

Palliative Care Quality Improvement Program (QIP) 2017-18 Measurement Specifications

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Published on: October 6, 2017

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Program Overview

Partnership HealthPlan of California (PHC) has value-based programs in the areas of primary care, hospital care, specialty care, long-term care, community pharmacy, and mental health. These value-based programs align with PHC's organizational mission to help our members and the communities we serve be healthy.

In 2015, Partnership HealthPlan of California (PHC) developed a pilot pre-hospice intensive palliative care program, called *Partners in Palliative Care*. The legislature of California passed a bill (SB 1004) in late 2015, requiring the development of a similar program as a state wide benefit for Medicaid. Implementation of this benefit will likely be on or after January 1, 2018. However, Partnership's board of directors has approved expansion of the pilot program, based on the favorable clinical and financial outcomes of our pilot program.

The expanded pilot implementation began on July 1, 2017, and includes incentives used in the pilot. The incentives in the expanded pilot will be monitored by the PHC Quality Department under the name "Palliative Care Quality Improvement Program (QIP)" consistent with terminology used for other value-based payment programs at PHC.

Participation Requirements

All contracted Intensive Outpatient Palliative Care provider sites participating in the expanded pilot will be automatically enrolled in the Palliative Care QIP, and therefor eligible for the Palliative Care QIP payments. Provider sites must be in good standing with state and federal regulators as of the month the payment is to be disbursed. Good standing means that the provider site is open, solvent, not under financial sanctions from the state of California or Centers for Medicare & Medicaid Services.

Patient Eligibility

Providers may earn incentives from the Palliative Care QIP based on care provided to PHC eligible members, 18 years or older, who have an approved Intensive Outpatient Palliative Care Treatment Authorization Request (TAR) on file. For more information about how members qualify for the program, please contact palliativeQIP@partnershiphp.org for a detailed policy.

Payment Methodology

The incentives provided through the Palliative Care QIP are separate and distinct from a palliative care provider site's usual reimbursement. Each provider site's earning potential is based on its volume of members approved for enrollment in the palliative care program. Please refer to the measure specifications for the incentive amount and payment calculation for each measure.

Program Timeline

The measurement set is approved for the full 2017-18 measurement year, which runs from July 1, 2017 to June 30, 2018. Within the measurement year, there are two six-month measurement periods, at the end of each performance is evaluated. Payment will be disbursed four months after the end of the measurement periods to allow time for payment calculation. This six-month – as opposed to annual – payment schedule is to ensure provider engagement during the transition from the PHC pilot to the implementation of the Medi-Cal benefit. To illustrate, for the measurement year 2017-18, there are two measurement periods:

Measurement Period I: July 1, 2017 – December 31, 2017, with payment date on April 30, 2018 Measurement Period II: January 1, 2018 – June 30, 2018, with payment date on October 31, 2018

Measure I. Avoiding Hospitalization and Emergency Room Visits

Description

The number of members enrolled in the Intensive Outpatient Palliative Care program who did not get admitted to the hospital or emergency department.

One goal of palliative care is to improve quality of life for both the patient and the family. For members who have serious illnesses and are in the palliative care program, we expect the palliative care team to be the first point of contact, which in turn minimizes unnecessary hospitalizations and emergency department visits.

Target

Zero admission or ED visit per member per month.

Measurement Period

Monthly, from July 1, 2017 to June 30, 2018.

Specifications

\$200 per member enrolled in the palliative care program per month only if there are no hospital admissions or ED visits that month.

Hospital admissions and ED visits are identified through data sources including encounters, claims, or treatment authorization requests (TARs) submitted to PHC. Observation stays are included.

Example

For a member who is enrolled in the program on February 25, seen in the emergency room on March 9, admitted from April 23 through April 30, and dies on June 2 at home, the number of months with no hospital encounters or ED visits is 3 (February, May and June). The palliative care provider site will be eligible for a total payment for avoiding hospitalization and ED visits of \$600.

Reporting

Reporting by palliative care provider sites to PHC is not required. PHC will send preliminary reports in the sixth month of the measurement period (i.e. December and June, prior to payment) to help providers monitor performance. Providers can also request member-level reports of admissions and ED visits on an ad hoc basis.

Measure II: Completion of POLST and use of Palliative Care Quality Network (PCQN) Tool

Description

To align best practices, the Palliative Care QIP includes an incentive for 1) completion of the Physician's Orders for Life Sustaining Treatment (POLST) in conjunction with 2) documentation of POLST and patient encounters in the Palliative Care Quality Network System (PCQN).

The POLST was designed for seriously ill patients with the goal of providing a framework for healthcare professionals so they can ensure the patient received the treatments they do want and avoid those treatments that they do not want. The PCQN tool is an online system where palliative care providers share data and from that data can identify possible quality improvement opportunities. This measure will incentivize providers in our program to contribute data, learn about best practices, and capture the key components of care delivery.

Measurement Period

Monthly, from July 1, 2017 to June 30, 2018.

Specifications

\$200 per member enrolled in the palliative care program per month upon completion of a POLST and documentation using the PCQN tool.

At least two entries reflecting patient contact into the PCQN tool must be submitted. Encounters must include minimum data elements exhibited in Appendix II: PCQN Data Elements.

Reporting

Palliative care sites are required to enter data elements in PCQN. Separate reporting by palliative care provider sites to PHC is not required. PHC will send preliminary reports in the sixth month of the measurement period (i.e. December and June, prior to payment) to help providers monitor performance. Providers can also request member-level reports on this measure on an ad hoc basis.

Example

For a member enrolled on February 25, with at least two visits documented on PCQN each month but the POLST completed and entered into PCQN on April 20, the number of months meeting this measure is 3 (April, May, and June). The palliative care provider site will be eligible for a total payment for using PCQN of \$600.

Appendix I: Palliative Care Quality Network Data Elements



	CORE DATASET ITEM	ITEM CHOICES		
	Location / Type of Visit	□ Clinic		
		□ Home		
		□ Telehealth		
		□ SNF / Nursing Home		
S		- Sivi / ivaising nome		
ER	Visit type	□ Initial consult		
별	Visit type	- Initial Consult		
DENTIFIERS	Date of Visit	{YYYY-MM-DD}		
₽	Medical Record Number	(1111 MM 55)		
	Encounter #			
	First Name, Last Name			
	Flist Name, Last Name			
	Gender	- Mala		
	Gender	□ Male		
		□ Female		
	Association of the	□ Unknown		
	Age at time of visit			
	Primary diagnosis leading to PC	□ Cancer (solid tumor)		
	consult	□ Cardiovascular		
		□ Pulmonary		
		□ Vascular		
		☐ Complex chronic conditions/failure to thrive		
		□ Renal		
0		□ Trauma		
Ä		□ Congenital/chromosomal		
<u> </u>		□ Gastrointestinal		
표		□ Hepatic		
₽		□ Hematology		
90		□ Infectious/ immunological/HIV		
Σ		□ In-utero complication/condition		
ä		□ Neurologic/stroke/ neurodegenerative		
<u> </u>		□ Dementia		
TIENT / DEMOGRAPHIC INFO		□ Other		
PAT	Reasons given by referring provider	☐ Goals of care discussion/Advance Care Planning		
_	for initial PC consult (check all)	□ Pain management		
		□ Other symptom management		
		□ Withdrawal of interventions		
		□ Comfort Care		
		☐ Hospice referral/discussion		
		□ No reason given		
		□ Support for patient/family		
		□ Other:		

	Referral Source	□ Ir	npatient Palliative Care		
	CORE DATASET ITEM		ITEM CHOICES		
			□ Other Inpatient Team		
			□ ED		
			□ Primary care		
			☐ Outpatient Palliative Care		
			□ Other Outpatient Specialist		
			□ Self		
			□ Other:		
			□ Unknown		
	Advance directive on chart at the time of		□ Yes		
	consult		□ No		
	POLST on chart at the time of consult		□ Yes		
		_	□ No		
	Palliative Performance Scale (PPS) at time	of	(0% to 100%)		
	consult	. 01	(070 to 10070)		
	ESAS Measures				
	Pain		0-10 scale (77= pt unable)		
	Tiredness		0-10 scale (77= pt unable)		
	Nausea		0-10 scale (77 = pt unable)		
	Depression		0-10 scale (77= pt unable)		
	Anxiety		0-10 scale (77= pt unable)		
	Drowsiness		0-10 scale (77= pt unable)		
	Appetite		0-10 scale (77= pt unable)		
	Well-being		0-10 scale (77= pt unable)		
NS	Short of breath		0-10 scale (77= pt unable)		
'AT	Constipation		0-10 scale (77= pt unable)		
s/status	Composite score		Auto calculated		
T0	Are you at peace?		□ Not at all		
ΜP			□ A little bit		
SYI			□ A moderate amount		
Z			□ Quite a bit		
PATIENT SYMPTOM			□ Completely		
PA			□ Patient unable to rate		
	How much distress have you been	2	0-10 scale (Distress thermometer) (77=pt unable to		
	experiencing the past week including tod	ay?	rate)		
	How would you rate your overall quality		□ Very poor		
	of life?		□ Poor		
			□ Fair		
			□ Good		
			□ Excellent		
			□ Patient unable to rate		

	PC team members involved in visit		Physician
			Clinical Nurse Specialist
ES/			Nurse Practitioner
PRO CES SES/			
CE			
õ			
P			
	CORE DATASET ITEM	ITE	M CHOICES
			Physician Assistant
			Nurse
			Social Worker
			Chaplain
			Pharmacist
			Psychologist/Psychiatrist
		П	Other
	Screening Status		Care
	Pain		Negative
			Positive
	Non-Pain Symptoms		Negative
	, ,		Positive
	Psychosocial needs		Negative
	•		Positive
			Patient/Family declined
			Patient/Family unable to be screened
	Spiritual needs		Negative
	•		Positive
			Patient/Family declined
			Patient/Family unable to be screened
	Advance care planning/Goals of care needs		Negative
			Positive
			Patient/Family declined
			Patient/Family unable to be screened
	Intervention		
	Pain		Yes
	Non-Pain Symptoms		Yes
	Psychosocial needs		Yes
	Spiritual needs		Yes
	Advance care planning/Goals of care needs		Yes
	Other outcomes	1	
	Preference for life-sustaining		Yes
	treatment clarified		
	Advance directive completed		Yes
	POLST completed		Yes
	Preference for life-sustaining		Full code
	treatment		Partial code
			DNR/DNI
			Unknown (default if no code status In system)

Surrogate decision maker identified	 Surrogate decision maker identified and
	documented
	 Attempted to identify but not confirmed
	□ Not addressed
Support for family/caregiver provided	□ Yes
	□ No caregiver present