

Instructions

KPI Measures have been separated by Population of Focus (PoF), please report on all sites/clinics/departments that are engaged in making changes as part of the EPT program over the three-year program.

These PoF measures are based on NCQA HEDIS measures, which are used to measure health plan performance. HEDIS uses the calendar year as the measurement period. To adapt this to practice-level reporting required by EPT, you may use a 12-month period off the calendar year cycle, e.g. July-June, or April-March. The specifications below sometimes refer to a certain month in the year; please shift your measurement period accordingly. For example, October refers to the 10th month of any given measurement period.

To obtain more detailed specifications, please visit the following:

- Partnership Health Plan QIP measure specifications (2024 measurement year)
- IEHP 2025 Global Quality for P4P (for PCPs)
- Reach out to your Medi-Cal Managed Care Plan to see if they have measure specifications to share!
- NCQA (Note: NCQA charges a fee for measure specifications)
- CMS Adult specs and code sets
- CMS Child specs and code sets

To jump to specific PoF, click below:

- Depression Screening (Chronic Care, Preventative Needs, Children/Youth)
- Preventative
- Chronic Care
- Children and Youth
- Behavioral Health
- Pregnant People



Depression Screening (Chronic Care, Preventative Needs, Children/Youth)

Measure	Description	Numerator	Denominator
Depression	% of patients	Those in the denominator who had a depression screening result using a	People 12 years of age and older at the
Screening and	aged 12 and	standard screening tool for depression:	start of the measurement period.
Follow-Up for	older who were	Patient Health Questionnaire (PHQ-9, PHQ9M, PHQ-2)	start or the measurement period.
Adolescents	screened for	DHCS recommended screening tool	*For children and youth
and Adults	depression using	Beck Depression Inventory (BDI-II) adults only	People between the age of 12 and 17 at
(DSF)	a standard	Beck Depression Inventory-Fast Screen (BDI-FS)	the start if the measurement period.
(20.7	screening tool	Center for Epidemiologic Studies Depression Scale-Revised	and start in the measurement period.
	and, if positive,	(CESD-R)	
	received follow-	Edinburgh Postnatal Depression Scale (EPDS)	
	up care within 30	PROMIS Depression	
	days.	Duke Anxiety-Depression Scale (DUKE-AD) adults only	
		Geriatric Depression Scale-Short Form & Long Form (GDS) adults	
		only	
		My Mood Monitor (M-3) adults only	
		Clinically Useful Depression Outcome Scale (CUDOS) adults only.	
*Follow up		Those in the denominator who had one of the following within 30 days:	All patients from the Depression
with positive		Outpatient, telephone or e-visit for follow-up for	Screening Numerator with a positive
screening		depression/behavioral health	depression result.
		Depression case management encounter	·
		Behavioral health encounter (assessment, therapy, collaborative	
		care, medication management)	
		A diagnosis of encounter for exercise counseling	
		Dispensed antidepressant medication	
		Documentation of a negative full-length depression screening on	
		the same day as a positive screen on a brief screening tool (i.e., a	
		negative PHQ-9 as a follow-up to a positive PHQ-2).	



Preventative

Measure	Description	Numerator	Denominator	Notes
Colorectal Cancer Screening (COL)	The percentage of patients 45-75 years of age who had an appropriate screening for colorectal cancer.	Those in the denominator who had one or more screenings for colorectal cancer including date and result: Fecal occult blood test (within the year) Stool DNA (sDNA) with FIT test (within past three years) Flexible sigmoidoscopy (within past five years) CT colonography (within past five years) Colonoscopy (within the past 10 years)	People 45–75 years as of the end of the measurement period.	Documentation must include: Need date and type of colorectal cancer screening(s) performed. A result is not required if the documentation is clearly part of the "medical history" section of the medical record. If it is not clear, results or findings need to be provided to show screening was performed and not just ordered. Forms of documentation accepted: Member-reported colorectal cancer screening documented in the medical history (e.g., member reports normal colonoscopy in 2015) Pathology report indicating the type of screening and screening date FOBT, depending on the number of samples and type of test: guaiac (gFOBT) or immunochemical (FIT)
Breast Cancer Screening (BCS)	Percentage of people 50 to 74 years of age who had at least one mammogram to screen for breast cancer in the past two years.	Those in the denominator who had one or more mammograms any time in the 27 months prior to, and including the last day of, the measurement period.	The percentage of patients 52–74 years of age who were recommended for routine breast cancer screening and had a mammogram to screen for breast cancer.	Documentation indicating a mammogram was completed and the date it was performed. Patient reported mammogram is acceptable. Document date in the history or preventive service section of the medical record. If the exact date of the last mammogram is unclear, do not use terms like "approximate" or "about" in your documentation. Instead, record the month and year, or just the year.



Measure	Description	Numerator	Denominator	Notes
Cervical	The percentage	Those in the denominator who had one	People 24–64 years of age	Documentation in the medical record must include
Cancer	of women 21-	or more screenings for cervical cancer,	by the end of the	both of the following:
Screening	64 years of age	including date and result:	measurement period who	A note indicating the date when the cervical
(CCS)	who were	Members 21–64 years of age by	were recommended for	cytology was performed
	screened for	the end of the measurement	routine cervical cancer	Lab result or findings
	cervical cancer	period who had cervical	screening.	
		cytology performed within the		Note: Do not count biopsies because they are
		last 3 years		diagnostic and therapeutic only and are not for
		Members 30–64 years of age by		primary cervical cancer screening. Evidence of
		the end of the measurement		hrHPV testing within the last 5 years also captures
		period who were recommended		patients who had co-testing.
		for routine cervical cancer		
		screening and had cervical high-		
		risk human papillomavirus		
		(hrHPV) testing performed		
		within the last 5 years.		
		Members 30–64 years of age by		
		the end of the measurement		
		period who were recommended		
		for routine cervical cancer		
		screening and had cervical		
		cytology/high-risk human		
		papillomavirus (hrHPV)		
		cotesting within the last 5 years.		



Chronic Care

Measure	Description	Numerator	Denominator	Notes
Controlling High Blood Pressure (CBP)	Percentage of patients, aged 18 to 85 yrs, with hypertension whose blood pressure was adequately controlled (<140/90 mm Hg)	Those in the denominator whose most recent BP reading was adequately controlled (<140/90 mm Hg). Controlled BP reading must occur on or after the date of the second HTN diagnosis. Blood pressure is not adequately controlled if: Systolic blood pressure is >140 mm Hg Diastolic blood pressure is >90 mm Hg No blood pressure is recorded Either the systolic or diastolic value is missing	18- to 85-year-old people with hypertension as of the end of the measurement period.	Documentation must include: Most recent systolic BP is <140mm Hg AND diastolic BP is <90 mm Hg. If multiple BP measurements occur on the same date or noted in the chart on the same date, use the lowest systolic and lowest diastolic BP reading. BP can be taken by any staff member or the PCP. Patient documented BP in the chart counts if patient used a digital device.
Glycemic Status Assessment for Patients with DM >9% (GSD)	Percentage of 18- to 75-year-old people with diabetes whose most recent glycemic status (hemoglobin A1c [HbA1c] or glucose management indicator [GMI]) was not under control (>9.0%).	Those in the denominator whose most recent glycemic status assessment (HbA1c or GMI), taken during the measurement period and after the diagnosis, was: Inadequately controlled (>9%) Missing a result (i.e., there is a date the test was conducted with no result). Glycemic status assessment was not done during the measurement period. If there are multiple glycemic status assessments on the same date, use the lowest result.	18- to 75-year-old people with diabetes as of the end of the measurement period.	**This is an inverse measure, meaning that a lower rate indicates higher performance** Documentation in medical record or lab result must include a note indicating the date when the glycemic assessment was performed and result or finding during the measurement year



Children and Youth

Measure	Description	Numerator	Denominator	Notes
Child Immunization Status (CIS) – Combo 10	Percentage of two-year-old children who have received the 10 recommended vaccines.	Those in the denominator who have received the 10 recommended vaccines: 4 DTAP (diphtheria, tetanus, acellular pertussis) 3 IPV (polio) 1 MMR (measles, mumps, rubella) 3 HIB (haemophilus influenza	Children who turned 2 during the measurement year.	Evidence obtained from the medical record counts where the antigen was rendered from one of the following: A note indicating the name of the specific antigen and the date of the immunization A certificate of immunization prepared by an authorized health care provider or agency including the specific dates and types of immunizations administered
		type B) 3 HEP B (hepatitis B) 1 VZV (chicken pox) 4 PCV (pneumococcal conjugate) 1 HEP A (hepatitis A) 2 or 3 RV (rotavirus—2 Rotarix; 3 Rota Teq) 2 Influenza (flu)		If an immunization is not given, documentation for that immunization must include: Allergic reaction to the vaccine or other contraindication History of illness (measles, mumps, rubella, chicken pox, hepatitis A, hepatitis B) Parent refusal does not meet compliance for any vaccines.
Well Child Visits in First 30 Months of Life (W30)	Percentage of children who have had six or more well child visits in their first 15 months of life, and two or more well child visits	First 15: Those in the denominator who had at least 6 well child visits in the last 15 months. Does not include telehealth visits, or sick visits (unless converted to well). The well-child visit must occur with a PCP, but the PCP does not have to be the practitioner assigned to the child.	First 15: Children who turned 15 months old during the measurement year 15-30 months: Children who turned 30 months old during the measurement year	Reporting by age group (first 15 months and 15-30 months) Components of a well child visit: Health history Physical exam Assessment of physical development Assessment of mental development



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	between 15 and			Anticipatory guidance related to
	30 months of life.	15-30 months:		preventive health
		Those in the denominator who had at		·
		least 2 well child visits in the last 15		
		months.		

		Does not include telehealth visits, or		
		sick visits (unless converted to well)		
		The well-child visit must occur with a		
		PCP, but the PCP does not have to be		
		the practitioner assigned to the child.		
Child and	Percentage of	Those in the denominator who had at	Children who are 3 to 21	Reported by age group
Adolescent	patients 3–21	least one well visit with a primary	years of age at the end of	
Well-Care	years of age who	care practitioner or an OB/GYN	the measurement period	Preventive services may be rendered on visits
Visits (WCV)	received one or	practitioner during the measurement		other than WCVs. Well-child preventive services
, ,	more well-care	vear		count toward the measure, regardless of the
	visit with a	, ,		primary intent of the visit, such as OB/GYN visits,
	11010 111011 0	The well-child visit must occur with a		
	primary care			but services that are specific to an acute or
	practitioner or an	PCP or an OB/GYN practitioner, but		chronic condition do not count toward the
	OB/GYN	the practitioner does not have to be		measure.
	practitioner	the practitioner assigned to the child.		
	during the			
	measurement			
	year.			



Behavioral Health

Measure	Description	Numerator	Denominator	Notes
Depression Remission or Response for Adolescents and Adults (DRR)	Percentage of members 12 years of age and older with a diagnosis of depression and an elevated PHQ-9 score, who had evidence of response or remission within 4–8 months of the elevated score.	Follow- Up: Those in the denominator who had a follow-up PHQ-9 score documented within 4–8 months after the initial elevated PHQ-9 score. Remission: Those in the denominator who achieved remission within 4–8 months after the initial elevated PHQ-9 score, as measured by a score of <5 on the most recent PHQ-9 in the response period. Response: Those in the denominator who showed response within 4–8 months after the initial elevated PHQ-9 score, as measured by a 50% reduction in the response period from the initial elevated PHQ-9 score.	People 12 years and older with a diagnosis of depression, and an elevated PHQ-9 score, from May 1 of the year prior to the measurement period through December 31 of the measurement period	
Pharmacotherapy for Opioid Use Disorder (POD)	Percentage of opioid use disorder (OUD) pharmacotherapy treatment events among members age 16 and older that continue for at least 180 days (6 months).	People 16 years of age and older with an active diagnosis of Opioid Use Disorder (OUD), between July 1 of the year prior to the measurement year and June 30 of the measurement year.	Those in the denominator who had pharmacotherapy for at least 180 days, without a gap in treatment of 8 or more consecutive days.	Opioid Use Disorder Treatment Medications include: Naltrexone (oral) Naltrexone (injectable) Buprenorphine (sublingual tablet) Buprenorphine (injection)



		Buprenorphine
		(implant)
		Buprenorphine/
		naloxone (sublingual
		tablet, buccal film,
		sublingual film)
		Methadone (oral)



Pregnant People

Measure	Description	Numerator	Denominator
Prenatal and	The percentage of	Those in the denominator who have a postpartum visit between 7 and 84	Live births or deliveries on or
Postpartum	deliveries that had	days after delivery, with an OB/GYN or other prenatal care clinician or PCP in	between October 8 of the year prior
Care (PPC) -	a postpartum visit	an outpatient setting. Postpartum visit must include at least one of the	to the measurement year and
Postpartum	on or between 7	following:	October 7 of the measurement year.
Care	and 84 days after	Pelvic exam.	Include deliveries that occur in any
	delivery.	Evaluation of breasts, weight, blood pressure and abdomen.	setting.
		Notation of breastfeeding, postpartum care, postpartum care check	
		or six-week check.	
		Preprinted postpartum care form with information from visit	
		documented.	
		Perineal or cesarean incision/wound check.	
		Screening for depression, anxiety, tobacco use, substance use	
		disorder or preexisting mental health disorders.	
		Glucose screening for members with gestational diabetes	
		Documentation of infant care or breastfeeding; resumption of	
		intercourse, birth spacing or family planning; sleep/ fatigue;	
		resumption of physical activity; and attainment of healthy weight.	
Prenatal and	The percentage of	Those in the denominator who had a prenatal visit with an OB/GYN or other	Live birth or deliveries on or between
Postpartum	deliveries that	prenatal care provider, or PCP (with a diagnosis of pregnancy) within the	October 8 of the year prior to the
Care (PPC) -	received a prenatal	required timeframe (first trimester/280–176 days prior to delivery). Prenatal	measurement year and October 7 of
Timeliness of	care visit in the first	visit must include at least one of the following:	the measurement year. Include
Prenatal Care	trimester, on or	When documenting a prenatal visit, include diagnosis of pregnancy,	deliveries that occur in any setting.
	before the	last menstrual period or estimated date of delivery	
	enrollment start	Basic physical obstetrical examination	
	date or within 42	Evidence that a prenatal care procedure was performed.	
	days of enrollment	Include prenatal care such as prenatal risk assessment, complete obstetrical	
	in the organization.	history, fetal heart tone and screening tests.	



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Postpartum	Percentage of	Those in the denominator who had a depression screening result using a	Live birth deliveries on or between
Depression	deliveries in which	standard screening tool for depression, within 7-84 days following delivery:	September 8 of the year prior to the
Screening	patients were	Patient Health Questionnaire (PHQ-9, PHQ9M, PHQ-2)	measurement year and September 7
(PDS-E)	screened for clinical	Beck Depression Inventory (BDI-II) adults only	of the measurement year. Include
	depression using a	Beck Depression Inventory-Fast Screen (BDI-FS)	deliveries that occur in any setting.
	standardized	Center for Epidemiologic Studies Depression Scale-Revised (CESD-R)	
	instrument during	Edinburgh Postnatal Depression Scale (EPDS)	
	the postpartum	PROMIS Depression	
	period and, if	Duke Anxiety-Depression Scale (DUKE-AD) adults only	
	positive, received	Geriatric Depression Scale—Short Form and Long Form (GDS) adults	
	follow-up care	only	
	within 30 days.	My Mood Monitor (M-3) adults only	
		Clinically Useful Depression Outcome Scale (CUDOS) adults only.	
*if PDS-E is	Percentage of	Those in the denominator who had one of the following within 30 days:	All patients from the Postpartum
positive	deliveries in which	Outpatient, telephone or e-visit for follow-up for	Depression Screening Numerator
	patients were	depression/behavioral health	(previous slide) with a positive
Postpartum	screened for clinical	Depression case management encounter	depression result.
Depression	depression using a	Behavioral health encounter (assessment, therapy, collaborative	
Screening	standardized	care, medication management)	
Follow-up	instrument during	A diagnosis of encounter for exercise counseling	
(PDS-E)	the postpartum	Dispensed antidepressant medication	
	period and, if	Documentation of a negative full-length depression screening on the	
	positive, received	same day as a positive screen on a brief screening tool (i.e., a	
	follow-up care	negative PHQ-9 as a follow-up to a positive PHQ-2).	
	within 30 days.		