Background

Primary care access and quality impact all 329 million Americans. Geographic access varies significantly and is often lower in areas with a higher proportion of people of color, adding to health disparities.

Defining Primary Care

Primary care, widely identified as the cornerstone of the health care system, serves as a usual source of care focused on both acute and chronic disease detection, management, treatment, and prevention.¹ Efforts to define primary care often start with a broad scope of services and general attributes and are often described in contrast to health care services provided for acute or urgent needs or within a hospital or surgical setting.

The concept of primary care was first introduced in the 1920s and described by the Institute of Medicine (IOM) in 1978 as being "accessible, comprehensive, coordinated, continuous, and accountable." Barbara Starfield further describes primary care as being characterized by first-contact care and being longitudinal and comprehensive. Primary care can further be defined as including advocacy, taking place in the context of a community context and family, including goal-oriented care and health promotion, being integrated, and being based on a relationship. In many studies, primary care is defined by four Cs: first-contact care that is comprehensive in addressing a wide variety of issues from sprains to behavioral health to prenatal care, is continuous with multiple touch-points over time, and is coordinated. In order to know whether primary care spend is increasing in the state, Washington must first develop an agreed upon definition of primary care that will allow for accurate measurement.

The IOM categorizes possible definitions into care provided by certain clinicians, a particular set of activities, a level or setting of care, the attributes themselves, or as a strategy for organizing a system.⁵ More simply, primary care can be defined broadly as consisting of the care provided by a subgroup of medical providers, the set of functions that providers within and outside of that subgroup perform, and/or a general orientation of a health delivery system.⁶ A family medicine physician may order a thyroid test which would be considered part of primary care while an endocrinologist ordering that same test may not necessarily be considered primary care. These provider, service, and system categories have been expanded by Millbank into:⁷

- Provider: All the services delivered by pre-defined primary care providers in an ambulatory setting.
- Service: Services that meet particular definitions including being: comprehensive, first-contact for a wide variety of (not limited) conditions, coordinated, and taking place over time (longitudinal).
- Service: All office visits and preventative services within a category independent of the provider type.
- Service and Provider: Based in claims, specific set of pre-defined services delivered by predefined primary care providers not limited to an ambulatory setting.
- Health systems: Primary care delivered at a system level, useful for capitated systems but most difficult to measure.

Access and Outcomes

Access to primary care has multiple dimensions; identified as: availability or resources, accessibility or how close a delivery site is to where a person lives or works, accommodation or the hours that the delivery site operates, affordability or cost of care and acceptability or how well the care that is offered matches a person's individual needs and preferences such as through the availability of care in different languages. Accessibility is the most well studied in association with individual and population health, consistently showing a positive impact. Early studies in the 1990s found an association between a higher ratio of primary care physicians at a state-level and population-level health outcomes such as lower all-cause mortality and mortality from heart disease, cancer, stroke, as well as infant mortality. Presence of primary care providers is also associated with increased life span, reduction in infant low birth weight, better overall patient experience, and a person's self-rated health. 11,12,13

Access to regular, high-quality care is a challenge for many. Analysis of urban census tracts show lower levels of access to primary care in areas with a higher proportion of black Americans.¹⁴ Those living in rural areas also have lower levels of access to primary care.¹⁵ A primary care delivery site may be located in close geographic proximity but may not be of high quality, may have hours that render it inaccessible, or the providers may not be taking new patients. In a report from the Primary Care Collaborative, the authors note that "consistent and growing evidence shows that primary care-oriented systems achieve better health outcomes, more health equity, and lower costs."¹⁶ A lack of a sufficient primary care workforce is a growing issue that impacts accessibility in Washington State as well as nationally.¹⁷

These issues are influenced and compounded by low reimbursement for primary care compared to specialty care and hospital care, with the United States spending between 5-7% of total health care expenditure on primary care and Washington State spending between 4.4% to 5.6% of total expenditure on primary care. 16,18

Workgroup

The workgroup met from January 2020 to X to develop standards for use in payment models for Washington State.

Recommendation Framework

The workgroup's goal is to foster a common understanding of primary care in order to increase primary care accessibility and availability.

Current Primary Care Definition (providers of, components of, locations of service)	Millbank: Based in claims, specific set of pre-defined services delivered by pre-defined primary care providers not limited to an ambulatory setting.
Aspirational Primary Care Definition	Millbank: Services that meet particular definitions including being: comprehensive, first-contact for a wide variety of (not limited) conditions, coordinated, and taking place over time (longitudinal)
Components of Primary Care with Large Impact	 Care coordination Screenings (e.g., behavioral health, cancer) Chronic care management Medication management
Primary Care Spend (A mechanism for measuring primary care spend)	

Defining a Primary Care Provider

In Washington State, primary care provider is defined as "a general practice physician, family practitioner, internist, pediatrician, osteopathic physician, naturopath, physician assistant, osteopathic physician assistant, and advanced registered nurse practitioner licensed under Title 18 RCW."¹⁹ The workgroup sought to operationalize the four C's described above to develop a standardized set of questions to define primary care providers and primary care.

If primary care is yes to ALL of the following:

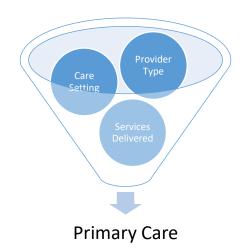
Accountability and advanced clinical judgement for a person's care/panel of patients is lies with one of the following:

- Doctor of Medicine General practice, Family Practice, Internal Medicine, Geriatrics, Pediatrics,
 Adolescent Medicine
- Doctor of Osteopathic Medicine General practice, Family Practice, Internal Medicine, Geriatrics, Pediatrics, Adolescent Medicine
- Advance Registered Nurse Practitioner Family, Adult, Pediatric, Women's Health
- Physician Assistant Family, Adult, Pediatric, Women's Health
- Osteopathic Physician Assistant Family, Adult, Pediatric, Women's Health
- Naturopath
- Certified nurse midwife When providing care outside of managing pregnancy

Team-Based – In addition to the accountable provider listed above, care should be managed by a team. The workgroup does not want to be prescriptive on who must be on the team but emphasizes that care must include behavioral health and a care coordination function. Team members can include: psychologist, psychiatrist, social worker, registered nurse, medical assistant, care coordinator, etc.

First Contact – Does the team assess, triage, and direct a person's health or health care issues as they arise?

Comprehensive – Does the team care for the whole person and provide services that address multiple organ systems including active management of chronic



physical (e.g., COPD, diabetes) and behavioral health (e.g., depression, anxiety, substance use disorder) conditions as well as USPSTF recommended screening and preventive services?

Continuous – Does the team maintain or attempt to develop a longitudinal relationship?

Coordinated – Does the team take responsibility for a person's care through managing a care plan in coordination with a multidisciplinary team and/or with offsite referrals?

Appropriate – Does the team provide evidence-based, person-centered medicine that includes behavioral health?

Figure 2: Care Provided Over the Course of a Person's Life: Doris

Pediatrician Asthma	Family Medicine Asthma		Family Medicine Asthma, high blood pressure	Geriatrician Asthma, HBP	Death
	Breaks leg, set in ED by Emergency Physician	High-risk pregnancy managed by OB/GYN	Breast cancer managed by oncologist	Hospice	

Measurement

Measurement is limited by available data, which is predominantly derived from fee for service claims. Developing a mechanism to measure primary care using the four Cs described above (first contact, comprehensive, continuous, and coordinated) is difficult using claims data as these aspects are reflected in codes that are billed. Further, the lack of a nationally accepted definition of primary care is a major impediment to assessing and increasing the primary care expenditures uniformly across states.

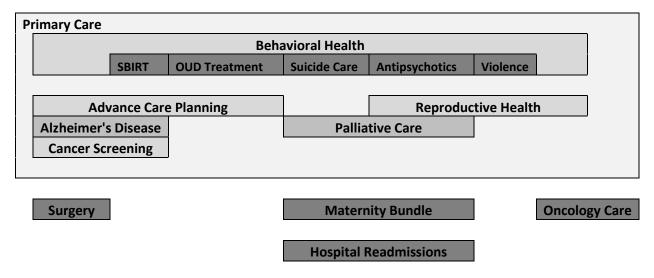
The Washington State Office of Financial Management (OFM) report, mandated by legislation and published in December 2019 notes that comparisons between Washington's percent expenditure and either the national average or other states' reported expenditures are likely misleading due to these differing definitions. For example, Washington does not currently include non-claims-based care (e.g., care coordination activities) in the measure of primary care, but Oregon and Rhode Island do; this may artificially lower Washington's numbers.¹⁸

In order to develop a proxy for what would be true primary care measurement, groups have operated on various assumptions. If defining by provider, the assumption is that a group of subspecialists (i.e., family medicine) <u>always</u> offers primary care and that other groups of subspecialists <u>never</u> provide primary care (i.e., emergency medicine). This assumption holds true for some but not all disciplines. Advance registered nurse practitioners and physician assistants practice in a multitude of settings, including surgical care, which is not reflected in a claim. The 2019 OFM report adjusted the total claims from ARNPs and PAs by 41% and 34%, respectively.

The OFM report presents narrow and broad definitions of primary care, differing based the types of providers who are assumed to be providing primary care. The narrow definition only includes providers who are traditionally considered to perform primary care while the broad definition includes a wider range of provider taxonomy codes includes behavioral health providers, clinical nurse specialists, registered nurses, midwives, and a host of other providers who are not typically considered general practitioners.¹⁸ The OFM stakeholder group also reviewed procedure codes and created both narrow and broad definitions of services qualifying as primary care. Only claims which met both the provider and service definitions of primary care were counted toward the state's total expenditure, with the narrow definition yielding 4.4% and the broad 5.6%.¹⁸

However, the OFM report noted that deficiencies inherent to the Washington All Payor Claims Database claims database, combined with lack of a firm definition for primary care, limit the report's accuracy in some regards. Claims data does not capture, for example, whether or not the location of services provided was a primary care clinic. As was mentioned earlier, Washington lacks a way to measure non-claims-based expenditures. The OFM report mentions a number of other systemic impediments to accurate measurement that may need to be addressed in order to calculate an accurate primary care expenditure percentage for the state.¹⁸

Alignment with Previous Recommendations



Attribution

Decisions for attribution:

unit of analysis (patient versus episode of care); signal for responsibility (professional costs versus number of evaluation and management visits); number of physicians that can be assigned responsibility (single physician versus multiple); and minimum threshold for assigning responsibility (majority of visits or costs versus plurality of visits or costs).²⁰

Recommendations for Stakeholders

Patients and Family Members

- Think about your broad health and wellness-related goals (e.g., being able to attend an upcoming family wedding).
- Where different options are available, like the areas noted in this document, give your provider(s) information about your values and preferences, and discuss options, tradeoffs, and implications of a decision together.
- Ask about whether a patient decision aid is available.
- Ask your care provider about the test or treatment options available, including the option of "doing nothing" or "watchful waiting."

Health Care Delivery Organizations and Systems

- Work with your clinical champion(s) to educate providers about the value of shared decision
 making and how to have a good conversation that uses the patient decision aid or references
 the patient decision aid if the aid will be distributed to patients prior to the visit.
- Select one of the 10 clinical areas to pilot (e.g., breast cancer screening).

Providers

 Participate in skills training. Shared decision making is a learned skill-set that is supported by patient decision aids.

Health Plans and/or Professional Liability Carriers

 Incorporate shared decision making requirements as standards for value-based models (e.g., Centers of Excellence).

Employers

- Incorporate shared decision making requirements as standards for value-based contracting (e.g., Centers of Excellence, Accountable Care Organizations).
- Talk to your health plan about the importance of shared decision making and how to report on use of shared decision making including how to ensure appropriate reimbursement.

Washington State Health Care Authority

• Encourage the patient decision aid developer community to develop patient decision aids for the ten priority areas publicly available at no cost.

Examples from Shared Decision Making Report

Options for tracking shared decision making are below including those aligning with value-based reimbursement models from the Bree Collaborative and Federal programs:

• Shared Decision Making Process

Steward: Massachusetts General Hospital NQF #2962

This measure assesses the extent to which health care providers actually involve patients in a decision-making process when there is more than one reasonable option. This proposal is to focus on patients who have undergone any one of seven common, important surgical procedures: total replacement of the knee or hip, lower back surgery for spinal stenosis of herniated disc, radical prostatectomy for prostate cancer, mastectomy for early stage breast cancer or percutaneous coronary intervention (PCI) for stable angina. Patients answer four questions (scored 0 to 4) about their interactions with providers about the decision to have the procedure, and the measure of the extent to which a provider or provider group is practicing shared decision making for a particular procedure is the average score from their responding patients who had the procedure.

Informed, Patient-Centered Hip and Knee Replacement Surgery

NQF #2958

Steward: Massachusetts General Hospital

The measure is derived from patient responses to the Hip or Knee Decision Quality Instruments. Participants who have a passing knowledge score (60% or higher) and a clear preference for surgery are considered to have met the criteria for an informed, patient-centered decision. The target population is adult patients who had a primary hip or knee replacement surgery for treatment of hip or knee osteoarthritis.

Appendix C: Guideline and Systematic Review Search Results

	Year	Title	Summary or Findings
AHRQ: Research Findings and Reports	2019	Achieving Health Equity in Preventive Services	No eligible studies evaluated effects of provider-specific barriers; 18 studies of population barriers provided low or insufficient evidence regarding insurance coverage, access, age, rural location, low income, language, low health literacy, country of origin, and attitudes. In 12 studies of clinician interventions, screening was higher for colorectal cancer with patient navigation, risk assessment and counseling, educational materials, and decision aids; breast and cervical cancer with reminders involving lay health workers; and cervical cancer with outreach and health education. Clinician-delivered interventions were effective for smoking cessation and weight loss. In 11 studies of health information technologies, automated reminders and electronic decision aids increased colorectal cancer screening, and web- or telephone-based self-monitoring improved weight loss, but other technologies were not effective. In 88 studies of health system interventions, evidence was strongest for patient navigation to increase screening for colorectal (risk ratio [RR] 1.64; 95% confidence interval [CI] 1.42 to 1.92; 22 trials), breast (RR 1.50; 95% CI 1.22 to 1.91; 10 trials), and cervical cancer (RR 1.11; 95% CI 1.05 to 1.19). Screening was also higher for colorectal cancer with telephone calls, prompts, other outreach methods, screening checklists, provider training, and community engagement; breast cancer with lay health workers, patient education, screening checklists, provider training, and community engagement; breast cancer with lay health workers, patient education, screening checklists, provider training, and community engagement; breast cancer with patient navigation. Trials of smoking cessation and obesity education and counseling had mixed results. In populations adversely affected by disparities, evidence is strongest for patient navigation to increase colorectal, breast, and cervical cancer screening; telephone calls and prompts to increase colorectal cancer screening; and reminders including lay health workers
	2016	Improving Cultural Competence to Reduce Health Disparities	None of the included studies measured the effect of cultural competence interventions on health care disparities. Most of the training interventions measured changes in professional attitudes toward the population of interest but did not measure the downstream effect of changing provider beliefs on the care delivered to patients. Interventions that altered existing protocols, empowered patients to interact with the formal health care system, or prompted provider behavior at the point of care were more likely to measure patient-centered outcomes. The medium or high risk of bias of the included studies, the heterogeneity of populations, and the lack of measurement consensus prohibited pooling estimates or commenting about efficacy in a meaningful or responsible way. The term "cultural competence" is not well defined for the LGBT and disability populations, and is often conflated with patient-centered or individualized care. There are many gaps in the literature; many large subpopulations are not represented.

Cochrane	2012	Narrow band imaging versus conventional white light colonoscopy for the detection of colorectal polyps	We could not find convincing evidence that NBI is significantly better than high definition WLC for the detection of patients with colorectal polyps, or colorectal adenomas. We found evidence that NBI might be better than standard definition WLC and equal to high definition WLC for detection the patients with colorectal polyps, or colorectal adenomas.
	2016	Interventions to encourage uptake of cancer screening for people with severe mental illness	A comprehensive search showed that currently there is no RCT evidence for any method of encouraging cancer screening uptake in people with SMI. No specific approach can therefore be recommended. High-quality, large-scale RCTs are needed urgently to help address the disparity between people with SMI and others in cancer screening uptake.
	2019	Follow-up strategies for patients treated for non-metastatic colorectal cancer	The results of our review suggest that there is no overall survival benefit for intensifying the follow-up of patients after curative surgery for colorectal cancer. Although more participants were treated with salvage surgery with curative intent in the intensive follow-up groups, this was not associated with improved survival. Harms related to intensive follow-up and salvage therapy were not well reported.
	2013	Personalised risk communication for informed decision making about	There is strong evidence from three trials that personalised risk estimates incorporated within communication interventions for screening programmes enhance informed choices. However the evidence for increasing the uptake of such screening tests with similar interventions is weak, and it is not clear if this increase is associated with informed choices. Studies included a diverse range of screening programmes. Therefore, data from this review do not allow us to draw conclusions about the best interventions to deliver personalised risk communication for enhancing informed decisions. The results are dominated by findings

		taking screening tests		topic area of mammography and colorectal cancer. Caution is therefore required in generalis rly for clinical topics other than mammography and colorectal cancer screening.	ing from these results, and
	2017	Decision aids for people facing health treatment or screening decisions	better in risk perc on health when de	ed to usual care across a wide variety of decision contexts, people exposed to decision aids feed formed, and clearer about their values, and they probably have a more active role in decision eptions. There is growing evidence that decision aids may improve values-congruent choices. In outcomes or satisfaction. New for this updated is evidence indicating improved knowledge accision aids are used either within or in preparation for the consultation. Further research is not example the chosen option, cost-effectiveness, and use with lower literacy populations.	making and more accurate There are no adverse effects and accurate risk perceptions
Veterans Administration Evidence- based Synthesis Program	2019	What is the Optimal Panel Size in Primary Care? A Systematic Review	physiciar the num	the Veterans Health Administration Handbook 1101.02 established a baseline panel size of 1, in a Patient Aligned Care Team (PACT). This number could be adjusted up or down based on per of examination rooms, and patient complexity. After adjustment for these factors, panels Health Administration (VHA) Directive 1406 reaffirmed both the baseline panel numbers and	availability of support staff, ranged from 1,000 to 1,400.
Health Technology Assessment Program	None				
Centers for Disease Control and Prevention	N/a				
Institute for Clinical and Economic Review	2008	Computed tomo (CT) colonograp	•	Given the possible benefits of introducing a widely available minimally-invasive option for colorectal cancer screening, there is considerable interest in CTC.	

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