few mentions of each of these categories of concerns and they are reasonably self-explanatory. Some people just wanted to feel healthier or happier, and generally have better wellbeing. A couple of people were concerned about their purpose in life and sense of belonging, and a couple of people's concerns related to their level of independence.

“I want to feel happier.”

“I want to start my own business. I need a sense of purpose again.”

“Remain independent.”

3.4.4 Concerns about physical health

Only 8% of residents put concerns about their physical health as their priority. It is likely that the residents are experiencing far more physical concerns but that the concerns about finances, access to food and housing were a greater priority.

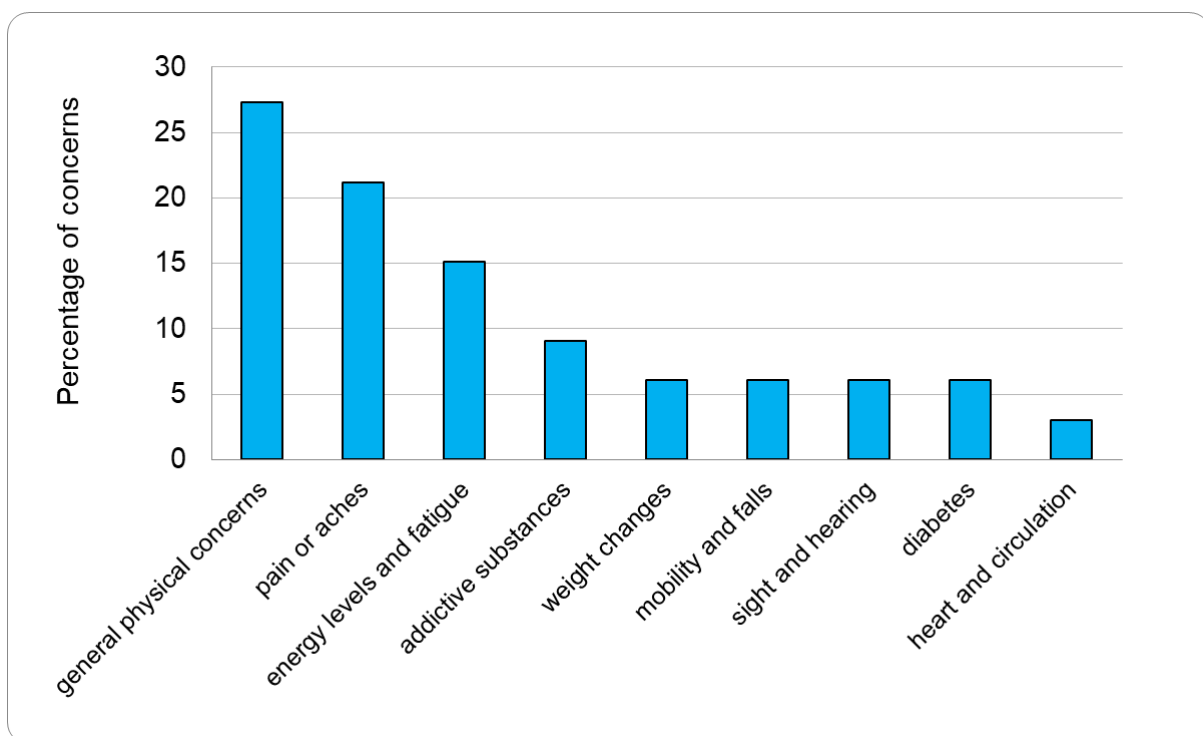


Figure 7. Breakdown of concerns about physical health.

For the largest group of people in the sample, these concerns were described just as ‘health concerns’ so there wasn’t enough detail to discern what they related to specifically.

There were then a range of physical concerns about weight, suffering with low energy levels and pain,

“Would like to lose weight as I think it will help me with my social anxiety.”

“I don’t want to feel so tired and in pain all the time.”

Other concerns each mentioned by one or two people related to issues with mobility, diabetes, sight, addiction.

"I am struggling with mobility and accessing activities."

"Diabetes and knowing what to eat."

"I am losing my sight and need help at home."

"I drink a lot of alcohol, I need some help to cut down so I don't fall as much."

3.4.5 Concerns about information and service provision

The final theme of categories relates to people needing specific support or information about particular services.

Only seven people had their priority concerns in this theme. Generally, people wanted very specific information about services e.g. support about adoption, support after a stroke, information on how to get a dentist, find a chiropodist and information about autism services.

3.5 What else is going on in your life?

On the follow up MYCaW form, we ask people if there is anything else going on in their life that is important, besides the concerns they have nominated. This data identifies other 'confounding variables' that can affect score changes and give more insight into the reality of residents. Of the 211 people who completed MYCaW, 130 (62%) people recorded a total of 139 other things going on in their life.

We can split the additional issues into positive and negative events, to give a sense of challenges still faced or improvements that are happening. Table 6 below shows that 42% of additional events were positive in nature, 49% were negative in nature. It was not clear whether the remaining 9% of comments were positive or negative in nature.

The biggest positive change for people (9%) was improvement in their finances or work situation. It was implied that the CHWWs had helped get benefit payments sorted, but predominantly people had either found a job that they were happy with or were actively looking for work now.

residents were also trying to improve their wellbeing through diet and physical activity (6%), were actively improving their environment (6%) and were benefiting from the increased support from the CHWWs as well as friends and family (6%). A further 6% of residents were getting more social activity and connections. All of these situations are known to have a positive impact on a person's health and wellbeing.

Sadly, the most frequent negative issue was ongoing issues with physical and mental health (20%) and this often related to issues with long term conditions and pain. A further 9% of residents were struggling with their finances and work issues, e.g., lack of work. The rest of the additional issues having a negative impact can be seen at the bottom of Table 6 below.

Research previously carried out has shown that in many cases where negative additional issues exist, it can make the overall score changes smaller for concerns and for wellbeing. Conversely where additional situations are positive, people may have larger score changes. We have not carried out this analysis for this piece of work, but please note that the MYCaW score changes are already showing a large effect size, so it is likely that these additional issues are not impacting the scores too much.

Impact	Category	n=139 (%)	example quotes
Positive impact	Work or finance	9 %	<i>"CHWW 's got me attendance allowance and HHSF to purchase some white goods". "returning to work"</i>
	Awareness of own wellbeing	6	<i>"I have made changes to my diet and stuck to this which has helped with my pain".</i>
	Positive change in environment	6	<i>"Really starting to get things sorted now with the decluttering and decorating. Slowly but surely, but I feel better mentally the more I do. We will have a much nicer home after all this".</i>
	Increased support	6	<i>"Your support has helped me to believe in myself more. Putting me in touch with The People Hub has really helped too"</i>

	<i>Increased social activity</i>	6	<i>I have made many social connections, through the neurodiversity group, chill and chat and i have even performed with my guitar for a singer at a concert. i have also started a level 2 autism awareness course which is very interesting".</i>
	<i>Taking physical activity or exercise</i>	3	<i>"Going to the gym is really helping - I am feeling stronger and I am managing to pay for it now at the reduced rate. I had to continue, because it does me good, mentally and physically"</i>
	<i>Improved family situation</i>	3	<i>"feel better now that my child is at nursery is at nursery".</i>
	<i>receiving support from health professionals</i>	3	<i>"getting help with pain management"</i>
<i>Negative impact</i>	<i>Continuing issues with health and long term conditions</i>	20%	<i>"I have depression and chronic back pain, so some days I cannot do anything. I don't even get up. It's so bad. I want to be doing things but I can't when I am in so much pain".</i>
	<i>Work or finance</i>	9%	<i>"Loss of employment". "rent arrears"</i>
	<i>Family issues</i>	6%	<i>"Fathers mental health concerns"</i>
	<i>loneliness and negative wellbeing issues</i>	4%	<i>"loneliness" "Things just get on top of me. Everything seems to go wrong. Haven't been taking my medication".</i>
	<i>Negative change in environment</i>	4%	<i>"Currently homeless and trying to find myself a flat".</i>
	<i>Bereavement</i>	3%	<i>"Recent bereavement over my wife, sometimes I am still very sad and lonely".</i>
	<i>Carer burden</i>	3%	<i>"I have a son with autism and a wife who suffers with seizures. I do worry about my son when we are no longer around".</i>

Table 4. What other things are going on in your life? - split by positive and negative impact.

3.6 What is important about the support you have received?

The final follow-up question on MYCaW asks what a person found most important about the service or support they received. 142/212 people responded to this question, stating 202 things that were important about the support they received from the CHWWs. Responses were categorised and are displayed in Figure 8 below.

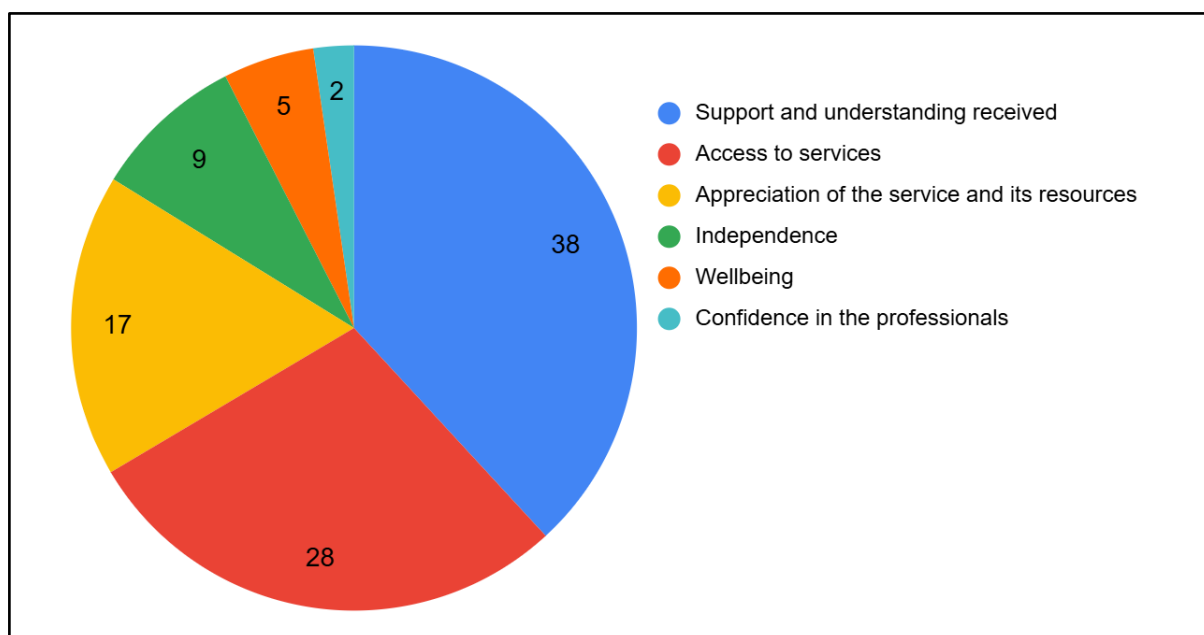


Figure 8. Percentage responses for people in response to “What was the most important thing about the service?”.

3.6.1 Support and understanding received

The most frequent response (38%) related to the support, understanding and provided by the CHWWs as well as the encouragement to make changes or improvements. Many householders were really appreciative of having CHWW to talk with, who listened to them and believed them. For many people this has helped them trust the CHWWs in a way that they don’t normally trust professionals.

“It got things sorted, but so much more to do. I don’t trust easily, but I can talk to you.”

“Having someone call in to see me just to check in and have a chat has been really great. My dog loves seeing the CHWW too as she never sees anyone else! I am very grateful!”

“I really do appreciate the time you spend listening to me going on about my funny little ways and my family issues. I don't have anyone else who can give me this time.”

3.6.2 Access to services and organisations

The second most frequently appreciated aspect of the CHWW service was getting access to services and additional support organisations. Often this included getting more financial advice and support as well as support regarding housing issues or the housing association. Additionally, people were put in touch with organisation that would provide social activities. Furthermore, several people are grateful that they now have a source of support they can contact in the future if they need it again.

“Having financial support has been so helpful, I could not have done this on my own. I was getting depressed and exasperated by paying bills and worrying about debt. The CHWW changed all that for me.”

“The Help I got from the CHWW has made a huge difference. They got me in touch with Safer Truro, and secured my home which made me feel a lot safer. I was listened to and helped immediately. Even helping me reduce some of my bills. I can't thank them enough.”

“Good to know support is there if I need it.”

3.6.3 General Appreciation of the Service

Many people appreciated the service in general citing a range of things they valued about it e.g.

“Having someone to listen and believe me. The CHWW met me at home and walked with me to groups and built up my confidence. Being able to be part of a social group again at my age, and getting to know my neighbours. All this would not have happened without the CHWW.”

There was a lot of appreciation for the availability of the access to the CHWW service, whether it be the home visits, the drop-in sessions that were provided or the regular contact that people had with the CHWWs

“The help I received from the CHWW has been a real help in my life. Knowing they are there open-ended if I need them, knowing they are on the end of the phone. I can rely on them to turn up when I feel everyone else lets me down.”

“Being able to drop in to session.”

“Someone who wanted to help me and actually actioned it! Being able to have the appointment at home, because I probably would have cancelled if it wasn't.”

3.6.4 Being more Independent

A small group of people (9%) were very clear that the support from the CHWWs had given them motivation and independence which in some cases has led to them being more active and having a much better social life.

“Giving me the confidence to do things myself and encouraging me and making me feel like I've got this! The wheelie bin numbers you got me helped too - people know which bin is mine now and they haven't taken it”.

“I am hardly ever home now, I go out a lot, have made friends and even played my guitar on stage with a band. My life has changed a lot really”.

The final areas mentioned by a few people related to them having improved wellbeing in general and also their trust in the professionals that they are working with now.

Overall, this section of data shows that the CHWW service is accessible to people, and the CHWWs are able to develop trusting relationships with the residents who are then going on to share their concerns and receive the support they need.

4.0 Discussion

This analysis aimed to understand the types of concerns that people wanted support with when using the CHWW service. Furthermore, we wanted to understand if there was any impact of the CHWWs that could be discerned from the MYCaW data.

As a proof of concept, the concerns collected alongside the score changes in concerns and wellbeing at follow-up show that this is an extremely valued and important service in Cornwall and IoS.

The types of concerns being recorded were predominantly those associated with the most basic of needs such as having access to food, financial issues and debt, problems with housing conditions and issues with social welfare and crime - it is well documented that people who are the most socioeconomically deprived have more of these issues which are associated with poverty and destitution. The findings of this report make for very sober reading and show the difficult lives of people living in Cornwall and IoS. This report echoes the issues of deprivation and destitution outlined in the introduction.

The average changes in MYCaW scores after receiving support from the CHWWs were very large and highly statistically significant. Whilst we can't attribute all of the change in scores to the CHWW service (a comparison group is needed for this), it is clear from the additional follow-up questions, that there are several key aspects to the service that are important to the residents.



Firstly, people appreciated the style of access to the service, having home visits, as well as drop-in session and continuity of support. A key enabler appears to be the ability of the CHWWs to establish trust with the householders, which led to them feeling they could talk to the CHWWs. Trust was further developed as people felt they were genuinely being listened to and then getting the support they needed. The type of support needed was often financial in nature, as well as helping people sort out

housing issues, and get out to social activities. Another aspect mentioned several times was that the CHWWs accompanied people to their activities for the first time and that they did what they said they were going to do.

It was noticed on the data collection spreadsheets that CHWWs from different locations may enter different information into the spreadsheet - particularly when reporting who completed the MYCaW form and the type of consultation. For future data collection and analysis, particularly with a larger dataset, it would be beneficial to add a drop-down menu choice on some of these questions, making it easier and quicker for data to be entered and improving the overall consistency of data collection.

Other tweaks to the data collection include ensuring that the concerns are written down as the residents talk about them, instead of just summarising their concerns with one or two words. There is a lot of detail which helps to demonstrate what matters to people and how the service is helping which is lost if a CHWW summarises the concern with one or two words only.

Finally, all scores should only be a whole number however there were some scores that were not whole numbers and therefore it is important that all CHWWs are clear on ensuring whole numbers only are entered.

5.0 Conclusion

The data analysis here shows that the CHWW project is a highly successful and valuable service supporting the poorest people in Cornwall and IoS to access what are considered their basic needs and rights to appropriate finance, housing, food and support with their physical and mental health. Given the statistical significance of the MYCaW score changes we conclude that this service should be continued, and more data collected to establish the longer-term impacts on the current residents.

6.0 Appendix 1: Discussion of Measures

In terms of the second aim of the evaluation, the data obtained from the MYCaW analysis can possibly indicate where other person reported outcome measures may be used going forwards. The most often picked MYCaW super-category was “practical concerns” and here the main issues were: problems with finance, housing, managing the household admin, managing to look after ones-self, social justice and safety issues, work and training, school and education and carer burden. Most of these concerns are social determinants of health. Other outcome measures for health and wellbeing may not be able to reach all of the areas that MYCaW reaches, especially in terms of social determinants of health. Table 5 below is adapted from Polley et al. (2020)¹⁴ which reviewed outcome measures for social prescribing. It is hoped that this information helps the reader in working out best ways forwards with outcome measures which may accompany MYCaW. Most useful measures are those which are short, free, sensitive and provide economic value data (the best tools are coloured in green).

There are several tools that can be used to derive economic data that we have listed below. The caveat here is that you want to use a tool that measures changes in the broad range of domains identified via MYCaW. If you only capture data on a proportion of domains that we know are relevant to residents, the score changes may not be statistically significant at follow-up and may not accurately represent the impact of the CHWW service. The ensuing economic calculations may also, therefore, not accurately represent the service. This would be seen as a false negative result.

Going forwards, therefore, it is important to test out any other outcome measures you are using to see if they are sensitive to the change your residents are experiencing. The change in wellbeing score from the MYCaW data would be a good benchmark for comparison. We know that people are rating their wellbeing as improving on average,

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https://www.london.gov.uk/sites/default/files/a_guide_to_selecting_outcomes_measures_in_social_prescribing_final.pdf

therefore, any other tool monitoring wellbeing or quality of life should also pick up this change. If it doesn't then it may not be suitable for your residents.

Another point to consider is maximising the amount of follow-up data captured. The higher the proportion of people who have baseline and follow up scores, the more representative your data is. Many studies have shown economic calculations to show return on investment, but the follow up data represented less than 50% of all residents. The conclusions, therefore, should read that in those 50% of residents there is a positive return on investment. But if the other 50% did not show a return on investment, overall the final result could be very different - but it remains unknown. Aiming for at least an 80% representation with follow-up data would be acceptable. Maximising the amount of follow-up collected data is always a good investment of time when evaluating services!

Finally, note that we have not included outcome measures that are specific to conditions or situations, e.g. pain or loneliness as people's concerns are far ranging. If you want to use an outcome measure that is specific to a condition or situation, e.g. loneliness or mental health, you could use the MYCaW concerns to guide you on when to use the other tool. For example, if people say their main concerns relate to their mental health, then a mental health scale would be appropriate. Similarly you could use a loneliness scale on people who have identified concerns about loneliness. This is a slightly more complicated approach to monitoring outcomes and requires buy-in from and training for those administering the outcome measures.

Table 5 Comparison of outcome measures used to monitor wellbeing or Health Related Quality of Life

Tool	Description	Licence	Notes
Measure Yourself Concerns and Wellbeing (MYCaW)	Concerns and Wellbeing. Person centred data. In line with the findings of this report	Yes, currently in place with the Cornwall & Isles of Scilly ICA	Suggest to use this ongoing as it reaches all areas of need and experience in the community. Provides personalised data. Does not provide economic value data (although may do in future).

36-Item Short Form Health Survey (SF-36)	36 items measuring eight domains of health: physical and mental aspects, including physical functioning, role limitations, pain, general health, vitality, social functioning, and emotional well-being.	No, free tool	Researchers often use the SF-36 to evaluate health-related quality of life and determine the cost-effectiveness of health treatments in health economics. Does not provide personalised data.
Office for National Statistics personal wellbeing scale (ONS4)	The Office for National Statistics (ONS) uses a set of four questions to assess personal well-being, often referred to as the ONS4. These questions capture three types of well-being: evaluative, eudemonic, and affective experience.	No, free tool	This tool is useful as you can benchmark against national datasets to show how a particular population compares. Some questions may be triggering in a negative way. Does not provide personalised data. Can be used in some economic analysis, but does not provide QUALYs
EQ-5D (EQ-53-3L) European Quality of Life 5 Dimensions 3 Level Version	Tool assesses five dimensions of health: mobility, self-care, usual activities, pain/discomfort, and anxiety/depression.	No, free tool	Perceived by some to be a bit reductionist and the domains do not cover the determinants of health very well. Does not provide personalised data. Can provide economic values, can be used to calculate QUALYs
DIALOGUE +	The DIALOG scale consists of 11 questions. People rate their satisfaction on a 7-point scale with eight life domains (such as relationships, daily activities, and housing) and three treatment aspects (like medication and	No, free tool; available within the NHS, also available via an app.	Does not provide personalised data. Does not provide economic value data. The tool is used to help mainly with mental health issues.

	therapy). Tool also extends beyond the scale and allows people to discuss concerns and then addressed them in a four-step approach (understand, identify, explore and agree action).		
The Quality of Life Scale (QOLS)	The Quality of Life Scale (QOLS) is a 15-item instrument that measures five conceptual domains: material and physical well-being, relationships with other people, social, community and civic activities, personal development and fulfilment, and recreation.	Free for students, fees apply for other projects	Used for long term illness, but can be used in non-illness contexts. Does not provide personalised data. Does not provide economic value data.
Patient Reported Outcomes Measurement Information System (PROMIS)	PROMIS is a system of highly reliable, precise measures that assess patient-reported health measures. Domains are: physical, mental, and social well-being.	No, free tool	Sensitive and adaptable, and applicable widely to lots of different contexts. Does not provide personalised data. Does not provide economic value data.

Report End

Communications and Engagement Plan – Macmillan Project

Background

Hywel Dda University Health Board (HDdUHB), in partnership with Macmillan Cancer Support and a wide range of statutory, voluntary and community sector organisations, is developing a Neighbourhood Prevention Programme to improve health and wellbeing outcomes and reduce health inequalities across Mid and West Wales. The programme responds to increasing levels of cancer and long-term conditions, rising pressures on health and care services, and the need to strengthen prevention, early intervention and community-based support.

The programme will deliver a neighbourhood-based model comprising **Health Coaches, Community Health and Wellbeing Workers, and Community Capacity and Local Infrastructure for Prevention**. Together, these components will support individuals to improve their health and wellbeing, identify and address unmet need earlier, strengthen community resilience, and improve access to local support and services. The programme is aligned with the principles of Community by Design, the Social Model of Health and Value-Based Healthcare, A healthier Mid and West Wales- Healthier Lives, Well Lived recognising that health outcomes are influenced by social, economic and environmental factors as well as clinical care.

Subject to approval by the Board at **Hywel Dda University Health Board and Macmillan**, mobilisation is anticipated from autumn 2026, followed by phased implementation Health Board wide.

This Communications and Engagement Plan set out the approach to engaging stakeholders, partners and communities throughout programme development, mobilisation and implementation. Recognising that the programme will evolve through learning, co-production and local delivery, a **continuous engagement approach** will be adopted to ensure stakeholders remain informed, involved and able to influence delivery. Regular engagement and feedback mechanisms will be used throughout the life of the programme to support continuous improvement, strengthen local ownership, and maximise the long-term sustainability and impact of the programme.

Engagement methods

Method	Rationale
Engagement period	A period of engagement will take place from July 2026 – September 2031
Broad Approach	Communications and engagement activity will be delivered as a continuous and phased programme throughout the development, mobilisation, implementation and evaluation of the Neighbourhood Prevention Programme. The approach recognises that the model will evolve over time through co-production, learning and continuous improvement, requiring ongoing engagement with stakeholders, partners and communities. During the business case development phase, engagement will focus primarily on internal stakeholders, strategic partners and potential delivery organisations to inform programme design and secure organisational commitment. Subject to approval by Hywel Dda University Health Board and Macmillan Cancer Support, communications and engagement activity will expand to include staff, partner organisations, communities, elected members and the wider public through a coordinated programme of stakeholder engagement, community involvement, media activity and digital communications. A neighbourhood-based co-production approach will be adopted throughout implementation, ensuring that local communities and partners help shape delivery within each neighbourhood while maintaining a consistent programme framework: • October–December 2026: Co-produce and mobilise Neighbourhood 1 with local stakeholders, communities and partners. • January 2027: Neighbourhood 1 goes live. • 2027: Continuous engagement, learning and evaluation to inform refinement of the model while co-designing Neighbourhood 2. • January 2028: Neighbourhood 2 goes live. • 2028: Ongoing engagement, evaluation and shared learning across Neighbourhoods 1 and 2 while co-designing Neighbourhood 3. • January 2029: Neighbourhood 3 goes live, achieving full programme implementation.

	• 2029–2031: Two years of continuous engagement, evaluation and benefits realisation to test, refine and demonstrate the effectiveness, impact and sustainability of the neighbourhood prevention model. This continuous engagement approach will ensure that stakeholders remain informed and involved, communities will have meaningful opportunities to influence local delivery, and learning from each phase is used to improve subsequent implementation. It will support local ownership, strengthen partnerships and help build the evidence required to demonstrate and sustain the model over the longer term.																						
Targeted Engagement	Specific activity for the project will include: <table> <tr> <th>Stakeholder</th><th>Activity</th></tr> <tr> <td>Macmillan</td><td>Programme meetings</td></tr> <tr> <td>Hywel Dda Executive</td><td>Executive briefings</td></tr> <tr> <td>Public Health</td><td>Design workshops</td></tr> <tr> <td>Primary Care</td><td>Cluster engagement</td></tr> <tr> <td>Local Authorities</td><td>Partnership meetings</td></tr> <tr> <td>Third Sector</td><td>Co-design sessions</td></tr> <tr> <td>Regional Partnership Board</td><td>Strategic updates</td></tr> <tr> <td>Community organisations</td><td>Local engagement sessions</td></tr> <tr> <td>Staff</td><td>Internal communications</td></tr> <tr> <td>Communities</td><td>Co-production</td></tr> </table>	Stakeholder	Activity	Macmillan	Programme meetings	Hywel Dda Executive	Executive briefings	Public Health	Design workshops	Primary Care	Cluster engagement	Local Authorities	Partnership meetings	Third Sector	Co-design sessions	Regional Partnership Board	Strategic updates	Community organisations	Local engagement sessions	Staff	Internal communications	Communities	Co-production
Stakeholder	Activity																						
Macmillan	Programme meetings																						
Hywel Dda Executive	Executive briefings																						
Public Health	Design workshops																						
Primary Care	Cluster engagement																						
Local Authorities	Partnership meetings																						
Third Sector	Co-design sessions																						
Regional Partnership Board	Strategic updates																						
Community organisations	Local engagement sessions																						
Staff	Internal communications																						
Communities	Co-production																						

Activity log

Pre-engagement activity				
Date	Event / Activity	Staff / resourcing required	Comments	Achieved by
June 2026	Stakeholder mapping / confirm stakeholder list	RE/JB		1/7/ 2026
June 2026	Draft focused communications and engagement plan	RE/JB	Draft to be included with Business Case	20/7/26
July 2026	Stakeholder workshops with third and statutory sector	JB/BB/RE		24/7/26

Phase 1 – Business Case Development**Purpose – build understanding and confidence in the proposed model, test assumptions and secure support for the business case****Engagement commences – July - August**

Date	Event / Activity	Staff / resourcing required	Comments	Achieved by
July	Develop communications pack (programme overview, FAQs, key messages)	BB/JB		July
	Develop stakeholder narrative and key messages	Programme team		July
	Stakeholder workshops	JB		15/7/26
	Executive briefings	RE/BB		Ongoing/July
	Socialise the emerging model with strategic stakeholders	BB/JB	Key contacts – Local Authority	July
	Steering group meetings	BB/RE/FJ		Ongoing
	One -to- one conversations with key partners	BB/JB		7/8/26
	Draft business case	RE		20/7/26
	Develop governance proposals	Programme team		
	Agree communications approach for mobilisation	BB/RE		September

Phase 2 – Mobilisation and service design for neighbourhood 1

Date	Event / Activity	Staff / resourcing required	Comments	Achieved by
Oct-Dec 2026	Partner workshops	Public Health/Programme lead	Focus of engagement: • Confirm delivery sites/neighbourhoods • Agree governance arrangements • Develop implementation timetable • Establish local implementation groups	November
	Clinical engagement with GP surgeries	Public Health/Programme lead/Primary Care lead		
	Community sector engagement	Programme lead/CVC representatives/Community partnerships lead		
	Local authority engagement	Public Health/Programme lead/Local Authority partners		
	Develop implementation plan and timeline	Public Health/Programme lead		
	Communications toolkit	Public Health/ Communications lead/Programme lead		
	Recruitment planning + commissioning structure for external elements	Public Health/ Communications lead/Programme lead		
	Brand development	Public Health/ Communications lead/Programme lead		
	Planning for launch	Public Health/ Communications lead/Programme lead		

Phase 2 – Mobilisation and service design for neighbourhood 1

Date	Audience	Engagement	Staff / resourcing required	Comments	Achieved by
Oct-Dec 2026	Clusters	- Cluster meetings - Clinical workshops - Operational planning - Referral pathway design - Staff communications	Programme lead/ Primary care lead	Purpose of engagement: - Build ownership of the model - Refine service model - Agree implementation - Ensure integration - Prepare workforce	December 2026
	Primary care managers		Programme lead/ Primary care lead		
	Practice managers		Programme lead/ Primary care lead		
	Health Coaches		Programme lead/ Primary care lead		
	Community teams		Programme lead/ Primary care lead		
	People living with and beyond cancer		Programme lead/ Primary care lead		
	Carers		Programme lead/ Primary care lead		
	Underserved groups		Programme lead/ Primary care lead		
	Volunteers		Programme lead/ Primary care lead		

Phase 2 – Mobilisation and service design for neighbourhood 1

Date	Audience	Engagement	Staff / resourcing required	Comments	Achieved by
Community Health & Wellbeing Workers					
Oct-Dec 2026	Communities	- Community workshops - Local listening events - Asset mapping - Co-design sessions - Community reference groups - Partnership meetings	Public Health/Programme lead/Community partnerships lead/CVC representatives/Local Authority partners	Purpose: Engagement to support operational design plus community codesign	December 2026
	Community leaders				
	Third sector				
	Local authorities				
	Community connector teams				
	Housing associations				
	Social prescribing teams				
	People living with and beyond cancer				
	Carers				
	Underserved groups				
	Volunteers				

Local neighbourhood infrastructure					
Date	Audience	Engagement	Staff / resourcing required	Comments	Achieved by
Oct-Dec 2026	CVCs	- Partnership forums - Coproduction workshops - community conversations - mapping existing provision - grant design workshops - governance arrangements - local planning sessions	Public Health/Programme lead/Community partnerships lead/CVC representatives/Local Authority partners	Purpose: Engagement to support broader strategic and community engagement to shape investment, governance and delivery Key: Mapping of existing assets and identifying gaps to underpin investment decisions	December 2027
	Third sector organisations – including social enterprises, charities				
	Town and Community Councils				
	Community groups				
	Volunteers				
	Local Authorities				
	Anchor organisations				
	Regional Partnership Board				
	People living with and beyond cancer				
	Carers				
	Underserved groups				

Phase 3 – Mobilisation and service design for neighbourhood 2

Date	Event / Activity	Staff / resourcing required	Comments	Achieved by
Jan-Dec 2027	Partner workshops	Public Health/Programme lead	Focus of engagement: • Confirm delivery sites/neighbourhoods • Agree governance arrangements • Develop implementation timetable • Establish local implementation groups	December 2027
	Clinical engagement with GP surgeries	Public Health/Programme lead/Primary Care lead		
	Community sector engagement	Programme lead/CVC representatives/Community partnerships lead		
	Local authority engagement	Public Health/Programme lead/Local Authority partners		
	Develop implementation plan and timeline	Public Health/Programme lead		
	Communications toolkit	Public Health/ Communications lead/Programme lead		
	Recruitment planning + commissioning structure for external elements	Public Health/ Communications lead/Programme lead		
	Brand development	Public Health/ Communications lead/Programme lead		
	Planning for launch	Public Health/ Communications lead/Programme lead		

Phase 3 – Mobilisation and service design for neighbourhood 2

Date	Audience	Engagement	Staff / resourcing required	Comments	Achieved by
Jan-Dec 2027	Clusters	- Cluster meetings - Clinical workshops - Operational planning - Referral pathway design - Staff communications	Programme lead/ Primary care lead	Purpose of engagement: - Build ownership of the model - Refine service model - Agree implementation - Ensure integration - Prepare workforce	December 2027
	Primary care managers		Programme lead/ Primary care lead		
	Practice managers		Programme lead/ Primary care lead		
	Health Coaches		Programme lead/ Primary care lead		
	Community teams		Programme lead/ Primary care lead		
	People living with and beyond cancer		Programme lead/ Primary care lead		
	Carers		Programme lead/ Primary care lead		
	Underserved groups		Programme lead/ Primary care lead		
	Volunteers		Programme lead/ Primary care lead		

Phase 3 – Mobilisation and service design for neighbourhood 2

Date	Audience	Engagement	Staff / resourcing required	Comments	Achieved by
Community Health & Wellbeing Workers					
Jan-Dec 2027	Communities	- Community workshops - Local listening events - Asset mapping - Co-design sessions - Community reference groups - Partnership meetings	Public Health/Programme lead/Community partnerships lead/CVC representatives/Local Authority partners	Purpose: Engagement to support operational design plus community codesign	December 2027
	Community leaders				
	Third sector				
	Local authorities				
	Community connector teams				
	Housing associations				
	Social prescribing teams				
	People living with and beyond cancer				
	Carers				
	Underserved groups				
	Volunteers				

Local neighbourhood infrastructure					
Date	Audience	Engagement	Staff / resourcing required	Comments	Achieved by
Jan-Dec 2027	CVCs	- Partnership forums - Coproduction workshops - community conversations - mapping existing provision - grant design workshops - governance arrangements - local planning sessions	Public Health/Programme lead/Community partnerships lead/CVC representatives/Local Authority partners	Purpose: Engagement to support broader strategic and community engagement to shape investment, governance and delivery Key: Mapping of existing assets and identifying gaps to underpin investment decisions	December 2027
	Third sector organisations – including social enterprises, charities				
	Town and Community Councils				
	Community groups				
	Volunteers				
	Local Authorities				
	Anchor organisations				
	Regional Partnership Board				
	People living with and beyond cancer				
	Carers				
	Underserved groups				

Phase 4 – Mobilisation and service design for neighbourhood 3

Date	Event / Activity	Staff / resourcing required	Comments	Achieved by
Jan-Dec 2028	Partner workshops	Public Health/Programme lead	Focus of engagement: • Confirm delivery sites/neighbourhoods • Agree governance arrangements • Develop implementation timetable • Establish local implementation groups	December 2028
	Clinical engagement with GP surgeries	Public Health/Programme lead/Primary Care lead		
	Community sector engagement	Programme lead/CVC representatives/Community partnerships lead		
	Local authority engagement	Public Health/Programme lead/Local Authority partners		
	Develop implementation plan and timeline	Public Health/Programme lead		
	Communications toolkit	Public Health/ Communications lead/Programme lead		
	Recruitment planning + commissioning structure for external elements	Public Health/ Communications lead/Programme lead		
	Brand development	Public Health/ Communications lead/Programme lead		
	Planning for launch	Public Health/ Communications lead/Programme lead		

Phase 4 – Mobilisation and service design for neighbourhood 3

Date	Audience	Engagement	Staff / resourcing required	Comments	Achieved by
Jan-Dec 2028	Clusters	- Cluster meetings - Clinical workshops - Operational planning - Referral pathway design - Staff communications	Programme lead/ Primary care lead	Purpose of engagement: - Build ownership of the model - Refine service model - Agree implementation - Ensure integration - Prepare workforce	December 2028
	Primary care managers		Programme lead/ Primary care lead		
	Practice managers		Programme lead/ Primary care lead		
	Health Coaches		Programme lead/ Primary care lead		
	Community teams		Programme lead/ Primary care lead		
	People living with and beyond cancer		Programme lead/ Primary care lead		
	Carers		Programme lead/ Primary care lead		
	Underserved groups		Programme lead/ Primary care lead		
	Volunteers		Programme lead/ Primary care lead		

Phase 4 – Mobilisation and service design for neighbourhood 3

Date	Audience	Engagement	Staff / resourcing required	Comments	Achieved by
Community Health & Wellbeing Workers					
Jan-Dec 2028	Communities	- Community workshops - Local listening events - Asset mapping - Co-design sessions - Community reference groups - Partnership meetings	Public Health/Programme lead/Community partnerships lead/CVC representatives/Local Authority partners	Purpose: Engagement to support operational design plus community codesign	December 2028
	Community leaders				
	Third sector				
	Local authorities				
	Community connector teams				
	Housing associations				
	Social prescribing teams				
	People living with and beyond cancer				
	Carers				
	Underserved groups				
	Volunteers				

Local neighbourhood infrastructure					
Date	Audience	Engagement	Staff / resourcing required	Comments	Achieved by
Jan-Dec 2028	CVCs	- Partnership forums - Coproduction workshops - community conversations - mapping existing provision - grant design workshops - governance arrangements - local planning sessions	Public Health/Programme lead/Community partnerships lead/CVC representatives/Local Authority partners	Purpose: Engagement to support broader strategic and community engagement to shape investment, governance and delivery Key: Mapping of existing assets and identifying gaps to underpin investment decisions	December 2028
	Third sector organisations – including social enterprises, charities				
	Town and Community Councils				
	Community groups				
	Volunteers				
	Local Authorities				
	Anchor organisations				
	Regional Partnership Board				
	People living with and beyond cancer				
	Carers				
	Underserved groups				

Phase 5 – Public launch & communications**October 2026 – September 2031****Key: Neighbourhood 1 (N1), Neighbourhood 2 (N2), Neighbourhood 3 (N3)**

Date	Event / Activity	Staff / resourcing required	Comments	Achieved by
Jan 2027 – N1 Jan 2028 – N2 Jan 2029 – N3	Communications preparation and stakeholder engagement	Public Health/Programme lead/Communication lead/Partnership development lead	Key activities to include: • develop partner communications toolkit • stakeholder briefing pack • elected member briefing • community information materials • website resources (key messages, FAQs, downloadable materials)	Jan 2027 – N1 Jan 2028 – N2 Jan 2029 – N3
	Launch and awareness raising		Key activities to include: • launch event with partners • press releases • local media campaign • social media launch activity • videos – (launch/promotional content)	
	Ongoing engagement		Key activities to include: • website (ongoing programme information and updates) • social media (ongoing updates engagement and amplification) • case studies and staff stories • partner communications toolkit (ongoing use and updates)	

Evidence reviews and literature synthesis

Scoping searches on health coaching and community health and wellbeing workers (CHWWs)

Two scoping searches were undertaken by a librarian in May 2026 to support preliminary work for this proposal.

HEALTH COACHING

A search of the internet identified the NHS England *Workforce Development Framework for Health and Wellbeing Coaches*, which contains potentially relevant references. Additional searches were conducted in Epistemonikos (a database of systematic reviews), the Health Management Information Consortium (HMIC), and MEDLINE. In total, one hundred and eight references were identified.

NHS England (2023) *Workforce development framework for health and wellbeing coaches*. London: NHS England. Available at: <https://www.england.nhs.uk/long-read/workforce-development-framework-for-health-and-wellbeing-coaches/> (Accessed: May 2026).

COMMUNITY HEALTH AND WELLBEING WORKERS (CHWWs)

A search of the internet exploring links between CHWWs and deprivation retrieved several relevant papers. This was followed by a search of the Health Management Information Consortium (HMIC), which identified a further three papers. Further searches were undertaken in the MEDLINE and Embase databases, identifying further studies; however, the majority were conducted outside the United Kingdom. MEDLINE uses a broad definition of community health workers, whereas Embase lacks a specific subject heading; therefore, the following phrase search was employed to help define CHWWs:

A total of two hundred and seventy-one references were identified.

[illegible]

THIS AGREEMENT is made on

BETWEEN:

- (1) **MACMILLAN CANCER SUPPORT**, a charity registered in England and Wales (registration number 261017), Scotland (registration number SC039907) and the Isle of Man (registration number 604), and a company limited by guarantee with registered number 02400969 whose registered office is 3rd Floor, Bronze Building, The Forge, 105 Sumner Street, London, SE1 9HZ (the **Investor**); and
- (2) **[HEALTH AND SOCIAL CARE TRUSTS, HSCT]** a statutory NHS organisation whose registered office is NHS [] (the **Commissioner**);

(the **parties**, each a **party**).

WHEREAS

- (A) The Investor has a long history of providing transformational support for health and social care. It now wishes to undertake social investments to assist and support organisations that are seeking to shift care closer to home and to work with partners to develop a social investment programme that will act as a tool for sustainable transformation in neighbourhoods.
- (B) The Commissioner is the statutory NHS body responsible for the provision of primary, community, mental health and acute hospital services to meet the needs of the # million people living in [geography].
- (C) The Investor and the Commissioner wish to work together and enter into a social impact outcome funding agreement ("Agreement") in relation to the provision of certain neighbourhood transformation services in accordance with the terms of this Agreement.

IT IS AGREED as follows:

1. Definitions and Interpretation

Conditions Precedent	means the conditions listed in clause 5.1;
Data Protection Legislation	means the Data Protection Act 2018, the UK General Data Protection Regulation, and all applicable laws and regulations relating to the processing of personal data and privacy and any regulations or secondary legislation made thereunder, in each case as amended or replaced from time to time;
Funded Programme Period	means the Funding Start Date to the Funding End Date;
Funding	means the funding of up to £10,000,000 to cover the Programme Costs over the Funded Programme Period that will be provided by the Investor to the Commissioner in accordance with this Agreement;
Funding End Date	Means 31st December 2028;

Funding Instalment	shall be as defined in clause 4;
Funding Period	means each funding period set out in clause 4.1;
Funding Start Date	means 1 st January 2026;
Improvement Plan	shall be as defined in clause 10;
Key Performance Indicators	means the schedule attached at Appendix 2 that sets out the Key Performance Indicators and quality markers required for the Service to be evaluated, including frequency and format in which such information is required;
Operational Group	means the working group that will be established for the Programme comprising representatives of the Investor, the Commissioner and the Service Providers;
Outcomes	means the outcomes achieved from the Service;
Outcomes Methodology	means the methodology for calculating Outcomes Payments set out in Appendix 1;
Outcomes Payment	means any outcomes payment that may be payable by the Commissioner to the Investor as calculated in accordance with the Outcomes Methodology and as defined in Appendix 1;
Outcomes Period	means each of the periods as detailed in clause 7.7, the last of which ends on the date which is twelve (12) months after the end of the Funded Programme Period;
Programme	means the programme to promote and deliver the Service over the Funded Programme Period;
Programme Board	means the board that will be established for the Programme comprising accountable representatives from the Commissioner, the Investor and the Service Providers;
Programme Budget	means the detailed budget for each year of the Programme set out in Appendix 3 that totals £10,000,000 and includes all the anticipated costs required to promote and deliver the Service in accordance with the Service Specification and this Agreement;
Programme Costs	means the actual costs required to promote and deliver the Service in accordance with the Service Specification and this Agreement;
Programme Milestones	means the milestones for the Programme set out in Appendix 4;

Qualifying Service Patient	Shall be defined in Appendix 1:
Service	means the proposed service known as the [] that aims to [reduce the number of Unplanned Bed Days] for the Target Cohort by implementing a new integrated model for [...] in [geography].
Service Agreement	means the contract variation letter and the relevant standard NHS contract, to be signed, prior to service commencement, between the Commissioner and the Service Provider whereby the Service Provider agrees to provide the Service;
Service Patient	means each person in the Target Cohort who uses the Service during the Funded Programme Period;
Service Provider	means [...] whose registered office is [...];
Service Specification	means the specification for the Service attached at Appendix 5;
Shared Aims	means the matters set out in clause 2;
Target Cohort	means those aged 18 and [...] in the geographical area covered by the Commissioner;
[Unplanned Hospital Bed Day]	means as defined in Appendix 6].

1.1. In this Agreement, unless otherwise specified:

- 1.1.1. references to clauses, subclauses, paragraphs and subparagraphs are to clauses, subclauses, paragraphs and subparagraphs of this Agreement;
- 1.1.2. a reference to any statute or statutory provision shall be construed as a reference to the same as it may have been, or may from time to time be, amended, modified or reenacted;
- 1.1.3. any reference to a day (including within the phrase working day) shall mean a period of 24 hours running from midnight to midnight;
- 1.1.4. a reference to any other document referred to in this Agreement is a reference to that other document as amended, restated, varied, novated or supplemented from time to time; and
- 1.1.5. the expressions “including”, “includes” and “include” have the meaning as if followed by “without limitation”.

1.2. All headings and titles are inserted for convenience only. They are to be ignored in the interpretation of this Agreement.

2. Shared Aims

- 2.1. The purpose of this Agreement is to provide a common basis for all parties to implement the Programme in a manner that best ensures the intended outcomes

are achieved, as well as to provide for the management and coordination of any remedies or disputes that may arise.

2.2. The parties agree that their aims for the Programme include:

- 2.2.1. demonstrating the system benefit delivered by the Programme such that the NHS or another appropriate body takes on the funding of the Programme to enable it to continue beyond the end of the Funded Programme Period; and
- 2.2.2. ensuring that everyone in the Target Cohort has the opportunity to benefit from the Service and that no-one within the Target Cohort will be discriminated against when using the Service, including on grounds of gender, race, disability, age, sexual orientation, religion, belief, gender reassignment, pregnancy, maternity or marital or civil partnership status.

2.3. The parties agree that the successful implementation of the Programme will depend on the ability of the Investor and Commissioner effectively to co-ordinate and combine their expertise, manpower and resources in order to deliver an integrated approach to service provision. To this end, the parties agree to develop a close working relationship between each other at all appropriate levels, based upon openness and trust in a transparent information and data sharing environment. The Commissioner shall ensure that the Service Provider is compliant with this provision.

3. Commissioner Undertakings

3.1. Recognising the interdependencies between this Agreement, the Service Specification, Service Agreement, Programme Budget and the Shared Aims, the Commissioner warrants and undertakes throughout the term of this Agreement:

- 3.1.1. that the Service Specification and Service Agreement is fit for purpose and consistent with the Commissioner's requirements for delivery according to the agreed business case for the Service;
- 3.1.2. that the Service Specification and Service Agreement is fully consistent and does not conflict with the terms of this Agreement;
- 3.1.3. that the Programme Budget contains all the anticipated costs required to promote and deliver the Service in accordance with the Service Specification and this Agreement;
- 3.1.4. that the Commissioner, Service Provider and Investor have the appropriate information governance and sharing agreements in place to enable the Investor and the Commissioner to monitor Outcomes and the Key Performance Indicators in accordance with this Agreement;
- 3.1.5. that the Funding will be used solely to pay for Programme Costs in accordance with the Programme Budget over the Funded Programme Period and not for any other purpose;
- 3.1.6. not to amend the Service Agreement, Service Specification, Service Agreement or Programme Budget without the Investor's prior written approval, and to make any request to make any such amendment to the Investor in reasonable time (and at least 30 days) before the proposed amendment would come into effect; and

- 3.1.7. that it has the power to enter into, deliver and perform this Agreement and the transactions contemplated by it, and it has obtained all authorisations necessary to enable it to enter into and comply with this Agreement.
- 3.2. The Commissioner further warrants and undertakes that it will, and will procure that the Service Provider will:
 - 3.2.1. promote proactively and monitor referrals into the Service, taking appropriate action to increase referrals as necessary;
 - 3.2.2. make a meaningful contribution to the pre-marketing and marketing of the Service to enable relevant stakeholders (e.g. GPs/Out of Hours services) to engage with the Programme;
 - 3.2.3. keep, and ensure that relevant parties keep, full, true and accurate records relating to the Programme, Outcomes and referrals and ensure that the Investor is given sufficient information from such records to enable the Investor to perform their roles and exercise their rights under this Agreement;
 - 3.2.4. comply with all applicable laws in relation to the Programme and to comply with anti-bribery, corruption and tax evasion laws and codes, including adopting and enforcing appropriate policies and procedures to prevent anti-bribery, corruption and tax evasion in accordance with relevant legislation and government guidance;
 - 3.2.5. ensure that the information regarding the Programme provided to the Investor is complete, true and accurate, and not misleading in any material respect.

4. Funding

- 4.1. Subject to the Conditions Precedent being satisfied and remaining satisfied on the first date of each Funding Period, and subject to the provisions of this clause 4 and clause 7.9, the Investor will provide the Funding to the Commissioner for the following periods in instalments (each a **Funding Instalment**). The actual amount of each Funding Instalment will be calculated in accordance with the rest of this clause 4, the following amounts are indicative only and amounts calculated in accordance with clause 4.2 shall prevail:

Funding Period (dates inclusive)	Indicative Amount
1 January 2026 – 31 March 2026	£833,337
1 April 2026 – 30 June 2026	£833,333
1 July 2026 – 30 September 2026	£833,333
1 October 2026 – 31 December 2026	£833,333
1 January 2027 – 31 March 2027	£833,333
1 April 2027 – 30 June 2027	£833,333
1 July 2027 – 30 September 2027	£833,333
1 October 2027 – 31 December 2027	£833,333
1 January 2028 – 31 March 2028	£833,333
1 April 2028 – 30 June 2028	£833,333
1 July 2028 – 30 September 2028	£833,333
1 October 2028 – 30 December 2028	£833,333

4.2. Within 30 days of the end of each Funding Period, the Commissioner will:

4.2.1. submit a statement of costs to the Investor to show a breakdown of the Programme Costs that it and / or the Service Provider has incurred in respect of such period (**relevant costs**); and

4.2.2. notify the Investor of the amount of the relevant Funding Instalment, being the amount of the relevant costs, all in a form satisfactory to the Investor.

4.3. Also, within 30 days of the end of each Funding Period, the Commissioner will send the Investor a valid invoice for the Funding Instalment. Subject to clause 5, the Investor will then pay such amount to the Commissioner within 30 days of receipt.

4.4. In the event that any statement of costs submitted under clause 4.2 indicates that not all the Funding is required or that more funding is required to deliver the Programme, the Investor and the Commissioner will work together to agree whether the amount of the Funding and future Funding Instalments should be amended accordingly.

4.5. No amendment to the amount of the Funding may be made without the Investor's prior written approval.

4.6. In the event that the Commissioner wishes the Service Provider to continue to provide the Service after the Funding End Date, the parties will work collaboratively to consider options for further funding. The aim would be for such options to be considered in principle by the date which is six (6) months prior to the Funding End Date. For the avoidance of doubt, nothing in this Agreement obliges the Investor to provide further funding or payments in addition to the Funding, including for any overspend or additional costs incurred by the Commissioner or to make up any shortfall in funding from elsewhere.

5. Conditions Precedent

- 5.1. The Investor shall not be obliged to provide any Funding Instalment to the Commissioner unless:
 - 5.1.1. the Investor has agreed the Service Specification and the Service Agreement;
 - 5.1.2. The Investor has received appropriate confirmation from the Service Provider that they are aware of the Investor's role in relation to the Programme and they will give due consideration to the Investor's views;
 - 5.1.3. any Improvement Plan has been implemented and adhered to by the Service Provider within 28 days of being agreed;
 - 5.1.4. the applicable Programme Milestones have been met or, failing which, the parties have agreed in writing that achievement was near enough and/or to amend one or more of such milestones, in which case any withheld Funding Instalment will be paid once the applicable Programme Milestone(s) are achieved;
 - 5.1.5. there is no unremedied breach of any of the Commissioner's warranties, undertakings or other obligations under this Agreement;
 - 5.1.6. there is no unremedied breach of any of the Service Provider's warranties, undertakings or other obligations under the Service Agreement;
 - 5.1.7. there is currently outstanding no investigation or inadequate rating from the Care Quality Commission in respect of the Service;
 - 5.1.8. the Commissioner has provided the Investor with a breakdown of the total Programme Costs that it has incurred in relation to the Service in respect of the relevant Funding Period; and
 - 5.1.9. the Investor has received a valid invoice from the Commissioner for the Funding Instalment in accordance with this Agreement.
- 5.2. In the event that any or all of the Conditions Precedent are not met, the Investor may, in its absolute discretion, withhold or reduce any Funding Instalments that have not yet been provided to the Commissioner. The Investor will inform the Commissioner as soon as reasonably practicable if it is aware this step is being considered so the Commissioner has an opportunity to remedy the issue.

6. Outcomes Reporting

- 6.1. The Commissioner will, and will procure that the Service Provider will:
 - 6.1.1 monitor the Outcomes and Key Performance Indicators as set out in Appendix 2 on a monthly basis, including through the use of a dashboard that will be built and operated by the Commissioner;
 - 6.1.2 provide, and will ensure that the Service Provider provide, to the Commissioner the necessary data for the dashboard using the Reporting Schedule within two months of the month end unless otherwise specified in the Reporting Schedule. The Commissioner will collate and analyse such data and share reports on a monthly basis with relevant stakeholder groups including the Programme Board and Operational Group to assist in the performance management of the Programme;

- 6.1.3 ensure the data provided to the Commissioner is as accurate as possible, and will further ensure that any additional metrics or case studies as may be agreed are provided on a timely basis to evaluate impact and support service development and learning;
- 6.1.4 be responsible for reporting on Outcomes and the Key Performance Indicators.

6.2. Subject to clauses 14 (Publicity) and 16 (Data Sharing), the Investor will be entitled to disclose the information it receives in relation to Outcomes to an independent validator appointed by the Investor to ascertain whether (i) the relevant outcomes have been recorded accurately; and/or (ii) the quality of evidence is sufficient for reasonable certainty that the relevant outcome has been achieved.

7. Payment Dependent on Outcomes

7.1 Subject to clause 7.3 and clause 12 (term and termination), the Commissioner will make payments to the Investor solely on the basis of Outcomes. For the avoidance of doubt, the total amount of Outcomes Payments payable to the Investor will not exceed the total amount of Funding received by the Commissioner under this Agreement.

7.2 In respect of each Outcomes Period, based on the data provided under this Agreement, the Commissioner will calculate in accordance with the Outcomes Methodology the amount of any Outcomes Payment that may be due. Investor will be entitled to check the amount of any Outcomes Payment and request clarification or verification, if necessary, provided that if the Investor does not query such amount within 30 days of being notified by the Commissioner, the amount will be deemed to be agreed.

7.3 In respect of each of the first [six] Outcomes Periods, the total Outcomes Payments payable by the Commissioner to the Investor will be restricted to an amount equal to the total Funding provided by the Investor for the same period (See Appendix 1 Outcomes Methodology)

That is provided that any Outcomes Payment due in excess of this amount (the **Excess**) will only be payable to the Investor in the event that the sum of all the restricted Outcomes Payments payable to the Investor in respect of the first [six] Outcomes Periods is less than the sum of the total Funding for the same periods provided by the Investor under this Agreement (the shortfall being the **Initial Shortfall**).

In such event, the Investor will submit after the end of the first [six] Outcomes Periods an invoice from the Investor for an amount which is the lesser of the amount of the accumulated Excess and the amount of the Initial Shortfall and the Commissioner will pay such amount within 30 days of issue.

7.4 If, following payment to the Investor of part or all of the Excess under clause 7.3, the amount paid by the Commissioner to the Investor is still less than the total Funding provided by the Investor under this Agreement (the **Continuing Shortfall**), two further Outcomes Payments will be payable by the Commissioner in respect of the two final Outcomes Periods. Such payments will be restricted to the amount required to enable the Investor to recover such Continuing Shortfall.

Service Patients for whom outcomes are calculated, will only include those who use

the service during the Funded Programme Period who die during the two final Outcome Periods.

7.5 The Investor will be entitled to check the data provided under this Agreement and request clarification or verification from the Commissioner, if necessary, provided that if the Investor does not query such data within 30 days of receipt, it will be deemed to be agreed.

7.6 Following the Commissioner's calculation of an Outcomes Payment under clause 7.2, the Investor will submit an invoice for the amount payable by the Commissioner and the Commissioner will pay such amount within 30 days of issue.

7.7 The anticipated invoice and payment timetable is as follows:

Outcomes Period (inclusive)	Investor invoice	Commissioner payment
1 January 2026 – 30 June 2026	30 September 2026	31 October 2026
1 July 2026 – 31 December 2026	31 March 2027	30 April 2027
1 January 2027 – 30 June 2027	30 September 2027	31 October 2027
1 July 2027 – 31 December 2027	31 March 2028	30 April 2028
1 January 2028 – 30 June 2028	30 September 2028	31 October 2028
1 July 2028 – 31 December 2028	31 March 2029	30 April 2029
*1 January 2025 – 31 December 2028	31 March 2020	30 April 2029
**1 January 2028 – 30 June 2029	30 September 2029	31 October 2029
**1 July 2029 – 31 December 2029	31 March 2030	30 April 2030

*Applicable where there is Excess to repay Initial Shortfall (clause 7.3)

**Applicable where there is a Continuing Shortfall (clause 7.4)

7.8 In the event that any party considers that the Outcomes Methodology no longer remains appropriate and/or it proves not possible accurately to measure the outcomes required to calculate the Outcomes Payments, it will notify the other parties in writing. Such notification will include the reasons for its views and, if appropriate, propose an alternative methodology. Each of the other parties will respond to such notice within fourteen (14) days of receipt stating whether or not it agrees. If any party does not agree, the parties will promptly enter into good faith discussions with a view to agreeing an amended Outcomes Methodology or proposing and agreeing an alternative suitable proxy for reduced healthcare expenditure based on changes in activity.

7.9 If no amendment to the Outcome Methodology or alternative suitable proxy can be agreed within thirty (30) days of one party giving notice to the others under clause 7.8 such that this Agreement can no longer be based on Outcomes, the repayment mechanism will default to a fixed repayment to the Investor and the Investor will not be required to provide any further Funding to the Commissioner. Such repayment will total an amount equal to the total Funding provided to the Commissioner by the Investor from the Funding Start Date to the date of repayment less the sum of all Outcomes Payments received by the Investor up until the date of repayment. The Commissioner will make

such payment in equal quarterly instalments over the period remaining until the Funding End Date.

7.10 No amendment to the Outcomes Methodology may be made without the Investor's nor the Commissioner's prior written approval.

8 Arrangements beyond the final Outcomes Period

8.1 In the event that, at the end of the final Outcomes Period and after the Investor has received any amounts from the Commissioner that may be due under clauses 7.3 and 7.4 to recover any Initial Shortfall and Continuing Shortfall, the total amount of Outcomes Payments paid to the Investor for all the Outcomes Periods is still less than the total Funding provided by the Investor to the Commissioner (the shortfall being the **Final Shortfall**), and if the Service is continued beyond the end of the Funded Programme Period, this Agreement will be automatically varied to enable the Commissioner to make further payments to the Investor until either the Investor has recovered the Final Shortfall or the date which is two (2) years after the end of the Funded Programme Period (whichever is the sooner). Such payments will be calculated in accordance with the Outcomes Methodology and at a rate to be reasonably approved by the Investor provided that, before making any such payments, the Commissioner will be entitled to deduct the costs paid to the Service Provider for providing the Service during such extension period (to the extent that evidence of which is provided to the Investor). Service Patients for whom outcomes are calculated, are those who use the service and die in the Final Shortfall Outcomes Period.

Final Shortfall Outcomes Period (inclusive)	Investor invoice	Commissioner payment
1 January 2029 – 31 December 2029	31 March 2029	30 April 2029
1 January 2030 – 31 December 2030	31 March 2031	30 April 2031

9. Programme Governance

9.1 A structure of Programme Board and Operational Group has been agreed to ensure sufficient oversight and support for the Programme from the Investor, the Commissioner and the Service Provider. The Terms of Reference (ToR) for each will be agreed in accordance with the Programme Milestones.

9.2 The Programme Board will provide strategic oversight and have ultimate accountability for the Programme and be a point of escalation and support for the Operational Group. It will meet bi-monthly for the first 6 months and then quarterly to discuss the Programme. Such discussions will include:

- 9.2.1 the performance of the Programme;
- 9.2.2 approving and reviewing annual development plans and budgets for the Programme prepared by the Operational Group;
- 9.2.3 opportunities to improve collaboration between the parties and the Service Provider;
- 9.2.4 opportunities to improve promotion and delivery of the Service and achieve better outcomes for Service users;

- 9.2.5 the quality and nature of information provided by the Commissioner;
 - 9.2.6 assistance with resolving any disputes between the Commissioner and the Investor;
 - 9.2.7 learning, to enable knowledge sharing across the system and wider end of life care community.
- 9.3 The Operational Group will oversee the operational management of the Programme and report to the Programme Board. It will meet monthly (unless otherwise agreed), make recommendations on operational issues, take forward any recommendations or actions on operational issues required by the Programme Board and report on progress against actions to the Programme Board, highlighting risks, issues and mitigations as required.

10 Improvement Plan

- 10.1 In the event of failure by the Service Provider to meet one or more of the Key Performance Indicators or Programme Milestones, or if the Investor considers for any other reason that the performance of the Service Provider in providing the Service or the quality of information reported by the Service Provider is inadequate, the Commissioner will procure that the Service Provider will work with the Operational Group to agree a plan to improve performance and/or the quality of information (the **Improvement Plan**). The Commissioner will procure that the Service Provider implements and adheres to the Improvement Plan within 28 days of such agreement.
- 10.2 The Improvement Plan will be approved, reviewed and amended, if necessary, at subsequent meetings of the Programme Board.
- 10.3 The Investor and the Commissioner will, and the Commissioner will procure that the Service Provider will, provide at least 7 days before any meeting all information that it reasonably believes to be relevant and any other information reasonably requested by another party to inform the process and discussion.
- 10.4 The Commissioner will, and will procure that the Service Provider will, attend and co-operate with any and all Programme Board, Operational Group and other meetings held pursuant to this Agreement.
- 10.5 For the avoidance of doubt, the processes set out in clauses 9 and 10 shall be separate and in addition to any performance review processes set out in the Service Specification and Service Agreement, and are without prejudice to any other rights that the Investor may have in relation to this Agreement.

11 General Assistance

- 11.1 The Investor and the Commissioner each undertake to work with each other, and the Commissioner undertakes to procure that the Service Provider works with the Investor and the Commissioner, to ensure that the Service optimises positive impact on patient care and value to the health and care system, including, but not limited to, looking for the Service to develop over time to respond to:
- 11.1.1 best clinical and operational practice;

- 11.1.2 learning from operations and impact over time;
 - 11.1.3 changes in other parts of the local health and care system and enhanced scope for integration; and
 - 11.1.4 challenges in the delivery of the Service and anticipated impact or savings.
- 11.2 The Investor and the Commissioner will, and the Commissioner will procure that the Service Provider will, immediately disclose to each other any matter or information that may have a material impact on the Programme.

12 Term and Termination

- 12.1 Unless terminated earlier or extended in accordance with its terms, and without prejudice to clause 7, this Agreement will terminate six (6) months after the end of the final Outcomes Period.
- 12.2 The Investor or the Commissioner may terminate this Agreement on giving not less than six (6) months' written notice to the other parties provided that if mutually agreed between the parties the notice period may be less than six (6) months.
- 12.3 This Agreement will automatically terminate in the event that the Commissioner ceases to commission the Service from the Service Provider. In the event that the Commissioner ceases to commission a substantial part (but not all) of the Service from the Service Provider, it will immediately notify the Investor and the Investor will have the right to terminate this Agreement with immediate effect by giving written notice to the other parties.
- 12.4 The Investor may terminate this Agreement with immediate effect by giving written notice to the other parties if the Commissioner or Service Provider:
- 12.4.1 commits any act which results in the Investor being brought into disrepute, or breaching any of its charitable objects or social investment duties;
 - 12.4.2 transfers to another person, suspends (or threatens to suspend) or ceases (or threatens to cease) to carry on, all or a substantial part of its operations; or
 - 12.4.3 takes any action or has any action taken in relation to it in connection with its entering administration, insolvency processes, provisional liquidation or any composition or arrangement with its creditors, or has a receiver appointed to any of its assets.
- 12.5 Termination of this Agreement shall be without prejudice to any accrued rights or obligations of the parties. Upon termination of this Agreement for any reason, the Commissioner will immediately pay any Outcomes Payments or other amounts that may be outstanding, including any Outcomes Payments and other amounts that may have accrued but remain unpaid as at the date of termination.
- 12.6 The parties' rights and obligations with regard to information sharing and confidentiality shall survive any termination of this Agreement for a period of five (5) years.

13 Disputes

- 13.1 The parties will each use reasonable endeavours to resolve any dispute that arises under this Agreement through a prompt, good faith discussion at a managerial level appropriate to the dispute in question.
- 13.2 In the event that a dispute is not resolved within five (5) days of it having been referred to a managerial level for discussion then any party may refer the dispute for resolution to an executive lead of each party with sufficient authority to make a decision on its behalf. Such executive leads will meet for discussion within ten (10) days or at a time that the parties agree as soon after as possible.
- 13.3 If the dispute is not resolved within ten (10) days of escalation to executive leads, the parties will apply for mediation at the Centre for Effective Dispute Resolution (CEDR) and will use their reasonable endeavours to conclude any such mediation as soon as possible.

14 Publicity and learning

- 14.1 The parties will agree together all publicity and press announcements in relation to the Programme and this Agreement and will not make any such publicity or press announcements without the other parties' agreement. Such announcements shall include any naming/badging of the Service and/or publicising the names of any professionals working in the Service.
- 14.2 The parties acknowledge that an important aim of the Programme and this Agreement is to share learning and will seek to be open and share learning by mutual consent where appropriate. The Investor will be entitled to use learning from the Programme and this Agreement in relation to other end of life care or health related services.

15 Contract variations

- 15.1 Any party may request in writing to the other parties an amendment to this Agreement. The other parties shall be required to respond to such request within 30 days of receipt by setting out their comments on the requested amendment including on cost, feasibility, other implications and why, if applicable, it does not accept such amendment request. No party is obliged to agree any amendment to this Agreement.
- 15.2 Amendments to this Agreement will only be effective if in writing and signed by the authorised representatives of the Investor and Commissioner.

16 Data sharing

- 16.1 The parties will comply with all applicable requirements of the Data Protection Legislation. This clause 16 is in addition to, and does not relieve, remove or replace, a party's obligations under the Data Protection Legislation.
- 16.2 The parties acknowledge that for the purposes of the Data Protection Legislation, the Investor is not the processor or controller of any personal data processed in the course of the Programme, unless otherwise agreed in writing.

- 16.3 If the Investor and the Commissioner agree there is a need to share personal data with the Investor, the relevant parties will enter into a separate data sharing agreement.

17 Freedom of Information

- 17.1 The parties acknowledge that the Commissioner may be subject to the requirements of the Freedom of Information Act 2000 ("FOIA") and the Environmental Information Regulations 2004. The Investor will assist and co-operate with the Commissioner to enable the Commissioner to comply with any relevant information request.
- 17.2 The Commissioner will be responsible for determining at its absolute discretion whether commercially sensitive information and/or any other information:
- 17.2.1 is exempt from disclosure in accordance with the provisions of the FOIA or the Environmental Information Regulations 2004;
 - 17.2.2 should be disclosed in response to an information request; and
 - 17.2.3 in no event shall the Investor respond directly to an information request unless expressly authorised to do so by the Commissioner.
- 17.3 The parties acknowledge that the Commissioner may, acting in accordance with regulations under FOIA or the Environmental Information Regulations, disclose information:
- 17.3.1 without consulting with the other parties; or
 - 17.3.2 following consultation with the other parties having taken their views into account.
- 17.4 The Investor will ensure that all information produced by it in the course of this collaboration is retained for disclosure for a period of six years after this Agreement is terminated. The Investor will permit the Commissioner to inspect on reasonable notice such relevant records as it may reasonably request from time to time.
- 17.5 The Investor acknowledges that any lists or schedules provided by it defining confidential information are of indicative value only. The Commissioner may nevertheless be obliged to disclose confidential information to meet statutory, legal, or parliamentary obligations.

18 Outcomes Records

- 18.1 The parties are committed to gaining a deep understanding of impact at an individual patient level to inform both service development and wider learning.
- 18.2 In accordance with the Programme Milestones, the Commissioner will procure that the Service Provider will undertake clinical reviews of random selections of patients who have been accepted into and supported by the Service and share these findings with the Programme Board and Operational Group.

19 Force Majeure

- 19.1 If any party acting in good faith, reasonably determines that, as a consequence of an act of God, governmental act, war, Covid-19 or other pandemic, epidemic, strike, national or local lockdown, the Service cannot operate as anticipated and/or the Investor cannot provide the Funding, it will notify the other parties whereupon the parties will consider in good faith other means of enabling the Programme or Funding to continue and the Investor will endeavour to continue to provide the support, however they will not be obliged to do so. In such event, any party will be entitled to request a variation to this Agreement and the terms of clause 15 (contract variations) shall apply.

20 Confidentiality

- 20.1 The terms of this Agreement are confidential and, except as required by law or as permitted in this Agreement, no party shall be entitled to disclose such terms to any other person except their professional advisers without the express prior written consent of the other parties.

21 Liability

- 21.1 As between the parties, the parties agree that the Commissioner is solely responsible for the Programme. Subject to clause 21.3, the Investor shall not be required to pay more than the total Funding, and the Investor does not accept liability for any consequences, whether direct or indirect, that may arise in connection with the Programme or otherwise from the use or withdrawal of any Funding or for any liability arising out of the acts or omissions of any person involved in the Programme (including without limitation the Commissioner and the Service Provider). The Commissioner shall ensure that the Service provider has appropriate insurance and indemnity arrangements in place and shall indemnify the Investor against any liability incurred by the Investor as a result of any claim against the Investor by the Service Provider.
- 21.2 Subject to clause 21.3, no party will be liable to another party for any loss of profits (other than, for the avoidance of doubt, the payment of Outcome Payments), loss of use, loss of production, increased operating costs, loss of business, loss of business opportunity, or goodwill or any other consequential or indirect loss of any nature, in connection with this agreement whether arising in tort or on any other basis
- 21.3 Nothing in this Agreement shall exclude or restrict the liability of any party:
- 21.3.1 for death or personal injury resulting from its negligence;
 - 21.3.2 for fraud or fraudulent misrepresentation; or
 - 21.3.3 in any other circumstances where liability may not be limited or excluded under any applicable law.

22 No Waiver

- 22.1 Any failure to exercise or delay by a party in exercising a right or remedy arising in connection with this Agreement or by law shall not constitute a waiver of such right or remedy or of any other rights or remedies. No waiver shall be effective unless

in writing and signed by the relevant party or on its behalf by a duly authorised representative. A waiver of a right or remedy on one occasion shall not constitute a waiver of the same right or remedy in the future.

23 No Partnership

23.1 Nothing in this Agreement shall be:

23.1.1 deemed to constitute a partnership, joint venture, representative or agency relationship between the parties; (or between the parties and the Service Provider); or

23.1.2 construed or have effect as constituting any relationship of employer and employee between the parties (or between the parties and the Service Provider).

23.2 No party shall have the authority to bind or pledge the credit of, or oblige another party in any way without obtaining such other party's prior written consent.

24 Notices

24.1 The contact details for the parties are:

The Investor: Macmillan Cancer Support
3rd Floor, Bronze Building, The Forge, 105 Sumner
Street, London, SE1 9HZ

For the attention of: Steven McIntosh, Chief
Partnership Officer
Tel: 07875724088
Email: smcintosh@macmillan.org.uk

For the attention of: Head of Legal
Email: legalenquiries@macmillan.org.uk

The
Commissioner: [...]
[...]

For the attention of: [...]
Tel:[...]
Email: [...]

24.2 All notifications and other communications under this Agreement will be in writing (which includes email).

24.3 Any party may change its contact details by notice to the other parties. Such change of contact details shall take effect five (5) days after they have been notified to the other parties.

24.4 Any notice or communication shall be deemed to have been received:

- 24.4.1 if delivered by hand, on signature of a delivery receipt or at the time the notice or communication is left at the proper address;
- 24.4.2 if sent by pre-paid first-class post or other next day delivery service, at 9.00 am on the second business day after posting or at the time recorded by the delivery service if earlier;
- 24.4.3 if sent by email, on the day of transmission if sent before 4.00 pm on the day of transmission (if a business day) or if not at 9.00 am on the next business day after transmission.

25 Assignment

- 25.1 No party may assign, novate or otherwise transfer its rights and liabilities under this Agreement without the consent of the other parties, not to be unreasonably withheld.

26 Miscellaneous

- 26.1 No one other than a party to this Agreement, their successors and permitted assignees, shall have any right to enforce any of its terms.
- 26.2 This Agreement constitutes the entire agreement between the parties in respect of the Programme and the Service and supersedes and extinguishes all previous agreements, promises, assurances, warranties, representations and understandings between them, whether written or oral, relating to its subject matter.
- 26.3 This Agreement may be executed in any number of counterparts, each of which when executed and delivered shall constitute a duplicate original, but all the counterparts shall together constitute the one agreement.
- 26.4 This Agreement and any dispute or claim (including non-contractual disputes or claims) arising out of or in connection with it or its subject matter or formation shall be governed by and construed in accordance with the law of England and Wales.
- 26.5 Each party irrevocably agrees that the courts of England and Wales shall have exclusive jurisdiction to settle any dispute or claim (including non-contractual disputes or claims) arising out of or in connection with this Agreement or its subject matter or formation.

This Agreement has been entered into by the following duly authorised representatives of the parties on the date stated at the beginning of it.

Signed by
for and on behalf of

MACMILLAN CANCER SUPPORT

.....

Signed by
for and on behalf of

NHS [...]

.....

APPENDIX 1 - Outcomes Methodology

APPENDIX 2 - Key Performance Indicators

APPENDIX 3 - Programme Budget

APPENDIX 4 - Programme Milestones

APPENDIX 5 - Service Specification

APPENDIX 6 - Baseline Methodology

Part 1 - Quality Impact Assessment Toolkit

Overview & Guidance

This assessment should be completed to quantify potential impacts on quality or safety aspects (positive and negative), from any strategic decisions e.g. policy decisions, business cases, service improvements and changes, or efficiency savings projects that will affect operational services.

When completing the assessment consider the impact that the change will bring about in the long term. Also consider any impacts that might occur whilst the change is being implemented. For example, the project may be to introduce a new clinical pathway into an existing team, this will reduce waiting times for patients and result in smaller caseloads which are both long-term positive impacts. However, to introduce the new pathway staff working arrangement will need to change which may increase staff turnover resulting in patient waiting times for treatment increasing both are short term negative impacts. The QIA should reflect both the short-term and long-term impacts.

For the approval process and to ensure you are using the current version of the tool, please visit the QIA SharePoint page
https://nhs.wales365.sharepoint.com/sites/HDD_Nursing-assurance-and-safety/SitePages/Quality-Impact-Assessment.aspx

The tool once approved by the Clinical Care Group Trumvirate e.g. Service Director, Assistant Director of Quality, Safety and Patient Experience and Associate Medical Director, should be submitted to patient.safety@wales.nhs.uk for consideration at the QIA Panel. The dates for the panel and submission deadlines are published on the QIA SharePoint page.

Strategic Decision / Organisational Activity / Project	
Title:	Macmillan Cancer Support Neighbourhood Investment Partnership
Name and role of lead:	Bruce Bolam, Bethan Lewis, Craig Jones
Clinical Care Group	Public Health Directorate

Description of Strategic Decision / Project:	
Broadly outline what is being proposed and the decision that needs to be made	This business case proposes a five-year Neighbourhood Prevention Programme, funded through Macmillan Cancer Support, to deliver neighbourhood-based prevention and wellbeing support through Health Coaching, Community Health and Wellbeing Workers, and investment in community capacity and local infrastructure. The programme aims to improve population health, reduce health inequalities, strengthen communities, and reduce avoidable demand on health and care services. The decision required is for approval of the business case and progression to formal funding, implementation, governance, and evaluation arrangements.
Why is the proposal / decision needed	The proposal is needed to improve prevention, early intervention and community-based support for people affected by cancer and other long-term conditions. It will help address health inequalities, improve population wellbeing, strengthen community resilience and support a shift from reactive healthcare to preventative, value-based Healthcare.
What are the drivers and influencing factors around the decision to be made? (e.g. legislation, national policy, professional body guidance, cost savings, ministerial priorities, quality standards, incidents etc)	The proposal is driven by increasing demand on NHS services, rising prevalence of cancer and other long-term conditions, and persistent health inequalities across the Hywel Dda population. It aligns with Welsh Government priorities for prevention and population health, Value-Based Healthcare principles, the HDdUHB A Healthier Mid and West Wales strategy, the Social Model for Health and Wellbeing, Community by Design, and the 20four7 Prevention Model. The programme is also supported by a significant investment opportunity from Macmillan Cancer Support and is intended to improve health outcomes, strengthen communities, and reduce avoidable healthcare utilisation through earlier intervention and prevention.
Who is directly affected by this proposal / decision? Please also consider people who may be indirectly affected	Directly affected groups include people living in the three neighbourhoods selected for the programme, particularly those affected by cancer, long-term conditions, lifestyle-related risk factors, and those living in areas of greater deprivation. Patients accessing Health Coaching services, Community Health and Wellbeing Worker support, and community-based prevention initiatives will be directly impacted by the proposed model. Indirectly affected groups include carers and families, primary care teams, community and voluntary sector organisations, local authority partners, cancer services, wider health and care staff, and the broader population who may benefit from improved community infrastructure, prevention services, and reduced pressure on NHS services.
How have you engaged with the people affected? If you have not yet engaged, what are your plans?	Engagement has commenced during the development of the business case and has included stakeholder mapping, executive briefings, steering group discussions, stakeholder workshops, and engagement with statutory, voluntary sector and community partners. The business case has also been reviewed and signed off by a patient representative to ensure the patient perspective has informed the proposal. A comprehensive Communications and Engagement Plan has been developed which sets out a continuous programme of engagement from July 2026 to September 2031. Subject to approval, this will include co-production with local communities, people living with and beyond cancer, carers, underserved groups, primary care teams, third-sector organisations and local authority partners through workshops, listening events, community reference groups, asset mapping and co-design sessions to shape implementation and ongoing service development.
How does the proposal / decision impact on delivery of the organisation's strategic objectives or ministerial priorities?	The proposal directly supports Hywel Dda University Health Board's strategic objective to deliver A Healthier Mid and West Wales by shifting the focus from reactive healthcare towards prevention, early intervention and community-based support. It aligns with the Health Board's Social Model for Health and Wellbeing, Community by Design approach, and 20four7 Prevention Model, with a strong focus on reducing health inequalities, improving population wellbeing and supporting value-based healthcare.
Is the proposal / decision planned to be temporary or permanent?	The proposal is a five-year programme running from October 2026 to September 2031, funded through Macmillan social investment and grant funding. Whilst the initial funding is time-limited, the programme is intended to demonstrate value, evaluate impact and develop sustainable commissioning arrangements beyond the funding period. This includes exploring pooled budgets and joint commissioning arrangements with key stakeholders to support the long-term sustainability of successful neighbourhood prevention services.

Has this Quality Impact Assessment been completed in collaboration with the clinical team(s) that the project will affect?	Yes / No
	Planned and Specialist Care Clinical Care Group; Community and Integrated Medicine Clinical Care Group; Primary Care Clinical Care Group

QUALITY IMPACT ASSESSMENT

Area	IMPACT
<i>Note - you can select more than one box per area if change may have multiple impacts e.g. both positive and negative</i>	<i>Description of the impact for each area (the impact mabe positive and/or negative)</i>
Safe How does this decision ensure that our health care system is a high quality, highly reliable and safe system that avoids preventable harm, maximising the things that go right and learning from when things go wrong to prevent them occurring again? How does this decision ensure that people’s health, safety and welfare are actively promoted and protected?	This proposal is expected to have a positive impact by improving prevention, enabling earlier identification of health and wellbeing needs, and providing proactive support to reduce avoidable harm and health inequalities. It will strengthen access to community-based services, improve self-management, and support people to stay healthier for longer. A robust monitoring, evaluation and quality improvement framework will support continuous learning and service improvement, helping to maximise positive outcomes and respond to emerging issues. The programme promotes and protects people's health, safety and welfare by improving access to preventative support, strengthening community resilience, and enhancing coordination between health, care and community services.
Timely How does this decision ensure that people have access to the high-quality advice, guidance and care they need quickly and easily, in the right place, first time? How does this decision ensure we care for those with the greatest health need first, and where treatment is identified as necessary, we treat people based on their identified and agreed clinical priority?	This proposal is expected to have a positive impact by improving access to preventative support and neighbourhood-based services, enabling people to receive advice, guidance and support earlier and closer to home. Health Coaching, Community Health and Wellbeing Workers, and strengthened community infrastructure will help people access the right support at the right time, improve navigation of services, and reduce barriers to care. The programme will also strengthen integration between health, care and community services, making it easier for people to access coordinated support tailored to their needs. This proposal is expected to have a positive impact by targeting support towards people and communities with the greatest health needs, particularly those experiencing deprivation and at higher risk of cancer and other long-term conditions. Prioritisation and triage processes will help ensure individuals receive support and access to services according to their level of need and clinical priority.
Effective How does this decision ensure care and treatment reflects evidence-based best practice? How does this decision ensure people receive the right care to achieve the optimal outcomes possible for them and that matter to them?	This proposal is expected to have a positive impact by implementing evidence-based approaches to prevention, behaviour change, community outreach and supported self-management. The programme has been designed using learning from existing Hywel Dda initiatives, established prevention models, published evidence, and best practice in neighbourhood health and value-based healthcare. It also includes a robust evaluation and quality improvement framework to continuously assess effectiveness and refine delivery based on emerging evidence and learning. This proposal is expected to have a positive impact by providing personalised, person-centred support through Health Coaching, Community Health and Wellbeing Workers, and community-based services. The programme focuses on what matters to individuals, helping them improve their health, wellbeing and self-management while connecting them to the most appropriate support to achieve outcomes that are important to them.

Opportunity / Consequence Rating*	*IIA Matrix		
	Consequence	Likelihood	Total Score
** positive rating			0
** negative rating			0
** positive rating			0
** negative rating			0
** positive rating			0
** negative rating			0

[illegible]

Efficient	How does this decision ensure we take a value-based approach to improve outcomes that matter most to people in a way that is as sustainable as possible and avoids waste?	This proposal is expected to have a positive impact by taking a value-based approach that focuses on prevention, wellbeing and outcomes that matter to people. By investing in early intervention, self-management and community-based support, the programme aims to improve health outcomes, reduce health inequalities and avoid unnecessary demand on healthcare services, supporting more sustainable use of resources and reducing waste.
	How does this decision ensure that there is the most effective use of resources to achieve best value in an efficient way?	This proposal is expected to have a positive impact by investing in prevention and early intervention to reduce avoidable demand on health services and improve long-term outcomes. The programme will use a combination of Health Coaching, Community Health and Wellbeing Workers and community-based support to make the best use of available resources, while a robust evaluation and benefits realisation framework will ensure value for money and inform future investment decisions.
Equitable	How does this decision ensure that everyone is provided with an equal opportunity to attain their full potential for a healthy life which does not vary in quality because of personal protected characteristics; the organisation that provides care; or location where care is delivered?	Has an Equality Impact Assessment been undertaken? Yes/No
		Has an Equity Health Impact Assessment been undertaken? Yes/No Please reflect a summary of the conclusion of the Equality Impact Assessment and the Equality Health Impact Assessment Guidance can be found on SharePoint
Person-centred	How does this decision ensure people's needs are met and that their preferences, needs and values guide decision-making that is made in partnership between individuals and the workforce?	This proposal is expected to have a positive impact by adopting a person-centred approach that focuses on what matters to individuals. Through personalised Health Coaching, community-based support and co-production, people will be supported to identify their own goals, improve self-management and access services that reflect their needs, preferences and circumstances. The programme also promotes partnership working between individuals, communities and professionals to support shared decision-making and improved outcomes.
	How does this decision ensure people and their families are at the centre of decisions, seeing them as experts working alongside professionals to get the best outcome and experience?	This proposal is expected to have a positive impact by adopting a co-produced, person-centred approach that recognises individuals and families as partners in their care. Through personalised Health Coaching, Community Health and Wellbeing Workers and community-led design, people will be supported to identify what matters most to them and make informed decisions about their health and wellbeing in partnership with professionals.

** positive rating			0
** negative rating			0
** positive rating			0
** negative rating			0
** positive rating			0
** negative rating			0

Metric - Current Baseline (date) - Trigger for escalation -	
Metric - Current Baseline (date) - Trigger for escalation -	
Metric - Current Baseline (date) - Trigger for escalation -	
Metric - Current Baseline (date) - Trigger for escalation -	
Metric - Current Baseline (date) - Trigger for escalation -	
Metric - Current Baseline (date) - Trigger for escalation -	

Considered and supported by (in line with Scheme of Delegation):	Name	Supported	Comments (if applicable)	Date
Clinical Service Group				
General Manager				
Head of Nursing				
Clinical Lead				
Clinical Care Group				
Service Director				
Assistant Director of Quality, Safety and Patient Experience				
Associate Medical Director				

Other Executive Function (if outside of a CCG e.g. corporate function)				
Director				

QIA Panel Use Only

Considered and supported by:	Name	Supported	Comments (if applicable)	Date
Deputy Director of Health Science	Jon Arthur			
Head of Strategic Partnerships (or nominated deputy)	Anna Bird / Helen Sullivan			
Associate Medical Director for Quality and Safety	Subhamay Ghosh			
Interim Assistant Director of Nursing, Assurance & Safeguarding	Cathie Steele			
Consultant in Public Health Medicine	Michael Thomas			

Considered and approved by Clinical Executive:	Name	Supported	Comments (if applicable)	Date
Director of Nursing, Quality and Patient Experience	Sharon Daniel			
Medical Director	Mark Henwood			
Director of Allied Health and Health Science	James Severs			

Date presented to panel	
Chair of Panel	
Notes of panel discussion	Note: QIA must be reviewed if there is a trigger for escalation is met
If supported by QIA panel, what is the recommended frequency for review	Date for review recommended by QIA panel:

IMPACT MATRIX

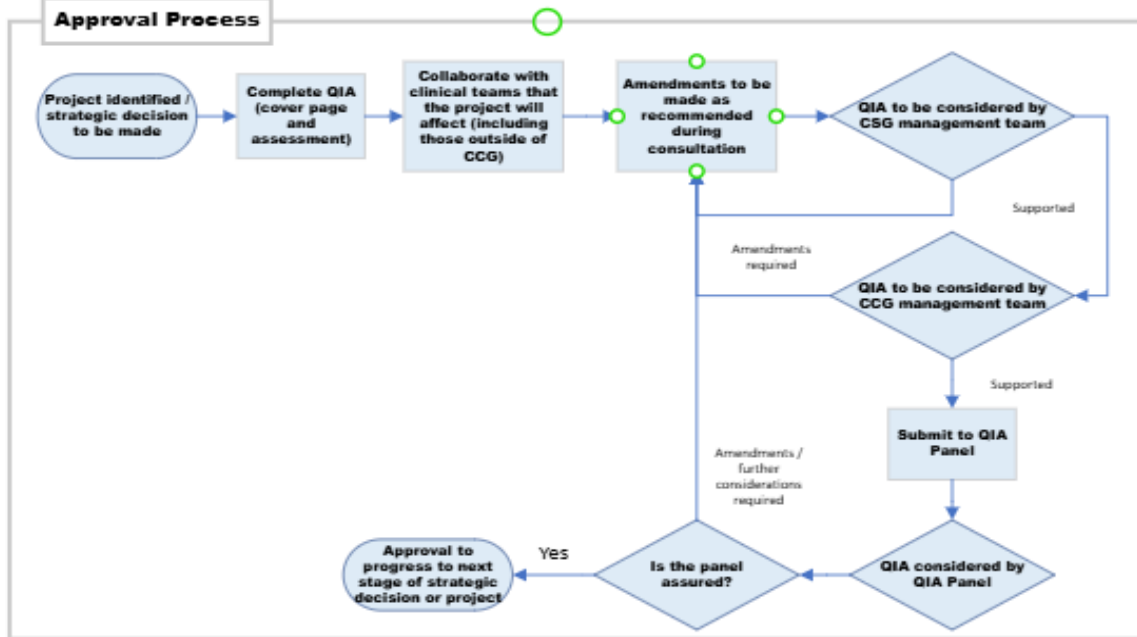
LIKELIHOOD		
1	Rare	Not expected to occur for years. Will occur in exceptional circumstances.
2	Unlikely	Expected to occur at least annually. Unlikely to occur
3	Possible	Expected to occur at least monthly. Reasonable chance of occurring.
4	Likely	Expected to occur at least weekly. Likely to occur.
5	Almost Certain	Expected to occur at least daily. More than likely to occur.

LIKELIHOOD	OPPORTUNITY						IMPACT				
		5	4	3	2	1	-1	-2	-3	-4	-5
	5	25	20	15	10	5	-5	-10	-15	-20	-25
	4	20	16	12	8	4	-4	-8	-12	-16	-20
	3	15	12	9	6	3	-3	-6	-9	-12	-15
	2	10	8	6	4	2	-2	-4	-6	-8	-10
	1	5	4	3	2	1	-1	-2	-3	-4	-5

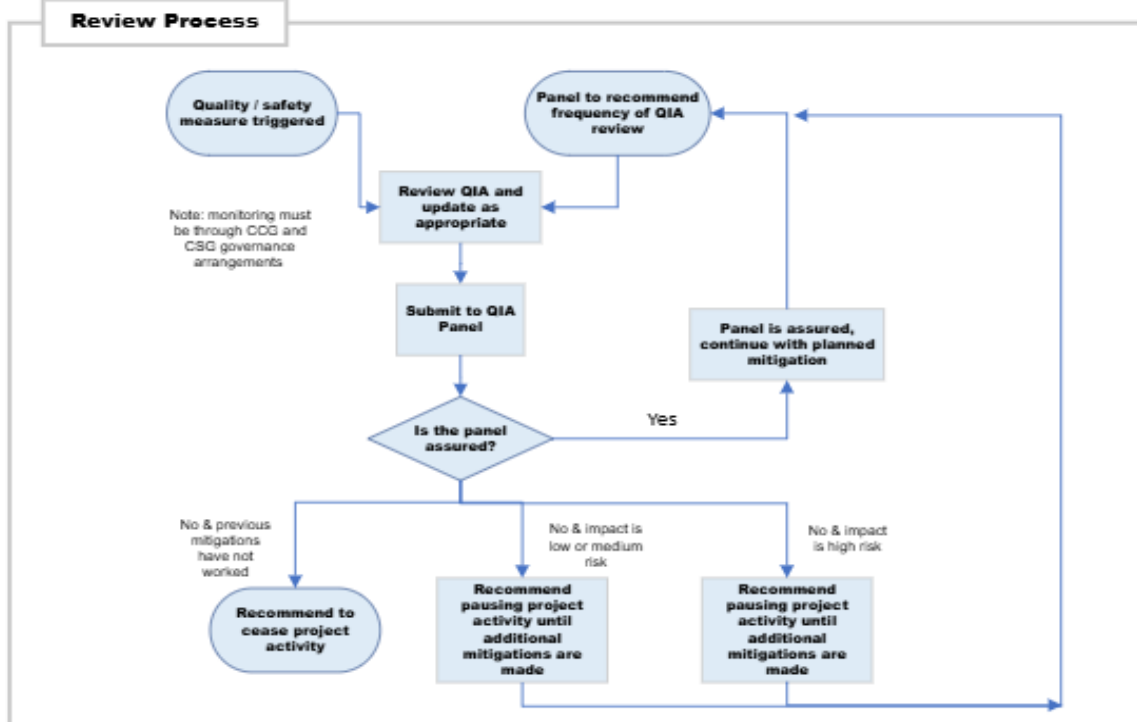
CATEGORY	
Excellent opportunity	Extreme risk
Good opportunity	High risk
Moderate opportunity	Moderate risk
Minor opportunity	Low risk

OPPORTUNITY AND IMPACT			
IMPACT	SCORE	The proposed change is anticipated to lead to the following level of opportunity and/or impact:	
Positive	5	Excellence	Multiple enhanced benefits including excellent improvement in access, experience and/or outcomes. Outstanding reduction in health inequalities by narrowing the gap in access, experience and/or outcomes between people with protected characteristic(s) and the general population. Leading to consistently improved standards of experience and an enhancement of public confidence. Significant improvements to service delivery, performance, financial and workforce sustainability.
	4	Major	Major benefit leading to long term improvements and access, experience and /or outcomes. Major reduction in health inequalities by narrowing the gap in access, experience and /or outcomes between people with protected characteristic(s) and the general population. Benefits include improvements in management of patients with long term effects and compliance with national standards. Substantial improvements to service delivery, performance, financial and workforce sustainability
	3	Moderate	Moderate benefits requiring professional intervention with moderate improvement in access, experience and /or outcomes. Moderate reduction in health inequalities by narrowing the gap in access, experience and /or outcomes between people with protected characteristic(s) and the general population. Moderate improvements to service delivery, performance, financial and workforce sustainability
	2	Minor	Minor improvement in access, experience and /or outcomes for people. Minor reduction in health inequalities by narrowing the gap in access, experience and /or outcomes between people with protected characteristic(s) and the general population. Minor improvements to service delivery, performance, financial and workforce sustainability.
	1	Negligible	Negligible improvement in access, experience and /or outcomes for people with this protected characteristic. Negligible reduction in health inequalities by narrowing the gap in access, experience and /or outcomes between people with protected characteristic(s) and the general population. Negligible improvements to service delivery, performance, financial and workforce sustainability.
Negative	-1	Negligible	Negligible negative impact on access, experience and /or outcomes for people. Negligible increase in health inequalities by widening the gap in access, experience and /or outcomes between people with protected characteristic(s) and the general population. Potential to result in minimal injury requiring no/minimal intervention or treatment, peripheral element of treatment suboptimal and/or informal complaint/inquiry. Negligible negative impacts to service delivery, performance, financial and workforce sustainability.
	-2	Minor	Minor negative impact on access, experience and /or outcomes for people . Minor increase in health inequalities by widening the gap in access, experience and /or outcomes between people with thi protected characteristic(s) and the general population. Potential to result in minor injury or illness requiring minor intervention, formal complaint and overall treatment suboptimal. Minor negative impacts to service delivery, performance, financial and workforce sustainability.
	-3	Moderate	Moderate negative impact on access, experience and /or outcomes for people. Moderate increase in health inequalities by widening the gap in access, experience and /or outcomes between people with protected characteristic(s) and the general population. Potential to result in moderate injury requiring professional intervention, formal complaint and failure to meet internal standards. Moderate negative impacts to service delivery, performance, financial and workforce sustainability.
	-4	Major	Major negative impact on access, experience and /or outcomes for people. Major increase in health inequalities by widening the gap in access, experience and /or outcomes between people with protected characteristic(s) and the general population. Potential to lead to major injury leading to long-term incapacity/disability, multiple complaints/independent review and critical report. Major negative impacts to service delivery, performance, financial and workforce sustainability.
	-5	Catastrophic	Catastrophic negative impact on access, experience and /or outcomes for people with this protected characteristic. Catastrophic increase in health inequalities by widening the gap in access, experience and /or outcomes between people with protected characteristic(s) and the general population. Potential to result in incident leading to death, multiple permanent injuries or irreversible health effects, an event which impacts on a large number of patients, totally unacceptable level or effectiveness of treatment, gross failure of safety and does not meet required standards. Significant negative impacts to service delivery, performance, financial and workforce sustainability.

Approval Process



Review Process



EQUITY IMPACT ASSESSMENT TOOLKIT

Developed by:

Dawn Esther-Davies, Principal Public Health Practitioner
Dr. Ardiana Gjini, Executive Director of Public Health
Dr. Jo McCarthy, Consultant in Public Health Medicine

Special thanks to the Primary Care Division, Public Health Wales

Dr Kerry Bailey, Consultant in Public Health
Fatima Sayed, Principal Public Health Practitioner
Victoria Tice, Senior Public Health Practitioner



INTRODUCTION:

Population health and wellbeing can be influenced by wider socio-economic and environmental factors and their impact vary considerably across different population groups as defined by age, sex, geography and deprivation.

Health equity is a fundamental principle that recognises that everyone deserves the opportunity to achieve their full health potential. However, inequalities in health outcomes continue to persist, disproportionately affecting marginalised and underserved individuals and communities.

WHY DOES HEALTH EQUITY MATTER?

Everyone has the right to live a healthy life, but systemic barriers and injustices such as where you were born, level of education, adverse childhood experiences, community cohesion and access to quality education and housing often prevent certain individuals and population groups from accessing the resources and opportunities they need to achieve optimal health.

Promoting health equity can also lead to better health outcomes for all, as improving overall access to quality healthcare and services can improve population health.

Equity is the absence of avoidable, unfair or remediable differences among groups of people, whether those groups are defined socially, economically, demographically or geographically or by other means of stratification.

World Health Organisation



The route to achieving equity will not be accomplished through treating everyone equally. It will be achieved by treating everyone justly according to their circumstances.

Race Matters Institute

HEALTH INEQUALITIES MAY BE DRIVEN BY:

- Different experiences and distribution of the wider determinants of health or structural factors. For example, the environment, community life, income or housing. In other words, the social economic and environmental conditions in which people live, work and play.
- Different exposure to social, economic and environmental stressors and adversities. These affect states of mind from an early age and throughout life. Stress and psychological wellbeing directly affect resilience, health conditions and health behaviours.
- Differences in health behaviours or other risk factors between groups, for example smoking, diet, and physical activity levels have different social distributions. Health behaviours may be influenced by wider determinants of health, like income.
- Unequal access to or experience of health and other services between social groups.

AIM & OBJECTIVES

To promote fairness, justice and equity in healthcare delivery to improve health outcomes for all

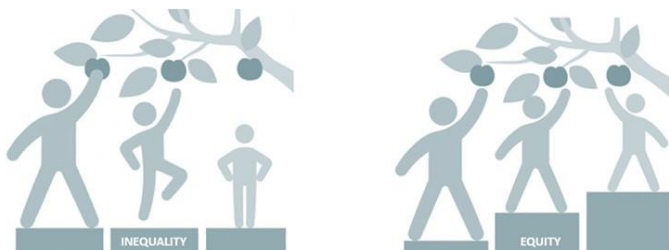
- Identify and understand the social determinants of health that contribute to health disparities.
- Assess the potential impact of health care policies, programmes and services on marginalised and vulnerable populations
- Assess the effectiveness of interventions in reducing health inequities and inequalities
- Engage with community members and stakeholders to ensure their perspectives and experiences are considered
- Develop recommendations and strategies to address identified disparities and promote equitable access to services
- Promote transparency and accountability in decision making processes

Links to Statutory Requirements and Policy

[A Healthier Wales: Long term plan for health and social care](#)
[Wellbeing of Future Generations \(Wales\) Act 2016](#)
[Socio-economic Duty](#)
[Equality Act 2010](#)
[Social Services & Wellbeing \(Wales\) Act 2014](#)
[The Primary Care Model for Wales](#)
[Welsh Language Standards](#)
[NHS Continuing Healthcare Framework](#)
[NHS Planning Framework 2022-2025](#)
[Digital Health and Care Wales: Equality & Diversity Policy](#)
[Public Health \(Wales\) Act 2017 \(legislation.gov.uk\)](#)

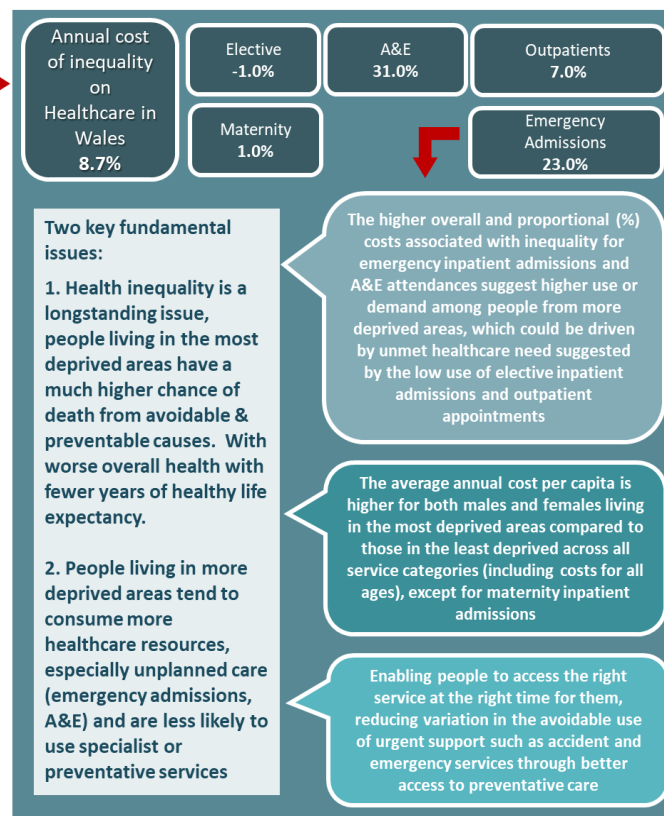
DEFINITIONS

INEQUALITY refers to differences in outcomes between different groups of people, often due to factors such as income, education, race, gender or geographic location. These differences can lead to unequal access to healthcare, education, employment and other essential services leading to a cycle of poverty and social exclusion. Inequality can perpetuate social divisions and create barriers to social mobility. Addressing inequality requires a focus on the social determinants of health and implementing policies and interventions that aim to reduce disparities and promote equity and inclusivity.



EQUITY is the concept of fairness and justice in the distribution of resources and opportunities, with a focus on addressing the needs and circumstances of individuals to ensure everyone has an equal chance to succeed. It involves recognising and acknowledging the systematic barriers that contribute to inequality and highlights the principle of ensuring that all individuals have equal access to services and opportunities to achieve their full potential regardless of their social or economic status. It is about addressing the root causes of inequalities and working towards eliminating barriers to and improving access and quality for all populations.

HEALTHCARE COSTS ATTRIBUTABLE TO INEQUALITIES



IN THE CONTEXT OF THIS TOOLKIT **PROPOSAL** IS USED AS AN **UMBRELLA TERM** TO DESCRIBE A NUMBER OF ACTIONS AND ACTIVITIES INCLUDING STRATEGY DEVELOPMENT AS WELL AS POLICY, PROGRAMME AND PROJECT DEVELOPMENT/REVIEW.

ACTION: DESCRIBE THE CONTEXT AND CONTENT OF THE PROPOSAL, INCLUDING STAKEHOLDERS.

[illegible]

ACTION: USE THE LIST BELOW TO IDENTIFY THE INTENDED TARGET POPULATION FOR THIS PROPOSAL AND IDENTIFY THOSE THAT MAY BE EXCLUDED



WILL THIS PROPOSAL AFFECT THE WHOLE POPULATION, OR WILL THERE BE A DIFFERENTIAL IMPACT ACROSS SUB-GROUPS?	WILL THE IMPACT BE ...				Do you know how to identify your population? (Yes/ No/ Not sure)	SUPPORTING EVIDENCE: What data do you have to help you understand your population/ patient profile? What additional data do you need?
	NEGATIVE	NEUTRAL	POSITIVE	UNKNOWN (E.G. Not yet considered as part of an option or specific project, pathway)		
Populations living in the most deprived communities. (see the WIMD tab)			x			
Rural Populations			x			
Those who miss appointment on a regular basis or are lost to follow-up?			x			
Those who access emergency services or A&E on a regular basis for non-urgent care that could be managed in the community?			x			
Those with Protected Characteristics under the Equality Act 2010						
- Age			x			
- Disability		x				
- Sex (male/female)		x				
- Marriage and Civil Partnership		x				
- Race & Ethnicity		x				
- Pregnancy & Maternity		x				
- Sexual Orientation		x				
- Religion or Belief		x				
- Gender Reassignment		x				
Vulnerable and Inclusion Health Groups						
People who are experiencing or are at risk of homelessness (may also include those in temporary accommodation or care leavers)			x			
People with drug and alcohol dependence			x			
Vulnerable migrants, refugees, asylum seekers, undocumented migrants			x			
Gypsy, Roma, and Traveller communities			x			
People in contact with the justice system			x			
Victims of modern slavery			x			
Sex workers			x			
Other Groups						
Welsh Language		x				
Carers		x				
Veterans		x				
Other Groups (Please list below):						

Please see DATA SOURCES tab to access information on how to identify your local population and other statistics

ACTION: USE THIS CHECKLIST TO ASSESS ANY POTENTIAL LINKS BETWEEN THE PROPOSAL, HEALTH AND EQUITY. USE THE RECOMMENDATIONS TABLE TO DETERMINE IF A COMPREHENSIVE EIA IS REQUIRED.

EQUITY IMPACT ASSESSMENT - CHECKLIST		Yes	Unsure	No
DOES THIS PROPOSAL CONSIDER ACCESS FOR...				
	Populations living in the most deprived communities (consider communities in the most deprived 20% of WIMD)?	x		
	Those with protected characteristics (Equality Act, 2010)?	x		
	Vulnerable and inclusion health groups (e.g. homeless, gypsy, Roma and traveller communities, sex workers, people in contact with the justice system)?	x		
	Rural populations?	x		
	Those who miss appointments on a regular basis?	x		
	Those who access urgent care on a regular basis for conditions that could be managed in the community?	x		
WILL THIS PROPOSAL ADDRESS PREVENTION ...				
	Through assessing the specific health risks and needs of our most vulnerable populations	x		
	By raising awareness of risk factors for certain health conditions, promoting healthy lifestyles and improving the availability of preventative health services (e.g. vaccinations, health check-ups)	x		
	Through promoting screening and early detection and addressing disparities in uptake/coverage between groups	x		
	Through improving care for those already suffering from chronic conditions (e.g. improving access and support mechanisms, monitoring outcomes in key groups)	x		
DOES THIS APPROACH OPTIMISE INTERVENTIONS AND ASSESS EFFECTIVENESS ...				
	Through using data to understand key factors in the local population that may contribute to poor health or high service utilisation. (e.g. an ageing population, areas of deprivation, higher prevalence of lifestyle risk factors)?	x		
	Through monitoring service utilisation to assess healthcare service use by various characteristics?	x		
	Through systematically assessing impact on inequalities and monitoring changes over time? (e.g. pilot testing, evaluation, research, establishing benchmarks)	x		
	Ensuring resource /financial implications of the programme/intervention/option in terms of workforce demand, skills, training and programme costs are considered?	x		
DOES THIS APPROACH CONSIDER PARTNERSHIPS AND SOCIAL ACCEPTABILITY THROUGH...				
	Strengthening and expanding partnerships to enhance health equity? (e.g. multi-sectoral, community working)	x		
	Working with those that are likely to be affected by the outcome to ensure they are engaged in the process? (e.g. Collaboration, coproduction, engagement)	x		
	Making a commitment to health equity through the development of inclusive policy and practice that is fair and transparent?	x		
	Ensuring that organisations and teams understand their responsibilities to tackling inequalities?	x		
	Collecting data to monitor patient experience?	x		
COULD THIS APPROACH WIDEN INEQUALITIES BY...				
	Not tackling the full spectrum of causes (e.g. behavioural risk factors, social determinants)?			x
	Not being co-designed (e.g. engaging stakeholders)?			x
	Relying on professional led interventions (e.g. can reinforce existing structures & barriers)?			x
	Not recognising the economic impact of ill health and treatment (e.g. travel cost, time off work)?			x
	Failing to ensure that health information is delivered in a way that empowers people to make informed decisions to meet their needs? (e.g. Health literacy, communications strategy)			x

RECOMMENDATIONS: Decisions should be based on the nature of the impact (e.g. a specific population groups or those with certain conditions are excluded etc). Which of the following recommendations apply to your initial assessment using the Rapid HEIA

RECOMMENDATION 1. There is likely to be only negligible or small health impacts following the implementation of this proposal. The effect is expected to be small and will not have an impact on equity	There is no need to adjust the proposal or to proceed to the comprehensive EIA. It may be useful to think about how this proposal will be monitored over time and what actions should be taken if a differences in health outcomes occur.
RECOMMENDATION 2. There is likely to be some potential effect as a result of the proposal. There is no need to adjust the proposal or to proceed with a full Health Equity Impact Assessment.	There is no need to adjust the proposal or to proceed with the comprehensive EIA, however, it is important to think about the key areas where a potential difference in outcomes may occur e.g. some groups have not been fully considered or there has been a lack of engagement with partners/service users). Ensure you develop a strategy to monitor health outcomes and consider what actions should be taken if a difference in health outcomes occur.
RECOMMENDATION 3. There is likely to be a significant effect on health outcomes. Further analysis should be undertaken to address difference in health outcomes and unmet needs.	Go to STEP 4 and complete the comprehensive EQUITY IMPACT ASSESSMENT

[illegible]

DATA SOURCES

Link to PHW Public Health Outcomes Framework

[Public Health Outcomes Framework \(2022\) - Public Health Wales \(nhs.wales\)](#)

ARCH Health Needs Assessment

[ARCH Health Needs Assessment](#)

Deprivation data:

[WIMD - Explore \(gov.wales\)](#)

[WIMD - Hywel Dda University Health Board \(gov.wales\)](#)

[Welsh Index of Multiple Deprivation \(WIMD\) 2025 results report \[HTML\] | GOV.WALES](#)

**Click on the links below to access population profile data by Unitary Authority,
Community or Middle Super output Area:**

[Pembrokeshire \(W06000009\) - ONS](#)

[Ceredigion \(W06000008\) - ONS](#)

[Carmarthenshire \(W06000010\) - ONS](#)

The Welsh Index of Multiple Deprivation (WIMD) is an official measure of relative deprivation for small areas in Wales. It ranks areas from 1 (most deprived) to 1909 (least deprived) and is designed to identify areas with the highest concentrations of different types of deprivation.

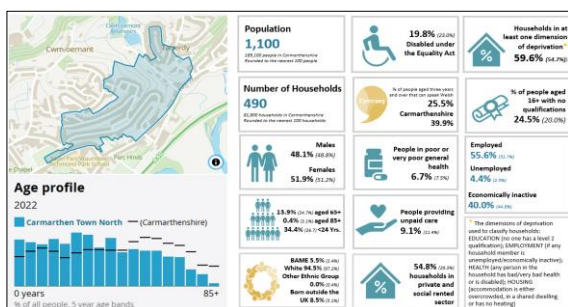
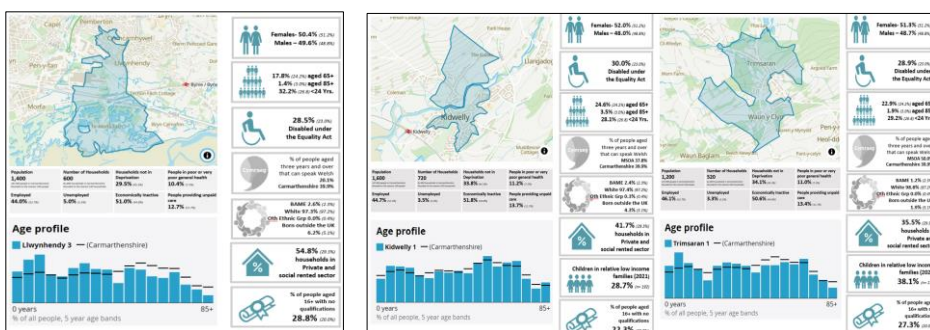
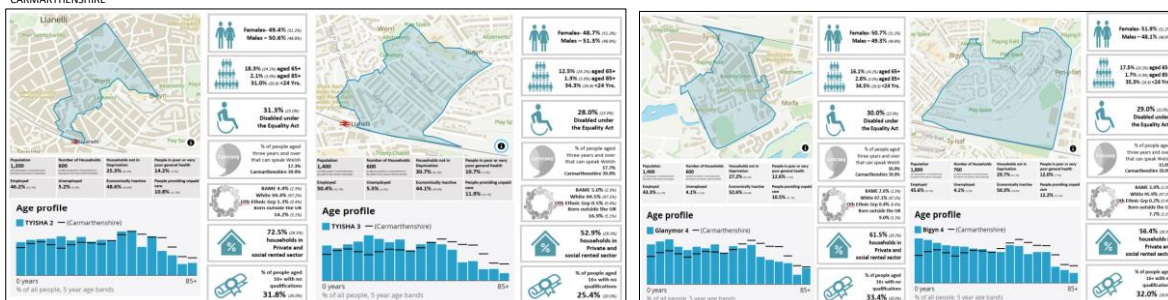
The profiles below are based on the previous release of WIMD data. For the most recent updates got to

[Welsh Index of Multiple Deprivation \(WIMD\) 2025 results report \[HTML\] | GOV.WALES](#)

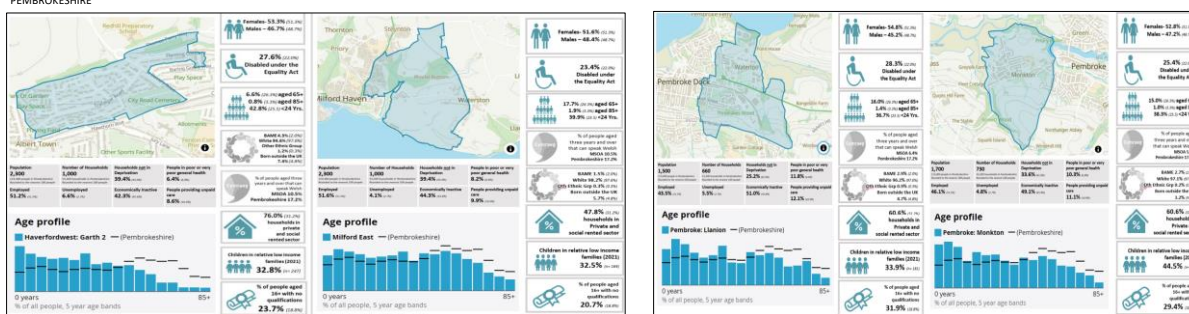
WIMD Indicator data from 2019

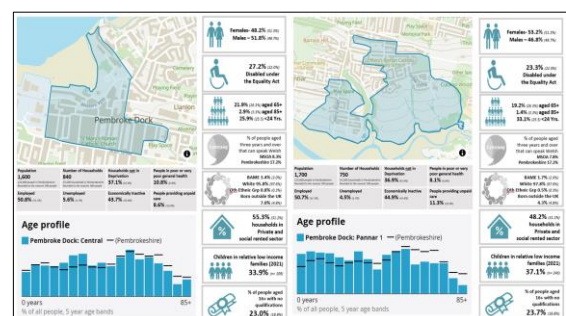
Communities in the most deprived 20% WIMD (RANK)

CARMARTHENSHIRE



PEMBROKESHIRE





Age profile

Carraig Ghaibhiann (Blue) | Ceredigion (Light Blue)

Age Group	Carraig Ghaibhiann (%)	Ceredigion (%)
0 years	~1.5	~1.5
1-4 years	~1.5	~1.5
5-9 years	~1.5	~1.5
10-14 years	~1.5	~1.5
15-19 years	~1.5	~1.5
20-24 years	~1.5	~1.5
25-29 years	~1.5	~1.5
30-34 years	~1.5	~1.5
35-39 years	~1.5	~1.5
40-44 years	~1.5	~1.5
45-49 years	~1.5	~1.5
50-54 years	~1.5	~1.5
55-59 years	~1.5	~1.5
60-64 years	~1.5	~1.5
65-69 years	~1.5	~1.5
70-74 years	~1.5	~1.5
75-79 years	~1.5	~1.5
80+ years	~1.5	~1.5

% of all people, 5 year age bands

Children in relative low income families

Carraig Ghaibhiann: 43.4%
Ceredigion: 33.3%

Age profile

Carraig Ghaibhiann (Blue) | Ceredigion (Light Blue)

Age Group	Carraig Ghaibhiann (%)	Ceredigion (%)
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35-39 years	~1.5	~1.5
40-44 years	~1.5	~1.5
45-49 years	~1.5	~1.5
50-54 years	~1.5	~1.5
55-59 years	~1.5	~1.5
60-64 years	~1.5	~1.5
65-69 years	~1.5	~1.5
70-74 years	~1.5	~1.5
75-79 years	~1.5	~1.5
80+ years	~1.5	~1.5

% of all people, 5 year age bands

Children in relative low income families

Carraig Ghaibhiann: 43.4%
Ceredigion: 33.3%

Use the table below to think about mitigation strategies to ensure interventions are effectively designed, at the intervention and organisational level and across sectors to ensure access to relevant groups and resources.

Intervention	Organisation	Alignment/Collaboration
Modifications that support or supplement a reduction in health inequities:	Modifications that support a reduction in health inequities:	With complementary initiatives that might help reduce inequities
Interventions and approaches that support access to programmes/services e.g. time and location	Population Health assessment (to understand needs and prioritise action)	Internal to the organisation
Stakeholder engagement & Co-design. Priority group participation in the service development, programme delivery or policy implementation	Surveillance - establish systems to monitor health outcomes by demographic groups, allowing for targeted interventions to address identified gaps.	local agencies and other services - partner with local organisations to provide support (e.g. housing issues, addressing risk factors)
Reducing barriers to those who can benefit most from the service e.g. time, location etc	Research and knowledge exchange	Local, regional and national stakeholders
Provision of additional supports e.g. Expand telehealth services to allow remote areas to consult with healthcare providers without the need for travel.	Programme evaluation	Integrated care models and pathways
Communication plan - to raise awareness of health issues or promote awareness of services. Create easy to understand resources to teach individuals how to navigate the healthcare system and manage their health.	Staff education and development - educate providers on the unique health challenges faced by specific population and strategies to reduce disparities in care.	
	Internal policies and procedures	

Addressing health equity in healthcare settings requires a multi-level approach that recognises the complex interplay of facilitators and barriers. By identifying and leveraging facilitators while working to mitigate the barriers, healthcare organisations could make a significant contribution toward achieving health equity for all

In planning your EIA think about some of the barriers and facilitators that may have an impact on implementing this proposal

	SYSTEM LEVEL	ORGANISATIONAL LEVEL	OPERATIONAL LEVEL
FACILITATORS	- Mandated use of the tools (e.g. Public Health (Wales) Act, 2017 - Policy frameworks – NHS long-term plans that emphasis reducing inequalities provide a framework for action. - Funding/ grants aimed at addressing health inequalities. - Increased political and public awareness about health equity – this can be used to pressure healthcare systems to prioritise these concerns - Embed health equity into performance management incentives - National/regional inequality targets	- Leaderships commitment – a strong commitment from healthcare leaders and management to prioritise health equity and drive organisational change. - Training and education for staff on health equity principles and cultural competency to enhance understanding and implementation - Interdisciplinary collaboration across departments and specialties to support a more comprehensive approach to health equity, - Integrate health equity metrics into performance evaluations and organisational aims.	- Community engagement through actively involving community members in the design and delivery of health services that can lead to more equitable outcomes - Develop targeted programmes that address the specific needs of diverse populations. - Use technology including telehealth and digital health solutions to improve access to care for underserved populations. - Implement systems for collecting feedback from patients about their health care experiences and identify gaps and inform improvements.
	SYSTEM LEVEL	ORGANISATIONAL LEVEL	OPERATIONAL LEVEL
BARRIERS	- Fragmentation of services and a lack of co-ordination between health and social care or primary and secondary care can hinder comprehensive approaches to health equity. - Insufficient data on health inequalities can make it difficult to identify needs and measure progress. - Resource constraints and budget cuts can restrict the ability to implement programmes focussed on health equity. - Policy inconsistencies including variability in health equity policies across different regions can lead to unequal implementation and outcomes. - Political pressures to adopt programmes or policies without equity consideration.	- Lack of organisational commitment and readiness and a resistance to change - Siloed decision-making which can limit sharing of best practice and collaborative approaches to addressing health inequalities - Lack of training and awareness among staff regarding health equity concepts - Competing priorities - Lack of buy in from top management - Lack of discussion on the importance of applying health equity approaches at management level.	- Access issues – including transportation, language and cultural differences can prevent marginalised groups from accessing healthcare services - Limited workforce diversity which may result in a lack of cultural competency necessary to effectively engage with diverse populations - Administrative burdens including a lack of resources (financial/human) - Short-term focus especially on immediate outcomes rather than long-term equity goals - Capacity of the health care sector to conduct research, access different types of information and analyse data. - Lack of data to support consensus - Time constraints

Equality Impact Assessment (EqIA) Screening Template

When to complete an EqIA Screening

An EqIA Screening Template must be completed when reviewing, changing and developing procedures/ proposals/ projects/ policies. This is a first step and is used to consider whether there are any negative impacts that may arise.

Purpose of an EqIA Screening Template

The purpose of this short exercise is to ensure that you have shown appropriate due regard when considering the impact for people with protected characteristics in your decision making. The screening process is designed to help you consider the circumstances and to inform evidence-based decisions.

If the proposal is of a significant nature and it is apparent from the outset that a full EqIA will be required, then it is not necessary to complete this Screening Template, you can proceed to complete the full [EqIA](#).

If no negative impacts are identified following completion of the EqIA screening then it is not necessary to undertake a full EqIA however, the decision and justification must be clearly recorded in this document.

On completion of the Screening Template:

- Ensure that all the white boxes within the screening are completed.
- Ensure that the Procedure/ Project/ Proposal/ Policy owner has signed and dated the Screening Template.
- Send a copy of the completed template along with the related policy or project proposal to Inclusion.hdd@wales.nhs.uk for the Diversity & Inclusion Team to review.
- Each Screening Template will be reviewed by the Diversity & Inclusion Team and feedback will be provided to the Procedure/ Project/ Proposal/ Policy owner. This may include recommendations for further action to inform robust decision-making.

Support

For further support please visit the [EqIA Sharepoint](#) or contact:

Email: Inclusion.hdd@wales.nhs.uk

Tel: 01554 899055

Director and Directorate	Ardiana Gjini – Executive Director of Public Health
Service Area	Public Health

Title of Procedure, Project, Proposal, Policy being screened:	Neighbourhood Prevention and Cancer Support Programme

Description of the Procedure/ Project/ Proposal/ Policy being screened (including key aims and objectives)

Hywel Dda University Health Board (HDdUHB), regional public and third sector partners are committed to a social model for health, recognising that health and wellbeing are shaped by social, economic and environmental conditions, not just illness or medical care. This includes recognition of the need to develop place-based integrated service models that meet specific community needs.

HDdUHB wants to put this approach into practice through the 20four7 prevention model. This means focusing support on the most deprived communities, addressing the main preventable risk factors for poor health (smoking, poor diet, harmful alcohol use and physical inactivity) and applying this work across seven priority areas, including cancer.

West Wales already has strong partnerships that can support better health and wellbeing. This proposal aims to build on those strengths by developing a coordinated neighbourhood model for rural communities, connecting prevention, community support, clinical services and local assets. This is important because one in three people in Wales live in rural areas. The proposal therefore offers an opportunity to test a Welsh neighbourhood-based model: one designed around the realities of smaller rural towns and dispersed populations, access barriers and local community strengths.

A three-year investment would allow partners to test and improve the model, gather evidence of impact, and lead change enabling longer-term funding. The aim is to improve cancer outcomes, reduce inequalities and strengthen community resilience in West Wales.

A proposal is being developed by HDdUHB, Macmillan Cancer Support and regional partners and communities to explore the use of social investment through the Neighbourhood Transformation Fund. The proposed three-year programme would act as a catalyst for system transformation, providing opportunity to develop, implement and evaluate an integrated neighbourhood model focused on cancer prevention, early intervention and wider population health improvement.

The programme would support implementation and testing of:

- Community Health & Wellbeing Workers
- Health Coaching and Lifestyle Medicine
- Community Capacity and Neighbourhood Infrastructure

Alongside direct service delivery, the investment would support community engagement, partnership development, workforce capability, project management and evaluation, and the strengthening of local neighbourhood infrastructure. The intention is to build clear evidence of impact, value and return on investment, creating a stronger basis for sustained action through Health Board investment, pooled budgets and wider partnership funding across West Wales. The learning would also have wider national relevance, including for other rural areas.

Evidence considered (including staff and population data, relevant research, expert and community knowledge etc.)

Scoping searches on health coaching and community health and wellbeing workers (CHWWs)

Two scoping searches were undertaken by a librarian in May 2026 to support preliminary work for this proposal.

HEALTH COACHING

A search of the internet identified the NHS England *Workforce Development Framework for Health and Wellbeing Coaches*, which contains potentially relevant references. Additional searches were conducted in Epistemonikos (a database of systematic reviews), the Health Management Information Consortium (HMIC), and MEDLINE. In total, one hundred and eight references were identified.

NHS England (2023) *Workforce development framework for health and wellbeing coaches*. London: NHS England. Available at: <https://www.england.nhs.uk/long-read/workforce-development-framework-for-health-and-wellbeing-coaches/> (Accessed: May 2026).

COMMUNITY HEALTH AND WELLBEING WORKERS (CHWWs)

A search of the internet exploring links between CHWWs and deprivation retrieved several relevant papers. This was followed by a search of the Health Management Information Consortium (HMIC), which identified a further three papers. Further searches were undertaken in the MEDLINE and Embase databases, identifying further studies; however, the majority were conducted outside the United Kingdom. MEDLINE uses a broad definition of community health workers, whereas Embase lacks a specific subject heading; therefore, the following phrase search was employed to help define CHWWs: A total of two hundred and seventy-one references were identified.

Assess which protected characteristics will potentially be affected by the proposal in the table below (please ✓ the relevant box to confirm positive, negative or no impact).

If at any point a negative impact has been identified (actual or potential), you do not need to proceed with the completion of this form, as a full EqlA must be undertaken: [Equality Impact Assessments \(EqlAs\) \(sharepoint.com\)](https://sharepoint.com)

Age			
Is it likely to affect older and younger people in different ways or affect one age group and not another?			
Positive Impact	✓	Negative Impact	No Impact
Justification of impact identified: The programme is designed to improve access to cancer prevention, early intervention, health coaching and community-based support within rural communities. Older people are more likely to be affected by cancer and long-term health conditions and may face barriers to accessing services due to isolation, transport difficulties or limited local provision. Younger people and working-age adults may also benefit from improved access to prevention, health promotion and lifestyle support aimed at reducing future cancer risk. By delivering neighbourhood-based support and strengthening community infrastructure, the			

programme has the potential to improve health outcomes, increase access to services and reduce inequalities across different age groups.					
Disability					
Is it likely to affect those with a physical disability, learning disability, sensory loss or impairment, mental health conditions, long-term medical conditions such as diabetes?					
Positive Impact		Negative Impact		No Impact	✓
Justification of impact identified:					
There is no evidence to suggest that the programme will discriminate against or disadvantage anyone because of a disability, including physical disability, learning disability, sensory impairment, mental health condition, or long-term medical condition.					
Gender Reassignment					
Is it likely to affect those who either:					
- Have undergone, intend to undergo or are currently undergoing gender reassignment. - Do not intend to undergo medical treatment but wish to live in a different gender from their gender at birth					
Positive Impact		Negative Impact		No Impact	✓
Justification of impact identified:					
There is no evidence to suggest that the programme will discriminate against or disadvantage anyone because of their gender. The programme is expected to apply equally and is not anticipated to affect men and women differently.					
Marriage / Civil Partnership					
Under the Equality Act, the characteristic of Marriage and Civil Partnerships is only protected in the workplace/ employment.					
Is it likely to affect those who are married or in a Civil Partnership? This means someone who is legally married or in a civil partnership.					
Positive Impact		Negative Impact		No Impact	✓
Justification of impact identified:					
This group is only in relation to workplace and employment and therefore has no relevance when using this programme. The proposal is not expected to affect people differently based on this protected characteristic.					
Pregnancy and Maternity					
Is it likely to affect those who are pregnant or have recently had a baby? Maternity covers the period of 26 weeks after having a baby, whether or not they are on Maternity Leave.					
Positive Impact		Negative Impact		No Impact	✓
Justification of impact identified:					
There is no evidence to suggest that the programme will discriminate against or disadvantage people who are pregnant or who have recently had a baby. The proposal is not expected to affect access, experience or outcomes differently for this group.					
Race / Ethnicity					
Is it likely to affect people of a different race, nationality, colour, culture or ethnic origin including non-English / Welsh speakers, Gypsies/Travellers, asylum seekers and migrant workers?					
Positive Impact		Negative Impact		No Impact	✓
Justification of impact identified:					
There is no evidence to suggest that the programme will discriminate against or disadvantage people because of race, nationality, colour, culture, ethnic origin, or					

language. Consideration should continue to be given to communication needs, including Welsh language and accessible information where relevant.					
Religion or Belief					
Is it likely to affect people who have a religion or belief? The term 'religion' includes a religious or philosophical belief.					
Positive Impact	☐	Negative Impact	☐	No Impact	✓
Justification of impact identified:					
There is no evidence to suggest that the programme will discriminate against or disadvantage people because of their religion, belief, or philosophical belief. The proposal is not expected to affect people differently on this basis.					
Sex					
Is it likely to affect people who are mostly male or female. Where it applies to both equally does it affect one differently to the other?					
Positive Impact	☐	Negative Impact	☐	No Impact	✓
Justification of impact identified:					
There is no evidence to suggest that the programme will discriminate against or disadvantage anyone because of their sex. The programme is expected to apply equally and is not anticipated to affect men and women differently.					
Sexual Orientation					
Whether a person's sexual attraction is towards their own sex, the opposite sex or either.					
Positive Impact	☐	Negative Impact	☐	No Impact	✓
Justification of impact identified:					
There is no evidence to suggest that the programme will discriminate against or disadvantage people because of their sexual orientation. The proposal is not expected to affect people differently on this basis.					
Armed Forces Community					
Consider whether this impacts on members of the Armed Forces and their families, whose health needs may be impacted long after they have left the Armed Forces and returned to civilian life. Also consider their unique experiences when accessing and using day-to-day public and private services compared to the general population. It could be through 'unfamiliarity with civilian life, or frequent moves around the country and the subsequent difficulties in maintaining support networks, for example, members of the Armed Forces can find accessing such goods and services challenging.'					
For a comprehensive guide to the Armed Forces Covenant Duty and supporting resource please see: Armed-Forces-Covenant-duty-statutory-guidance					
Positive Impact	☐	Negative Impact	☐	No Impact	✓
Justification of impact identified:					
There is no evidence to suggest that the programme will discriminate against or disadvantage members of the Armed Forces Community, including serving personnel, reservists, veterans and their families. The programme is not expected to create barriers linked to Service life, such as frequent relocation, unfamiliarity with civilian services, or difficulties maintaining support networks.					
Socio Economic Duty					
Consider those on low income, economically inactive, unemployed or unable to work due to ill-health. Also consider people living in areas known to exhibit poor economic and/or health indicators and individuals who are unable to access services and facilities. Food / fuel poverty and personal or household debt should also be considered.					

For a comprehensive guide to the Socio-Economic Duty in Wales and supporting resources please see: more-equal-wales-socio-economic-duty			
Positive Impact	✓	Negative Impact	No Impact
Justification of impact identified:			
The programme has the potential to have a positive impact on people experiencing socio-economic disadvantage by supporting improved access, experience and/or outcomes for individuals and communities who may face barriers linked to ill-health, food poverty or living in areas with poorer economic and health indicators. Implementation will consider how the programme can support fair access and reduce inequality of outcome where possible. Any emerging evidence of unequal access or disadvantage for socio-economically disadvantaged groups will be monitored and addressed through programme governance arrangements.			
Welsh Language			
Is it likely to impact on opportunities for people to use the Welsh language? The Welsh language should be treated no less favourably than the English language.			
Positive Impact		Negative Impact	No Impact ✓
Justification of impact identified:			
There is no evidence to suggest that the programme will negatively impact opportunities for people to use the Welsh language. The Welsh language will continue to be treated no less favourably than the English language, and bilingual communication requirements will be considered where relevant. If the programme involves public-facing materials, engagement, communication, training, digital content or service delivery, Welsh language requirements will be reviewed to ensure compliance with organisational Welsh language duties and standards.			

If a negative impact has been identified, you are not required to complete this form as a full EqlA must be undertaken. A full EqlA template and guidance can be found on the following link: [Equality Impact Assessments \(EqlAs\) \(sharepoint.com\)](#)

Screening Completed by:	Name	Felix Johnbritto
	Title	Project Manager
	Contact details	Felix.Johnbritto2@wales.nhs.uk
	Date	20/07/2026
Screening Authorised by: (Directorate level owner of the procedures/ proposals/ projects/ policy)	Name	Bruce Bolam
	Title	Deputy Director of Public Health
	Contact details	Bruce.Bolam@wales.nhs.uk
	Date	22/07/2026
Guidance has been provided by Diversity & Inclusion Team:	Name	Alan Winter
	Title	Senior Diversity & Inclusion Officer
	Contact details	Alan.winter@wales.nhs.uk
	Date	24/7/2026
Diversity and Inclusion Team additional Comments:		

Please note: The D&I team will save a copy of the completed form for reference. If any changes are made after the date of review, it is the directorate's responsibility to update the EqIA and inform the D&I team.