

NATIONAL QUALITY FORUM

Measure Submission and Evaluation Worksheet 5.0

This form contains the information submitted by measure developers/stewards, organized according to NQF's measure evaluation criteria and process. The evaluation criteria, evaluation guidance documents, and a blank online submission form are available on the [submitting standards web page](#).

NQF #: 0561	NQF Project: Cancer Project
(for Endorsement Maintenance Review)	
Original Endorsement Date: Oct 30, 2009 Most Recent Endorsement Date: Oct 30, 2009 Last Updated Date: Jul 17, 2015	
BRIEF MEASURE INFORMATION	
De.1 Measure Title: Melanoma Coordination of Care	
Co.1.1 Measure Steward: AMA-convened Physician Consortium for Performance Improvement	
De.2 Brief Description of Measure: Percentage of patient visits, regardless of age, seen with a new occurrence of melanoma who have a treatment plan documented in the chart that was communicated to the physician(s) providing continuing care within one month of diagnosis.	
2a1.1 Numerator Statement: Patient visits with a treatment plan documented in the chart that was communicated to the physician(s) providing continuing care within one month of diagnosis	
2a1.4 Denominator Statement: All visits for patients, regardless of age, diagnosed with a new occurrence of melanoma	
2a1.8 Denominator Exclusions: Documentation of patient reason(s) for not communicating treatment plan (eg, patient asks that treatment plan not be communicated physician(s) providing continuing care); Documentation of system reason(s) for not communicating treatment plan to the primary care provider(s) (eg, patient does not have a primary care provider or referring physician)	
1.1 Measure Type: Process	
2a1. 25-26 Data Source: Administrative claims, Electronic Clinical Data, Electronic Clinical Data : Electronic Health Record, Electronic Clinical Data : Registry, Paper Records	
2a1.33 Level of Analysis: Clinician : Group/Practice, Clinician : Individual, Clinician : Team	
1.2-1.4 Is this measure paired with another measure? No	
De.3 If included in a composite, please identify the composite measure (title and NQF number if endorsed): This measure is not included in a composite measure.	

STAFF NOTES (issues or questions regarding any criteria)
Comments on Conditions for Consideration:
Is the measure untested? Yes <input checked="" type="radio"/> No <input checked="" type="radio"/> If untested, explain how it meets criteria for consideration for time-limited endorsement:
1a. Specific national health goal/priority identified by DHHS or NPP addressed by the measure (check De.5):
5. Similar/related endorsed or submitted measures (check 5.1):

Other Criteria:

Staff Reviewer Name(s):

1. IMPACT, OPPORTUNITY, EVIDENCE - IMPORTANCE TO MEASURE AND REPORT

Importance to Measure and Report is a threshold criterion that must be met in order to recommend a measure for endorsement. All three subcriteria must be met to pass this criterion. See [guidance on evidence](#).

Measures must be judged to be important to measure and report in order to be evaluated against the remaining criteria. ([evaluation criteria](#))

1a. High Impact: **H● M● L● I●**

(The measure directly addresses a specific national health goal/priority identified by DHHS or NPP, or some other high impact aspect of healthcare.)

De.4 Subject/Topic Areas (Check all the areas that apply): [Cancer, Cancer : Skin](#)

De.5 Cross Cutting Areas (Check all the areas that apply): [Care Coordination](#)

1a.1 Demonstrated High Impact Aspect of Healthcare: [Affects large numbers, Patient/societal consequences of poor quality](#)

1a.2 If "Other," please describe:

1a.3 Summary of Evidence of High Impact (Provide epidemiologic or resource use data):

[In the year 2010, an estimated 68,130 new cases of melanoma were diagnosed and about 8,700 patients died of the disease in the United States. However, these figures for new cases may represent a substantial underestimation, because many superficial and in situ melanomas treated in the outpatient setting are not reported. The incidence of melanoma continues to increase dramatically. Melanoma is increasing in men more rapidly than any other malignancy and, in women more rapidly than any other malignancy except lung cancer. The lifetime risk of developing melanoma in the year 2005 for someone born in the United States may be as high as one in 55. The median age at diagnosis is 59 years. As such, melanoma ranks second to adult leukemia in terms of loss of years of potential life, per death.\(1\)](#)

[Coordination of care means that the primary care practice must integrate all aspects of care when patients must be seen elsewhere. Because 13 to 20 percent \(depending on various assumptions\) of an average practice population requires a referral each year, this burden is considerable.\(2\)](#)

[Melanoma is among the top 10 new cancer diagnoses for both American men and women. Nationally, melanoma incidence has increased 2.4% annually in the last decade. \(3\)](#)

1a.4 Citations for Evidence of High Impact cited in 1a.3: [1. National Comprehensive Cancer Network \(NCCN\). Clinical Practice Guidelines in Oncology: Melanoma. 2011. Available at: www.nccn.org](#)

[2. Starfield B, Shi L, Macinko J. Contribution of Primary Care to Health Systems and Health. The Milbank Quarterly 2005;83\(3\):457-502.](#)

[3. Hu A, Parmet Y, Allen G, Parker DF, et al. Disparity in Melanoma. A Trend Analysis of Melanoma Incidence and Stage at Diagnosis Among Whites, Hispanics, and Blacks in Florida. Arch Dermatol. 2009;145\(12\):1369-1374.](#)

1b. Opportunity for Improvement: **H● M● L● I●**

(There is a demonstrated performance gap - variability or overall less than optimal performance)

1b.1 Briefly explain the benefits (improvements in quality) envisioned by use of this measure:

The desirable outcome for patients with a new diagnosis or history of melanoma is the prevention of a recurrent melanoma diagnosis and mortality reduction. Requiring that the physician or other healthcare professional who is providing the primary treatment for melanoma communicates in a timely manner to the primary care physician (PCP) ensures better coordination of care, potentially reduces errors, and allows the PCP to encourage appropriate follow up. This utilization of coordination of care across clinicians and settings has been shown to result in greater efficiency and better clinical outcomes.

1. Kohn L, Corrigan J, Donaldson M, eds. Crossing the Quality Chasm: A New Health System for the 21st Century. Washington, DC: Committee on Quality of Health Care in America, Institute of Medicine. National Academy Press; 2001;134.

1b.2 Summary of Data Demonstrating Performance Gap (*Variation or overall less than optimal performance across providers*): [**For Maintenance** – *Descriptive statistics for performance results for this measure - distribution of scores for measured entities by quartile/decile, mean, median, SD, min, max, etc.*]

1. Deficits in communication have clearly been shown to adversely affect post-discharge care transitions. A recent summary of the literature found that direct communication between hospital physicians and primary care physicians occurs infrequently (in 3%-20% of cases studied), the availability of a discharge summary at the first post-discharge visit is low (12%-34%) and did not improve greatly even after 4 weeks (51%-77%), affecting the quality of care in approximately 25% of follow-up visits. This systematic review of the literature also found that discharge summaries often lack important information such as diagnostic test results, treatment or hospital course, discharge medications, test results pending at discharge, patient or family counseling, and follow-up plans.

2. This measure was included in the CMS Physician Quality Reporting Initiative/System (PQRI/S) in 2009 through 2011 in the claims and registry options for 2009 (and registry option only from 2010 and beyond) as PQRI/S #138 (Melanoma Coordination of Care). The number of professionals reporting on this measure in 2009 was approximately 1,028.

The 2009 PQRI/S Performance Rate reveals that there is a gap in care as shown by the following data: 12.72%% of patients reported on did not receive the optimal care.

1b.3 Citations for Data on Performance Gap: [**For Maintenance** – *Description of the data or sample for measure results reported in 1b.2 including number of measured entities; number of patients; dates of data; if a sample, characteristics of the entities included*]

1. Kripalani S, LeFevre F, Phillips CO, Williams MV, Basaviah P, Baker DW. Deficits in Communication and Information Transfer Between Hospital-Based and Primary Care Physicians. Implications for Patient Safety and Continuity of Care. JAMA. 2007;297:831-841.

2. Appendix B. 2009 Physician Quality Reporting System and eRx Experience Report Detailed Tables

1b.4 Summary of Data on Disparities by Population Group: [**For Maintenance** – *Descriptive statistics for performance results for this measure by population group*]

At least two of the reviewed analyses in urban counties showed that the supply of primary care physicians is less closely related to the health of urban African Americans than it is for urban whites or for African Americans in rural areas. This is likely due to the poorer distribution of primary care physicians in more deprived urban areas, with the consequently greater need to seek care in such places as hospital outpatient units and emergency rooms.(1)

Research and public education efforts have focused on melanoma prevention in white populations because of their higher risk of developing melanoma. Improved secondary prevention measures with earlier detection of thin (early-stage) melanoma likely account for the improved survival among whites from 68% in the early 1970s to 92% in recent years. Such advances, however, have not occurred in other racial and

ethnic groups in the United States. Emerging data call attention to disparity in melanoma diagnosis and survival in minorities such as Hispanics and blacks. Multiple reports found that US blacks have more advanced melanoma in association with worse survival rates; however, melanoma disparity among Hispanics is less recognized. The dearth of studies on melanoma among Hispanics partly reflects the small number of cases in many areas of the United States, as well as limitations of ethnicity information in cancer registries. In fact, the National Cancer Institute's Surveillance, Epidemiology, and End Results (SEER) program and most other cancer registries did not begin classifying data for "Hispanic" until the late 1990s. As a result, few studies included data regarding Hispanics.

1b.5 Citations for Data on Disparities Cited in 1b.4: *[For Maintenance – Description of the data or sample for measure results reported in 1b.4 including number of measured entities; number of patients; dates of data; if a sample, characteristics of the entities included]*

1. Starfield B, Shi L, Macinko J. Contribution of Primary Care to Health Systems and Health. The Milbank Quarterly 2005;83(3):457-502.

2. Hu A, Parmet Y, Allen G, Parker DF, et al. Disparity in Melanoma. A Trend Analysis of Melanoma Incidence and Stage at Diagnosis Among Whites, Hispanics, and Blacks in Florida. Arch Dermatol. 2009;145(12):1369-1374.

1c. Evidence (Measure focus is a health outcome OR meets the criteria for quantity, quality, consistency of the body of evidence.)

Is the measure focus a health outcome? Yes ☐ No ☐ If not a health outcome, rate the body of evidence.

Quantity: H ☐ M ☐ L ☐ I ☐ Quality: H ☐ M ☐ L ☐ I ☐ Consistency: H ☐ M ☐ L ☐ I ☐

Quantity	Quality	Consistency	Does the measure pass subcriterion 1c?
M-H	M-H	M-H	Yes <input type="radio"/>
L	M-H	M	Yes <input type="radio"/> IF additional research unlikely to change conclusion that benefits to patients outweigh harms: otherwise No <input type="radio"/>
M-H	L	M-H	Yes <input type="radio"/> IF potential benefits to patients clearly outweigh potential harms: otherwise No <input type="radio"/>
L-M-H	L-M-H	L	No <input type="radio"/>

Health outcome – rationale supports relationship to at least one healthcare structure, process, intervention, or service

Does the measure pass subcriterion 1c?
Yes ☐ IF rationale supports relationship

1c.1 Structure-Process-Outcome Relationship (Briefly state the measure focus, e.g., health outcome, intermediate clinical outcome, process, structure; then identify the appropriate links, e.g., structure-process-health outcome; process-health outcome; intermediate clinical outcome-health outcome):

This is a process measure, which encourages communication, within one month of diagnosis, to the physician providing continuing care to patients with a new occurrence of melanoma. Communication between physicians within a timely manner will lead to improved outcomes, by closing the loop of continuous care, thereby reducing morbidity and mortality rates due to delays in treatment and/or follow-up care.

1c.2-3 Type of Evidence (Check all that apply):

Clinical Practice Guideline, Systematic review of body of evidence (other than within guideline development)

1c.4 Directness of Evidence to the Specified Measure (State the central topic, population, and outcomes)

See Guidance for Definitions of Rating Scale: H=High; M=Moderate; L=Low; I=Insufficient; NA=Not Applicable
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addressed in the body of evidence and identify any differences from the measure focus and measure target population):

The evidence focuses on the need for more frequent, timely, and inclusive communication between physicians during a patient's transition. The evidence also outlines important information that may be included in physician communications. Therefore, the evidence is aligned with the specified measure.

1c.5 Quantity of Studies in the Body of Evidence *(Total number of studies, not articles):* The systematic review of the literature, cited in the policy statement, states the following:

Of the 1064 citations identified through the initial electronic search and screened for possible inclusion, 170 publications were judged to warrant full review. A hand search of references from relevant articles yielded an additional 43 articles for review, and the updated literature search identified 1 more intervention. After exclusion of 2 duplicate or similar publications a total of 73 studies met inclusion criteria, including 55 observational studies (21 medical record audits, 23 physician surveys, and 11 combined audit-surveys) and 18 trials of controlled interventions (3 randomized, 7 nonrandomized with concurrent control, and 8 with pre-post design).

Kripalani S, LeFevre F, Phillips CO, Williams MV, Basaviah P, Baker DW. Deficits in Communication and Information Transfer Between Hospital-Based and Primary Care Physicians. Implications for Patient Safety and Continuity of Care. JAMA. 2007;297:831-841.

1c.6 Quality of Body of Evidence *(Summarize the certainty or confidence in the estimates of benefits and harms to patients across studies in the body of evidence resulting from study factors. Please address: a) study design/flaws; b) directness/indirectness of the evidence to this measure (e.g., interventions, comparisons, outcomes assessed, population included in the evidence); and c) imprecision/wide confidence intervals due to few patients or events):* The systematic review of the literature, cited in the policy statement, states the following:

The primary limitations of this review relate to the consistency and quality of this body of evidence. First, interpretation and synthesis of the findings are restricted by the high degree of variability among studies in their patient populations, outcome measures, and types of interventions tested. Second, interventions to improve the quality of discharge summaries were more difficult to interpret and synthesize because the available trials contained limited outcome data and used different metrics for their outcomes, some of which had unclear clinical significance. Third, there was a relative lack of high-quality investigations, with only 3 randomized controlled trials identified. The nonrandomized studies are subject to numerous biases, including the possible noncomparability of groups at baseline, and pre-post studies may also be affected by the confounding effects of secular time trends. Many studies did not include an appropriate analysis of outcomes, lacking statistical testing for example, and few investigations made attempts to measure and control for potential confounding variables.

Kripalani S, LeFevre F, Phillips CO, Williams MV, Basaviah P, Baker DW. Deficits in Communication and Information Transfer Between Hospital-Based and Primary Care Physicians. Implications for Patient Safety and Continuity of Care. JAMA. 2007;297:831-841.

1c.7 Consistency of Results across Studies *(Summarize the consistency of the magnitude and direction of the effect):* The systematic review of the literature, cited in the policy statement, states the following:

The primary limitations of this review relate to the consistency and quality of this body of evidence.

The generalizability of these results is uncertain. Most of the included studies were conducted outside the United States in countries with a single-payer or national health system. There may be substantial differences in the feasibility and impact of interventions if implemented in a heterogeneous medical system

as found in the United States. Even within a single medical system, differences in logistics and efficiency across hospitals may create unique challenges and opportunities that need to be addressed at the local level.

Kripalani S, LeFevre F, Phillips CO, Williams MV, Basaviah P, Baker DW. Deficits in Communication and Information Transfer Between Hospital-Based and Primary Care Physicians. Implications for Patient Safety and Continuity of Care. JAMA. 2007;297:831-841.

1c.8 Net Benefit (*Provide estimates of effect for benefit/outcome; identify harms addressed and estimates of effect; and net benefit - benefit over harms*):

Communication between physicians within a timely manner will close the loop of continuous care, thereby reducing morbidity and mortality rates due to delays in treatment and/or follow-up care. No harms have been identified.

1c.9 Grading of Strength/Quality of the Body of Evidence. Has the body of evidence been graded? **No**

1c.10 If body of evidence graded, identify the entity that graded the evidence including balance of representation and any disclosures regarding bias: **Not applicable**

1c.11 System Used for Grading the Body of Evidence: **Other**

1c.12 If other, identify and describe the grading scale with definitions: **Not applicable**

1c.13 Grade Assigned to the Body of Evidence: **Not applicable**

1c.14 Summary of Controversy/Contradictory Evidence: **No controversial or contradictory evidence has been identified.**

1c.15 Citations for Evidence other than Guidelines(Guidelines addressed below):

Ensure that care information is transmitted and appropriately documented in a timely manner and in a clearly understandable form to patients and to all of the patient's health care providers/professionals, within and between care settings, who need that information to provide continued care.

National Quality Forum. Safe Practices for Better Healthcare 2010 Update.
www.qualityforum.org/About_NQF/CSAC/Safe_Practices_Table.aspx

1c.16 Quote verbatim, the specific guideline recommendation (*Including guideline # and/or page #*):

Communication and information exchange between the medical home and the receiving provider should occur in an amount of time that will allow the receiving provider to effectively treat the patient. This communication and information exchange should ideally occur whenever patients are at a transition of care; e.g., at discharge from the inpatient setting. The timeliness of this communication should be consistent with the patient's clinical presentation and, in the case of a patient being discharged, the urgency of the follow-up required.

Communication and information exchange between the MD and other physicians may be in the form of a call, voicemail, fax or other secure, private, and accessible means including mutual access to an EHR.

The TOCCC proposed a minimal set of data elements that should always be part of the transition record and be part of any initial implementation of this standard. That list includes the following:

- Principle diagnosis and problem list
- Medication list (reconciliation) including over the counter/ herbals, allergies and drug interactions
- Clearly identifies the medical home/transferring coordinating physician/institution and their contact information

- Patient's cognitive status
- Test results/pending results

The TOCCC recommended the following additional elements that should be included in an "ideal transition record" in addition to the above:

- Emergency plan and contact number and person
- Treatment and diagnostic plan
- Prognosis and goals of care
- Advance directives, power of attorney, consent
- Planned interventions, durable medical equipment, wound care etc
- Assessment of caregiver status
- Patients and/or their family/caregivers must receive, understand and be encouraged to participate in the development of their transition record which should take into consideration the patient's health literacy, insurance status and be culturally sensitive.

1c.17 Clinical Practice Guideline Citation: Snow V, Beck D, Budnitz T, Miller DC, Potter J, Wears RL, Weiss KB, Williams MV. Transitions of Care Consensus Policy Statement: American College of Physicians-Society of General Internal Medicine-Society of Hospital Medicine-American Geriatrics Society-American College of Emergency Physicians-Society of Academic Emergency Medicine. J Gen Intern Med 2009 Apr 3.

1c.18 National Guideline Clearinghouse or other URL:
<http://psnet.ahrq.gov/resource.aspx?resourceID=9929>

1c.19 Grading of Strength of Guideline Recommendation. Has the recommendation been graded? **No**

1c.20 If guideline recommendation graded, identify the entity that graded the evidence including balance of representation and any disclosures regarding bias:

1c.21 System Used for Grading the Strength of Guideline Recommendation: **Other**

1c.22 If other, identify and describe the grading scale with definitions: **N/A**

1c.23 Grade Assigned to the Recommendation: **N/A**

1c.24 Rationale for Using this Guideline Over Others: It is the PCPI policy to use guidelines, which are evidence-based, applicable to physicians and other health-care providers, and developed by a national specialty organization or government agency. In addition, the PCPI has now expanded what is acceptable as the evidence base for measures to include documented quality improvement (QI) initiatives or implementation projects that have demonstrated improvement in quality of care.

Based on the NQF descriptions for rating the evidence, what was the developer's assessment of the quantity, quality, and consistency of the body of evidence?

1c.25 Quantity: **Moderate** **1c.26 Quality:** **Moderate** **1c.27 Consistency:** **Moderate**

1c.28 Attach evidence submission form:

1c.29 Attach appendix for supplemental materials:

Was the threshold criterion, *Importance to Measure and Report*, met?

(1a & 1b must be rated moderate or high and 1c yes) Yes ☐ No ☒

Provide rationale based on specific subcriteria:

For a new measure if the Committee votes NO, then STOP.

For a measure undergoing endorsement maintenance, if the Committee votes NO because of 1b. (no opportunity for improvement), it may be considered for continued endorsement and all criteria need to be evaluated.

2. RELIABILITY & VALIDITY - SCIENTIFIC ACCEPTABILITY OF MEASURE PROPERTIES

Extent to which the measure, as specified, produces consistent (reliable) and credible (valid) results about the quality of care when implemented. (**evaluation criteria**)

Measure testing must demonstrate adequate reliability and validity in order to be recommended for endorsement. Testing may be conducted for data elements and/or the computed measure score. Testing information and results should be entered in the appropriate field. Supplemental materials may be referenced or attached in item 2.1. See [guidance on measure testing](#).

S.1 Measure Web Page (*In the future, NQF will require measure stewards to provide a URL link to a web page where current detailed specifications can be obtained*). Do you have a web page where current detailed specifications for this measure can be obtained? **Yes**

S.2 If yes, provide web page URL: www.physicianconsortium.org

2a. RELIABILITY. Precise Specifications and Reliability Testing: H● M● L● I●

2a1. Precise Measure Specifications. (*The measure specifications precise and unambiguous.*)

2a1.1 Numerator Statement (*Brief, narrative description of the measure focus or what is being measured about the target population, e.g., cases from the target population with the target process, condition, event, or outcome*):

[Patient visits with a treatment plan documented in the chart that was communicated to the physician\(s\) providing continuing care within one month of diagnosis](#)

2a1.2 Numerator Time Window (*The time period in which the target process, condition, event, or outcome is eligible for inclusion*):

[1 month after each visit within measurement period](#)

2a1.3 Numerator Details (*All information required to identify and calculate the cases from the target population with the target process, condition, event, or outcome such as definitions, codes with descriptors, and/or specific data collection items/responses*):

Numerator Definition:

[Communication: may include documentation in the medical record that the physician treating the melanoma communicated \(eg, verbally, by letter, copy of treatment plan sent\) with the physician\(s\) providing the continuing care OR a copy of a letter in the medical record outlining whether the patient was or should be treated for melanoma.](#)

[Treatment plan: for the purposes of this measure, should include the following elements: diagnosis, tumor thickness, and plan for surgery or alternate care.](#)

For EHR:

[eSpecification and eMeasure are currently under development \(expected completion end of Q1 2012\). See attached data elements \(using Quality Data Model\) required for the measure.](#)

For Claims/Administrative:

Report CPT Category II Code:

[5050F - Treatment plan communicated to provider\(s\) managing continuing care within one month of diagnosis](#)

2a1.4 Denominator Statement (*Brief, narrative description of the target population being measured*):
[All visits for patients, regardless of age, diagnosed with a new occurrence of melanoma](#)

2a1.5 Target Population Category (*Check all the populations for which the measure is specified and tested if any*): [Senior Care](#)

2a1.6 Denominator Time Window (*The time period in which cases are eligible for inclusion*):
12 consecutive months

2a1.7 Denominator Details (*All information required to identify and calculate the target population/denominator such as definitions, codes with descriptors, and/or specific data collection items/responses*):

For Claims/Administrative:

Option 1

ICD-9-CM diagnosis codes: 172.0, 172.1, 172.2, 172.3, 172.4, 172.5, 172.6, 172.7, 172.8, 172.9

AND

CPT codes for excision of malignant melanoma: 11600, 11601, 11602, 11603, 11604, 11606, 11620, 11621, 11622, 11623, 11624, 11626, 11640, 11641, 11642, 11643, 11644, 11646, 14000, 14001, 14020, 14021, 14040, 14041, 14060, 14061, 14301, 14302, 17311, 17313

OR

Option 2

ICD-9-CM diagnosis codes: 172.0, 172.1, 172.2, 172.3, 172.4, 172.5, 172.6, 172.7, 172.8, 172.9

AND

CPT codes: 99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99241, 99242, 99243, 99244, 99245

For EHR:

eSpecification and eMeasure are currently under development (expected completion end of Q1 2012). See attached data elements (using Quality Data Model) required for the measure.

2a1.8 Denominator Exclusions (*Brief narrative description of exclusions from the target population*):

Documentation of patient reason(s) for not communicating treatment plan (eg, patient asks that treatment plan not be communicated physician(s) providing continuing care);

Documentation of system reason(s) for not communicating treatment plan to the primary care provider(s) (eg, patient does not have a primary care provider or referring physician)

2a1.9 Denominator Exclusion Details (*All information required to identify and calculate exclusions from the denominator such as definitions, codes with descriptors, and/or specific data collection items/responses*):

The PCPI methodology uses three categories of reasons for which a patient may be excluded from the denominator of an individual measure. These measure exception categories are not uniformly relevant across all measures; for each measure, there must be a clear rationale to permit an exception for a medical, patient, or system reason. Examples are provided in the measure exception language of instances that may constitute an exception and are intended to serve as a guide to clinicians. For this measure: patient reason(s) (eg, patient asks that treatment plan not be communicated to the physician(s) providing continuing care) or system reason(s) (eg, patient does not have a primary care physician or referring physician)]. Where examples of exceptions are included in the measure language, these examples are coded and included in the eSpecifications. Although this methodology does not require the external reporting of more detailed exception data, the PCPI recommends that physicians document the specific reasons for exception in patients' medical records for purposes of optimal patient management and audit-readiness. The PCPI also advocates the systematic review and analysis of each physician's exceptions data to identify practice patterns and opportunities for quality improvement. For example, it is possible for implementers to calculate the percentage of patients that physicians have identified as meeting the criteria for exception. Additional details by data source are as follows:

For EHR:

eSpecification and eMeasure are currently under development (expected completion end of Q1 2012). See attached data elements (using Quality Data Model) required for the measure.

For Claims/Administrative:

Documentation of patient reason(s) for not communicating the treatment plan

- Append modifier to CPT Category II code: 5050F-2P

Documentation of system reason(s) for not communicating the treatment plan

- Append modifier to CPT Category II code: 5050F-3P

2a1.10 Stratification Details/Variables *(All information required to stratify the measure results including the stratification variables, codes with descriptors, definitions, and/or specific data collection items/responses):*

We encourage the results of this measure to be stratified by race, ethnicity, gender, and primary language, and have included these variables as recommended data elements to be collected.

2a1.11 Risk Adjustment Type *(Select type. Provide specifications for risk stratification in 2a1.10 and for statistical model in 2a1.13):* No risk adjustment or risk stratification **2a1.12 If "Other," please describe:**

2a1.13 Statistical Risk Model and Variables *(Name the statistical method - e.g., logistic regression and list all the risk factor variables. Note - risk model development should be addressed in 2b4.):*

No risk adjustment or risk stratification.

2a1.14-16 Detailed Risk Model Available at Web page URL (or attachment). Include coefficients, equations, codes with descriptors, definitions, and/or specific data collection items/responses. Attach documents only if they are not available on a webpage and keep attached file to 5 MB or less. NQF strongly prefers you make documents available at a Web page URL. Please supply login/password if needed:

2a1.17-18. Type of Score: Rate/proportion

2a1.19 Interpretation of Score *(Classifies interpretation of score according to whether better quality is associated with a higher score, a lower score, a score falling within a defined interval, or a passing score):*
Better quality = Higher score

2a1.20 Calculation Algorithm/Measure Logic *(Describe the calculation of the measure score as an ordered sequence of steps including identifying the target population; exclusions; cases meeting the target process, condition, event, or outcome; aggregating data; risk adjustment; etc.):*

To calculate performance rates:

- 1) Find the patients who meet the initial patient population (ie, the general group of patients that the performance measure is designed to address).
- 2) From the patients within the initial patient population criteria, find the patients who qualify for the denominator (ie, the specific group of patients for inclusion in a specific performance measure based on defined criteria). Note: in some cases the initial patient population and denominator are identical.
- 3) From the patients within the denominator, find the patients who qualify for the Numerator (ie, the group of patients in the denominator for whom a process or outcome of care occurs). Validate that the number of patients in the numerator is less than or equal to the number of patients in the denominator
- 4) From the patients who did not meet the numerator criteria, determine if the physician has documented that the patient meets any criteria for denominator exception when exceptions have been

specified [for this measure: patient reason(s) (eg, patient asks that treatment plan not be communicated to the physician(s) providing continuing care) or system reason(s) (eg, patient does not have a primary care physician or referring physician)]. If the patient meets any exception criteria, they should be removed from the denominator for performance calculation. --Although the exception cases are removed from the denominator population for the performance calculation, the number of patients with valid exceptions should be calculated and reported along with performance rates to track variations in care and highlight possible areas of focus for QI.

2a1.21-23 Calculation Algorithm/Measure Logic Diagram URL or attachment:

Attachment

[AMA-PCPI_Measure Calculation-Standard Measures.pdf](#)

2a1.24 Sampling (Survey) Methodology. If measure is based on a sample (or survey), provide instructions for obtaining the sample, conducting the survey and guidance on minimum sample size (response rate):

Not applicable. This measure does not require sampling or a survey.

2a1.25 Data Source (*Check all the sources for which the measure is specified and tested*). If other, please describe:

Administrative claims, Electronic Clinical Data, Electronic Clinical Data : Electronic Health Record, Electronic Clinical Data : Registry, Paper Records

2a1.26 Data Source/Data Collection Instrument (*Identify the specific data source/data collection instrument, e.g. name of database, clinical registry, collection instrument, etc.*): Not Applicable

2a1.27-29 Data Source/data Collection Instrument Reference Web Page URL or Attachment:

2a1.30-32 Data Dictionary/Code Table Web Page URL or Attachment:

Attachment

[AMA-PCPI_0561_MELANOMA.CoordCare_DATAELEMENTS.pdf](#)

2a1.33 Level of Analysis (*Check the levels of analysis for which the measure is specified and tested*):

Clinician : Group/Practice, Clinician : Individual, Clinician : Team

2a1.34-35 Care Setting (*Check all the settings for which the measure is specified and tested*): Ambulatory Care : Clinician Office

2a2. Reliability Testing. (*Reliability testing was conducted with appropriate method, scope, and adequate demonstration of reliability.*)

2a2.1 Data/Sample (*Description of the data or sample including number of measured entities; number of patients; dates of data; if a sample, characteristics of the entities included*):

AAD Testing Project

3 dermatology practice sites representing various types, locations and sizes were identified to participate in testing the melanoma measures.

- Site A was a dermatology practice with 3 physicians that cared for mainly geriatric patients.
- Site B was a general dermatology practice with 9 physicians that cared for a pediatric to geriatric patient population.
- Site C was a medical dermatology practice with 3 physicians that cared for a pediatric to geriatric patient population.

- Site A was paper chart-based. Sites B and C utilized EHRs.
- Sites were located in the northeastern and southern regions of the United States.
- Patient visit volume was 125 visits per day at site A and 115-130 visits per day at site C. For site B, physician 1 had 40 patient visits per day, physician 2 had 38 patient visits per day and patient 3 had 57 patient visits per day.
- All sites were instructed to select the number of charts that they had entered into the American Academy of Dermatology (AAD) 2011 PQRS Registry. For this measure, abstractors reviewed 476 patient visits.
- The measurement period (data collected from patients seen) was January 2011 through July 2011.
- Chart auditing was performed mid-July 2011 through mid-August 2011 (1 month total).

2a2.2 Analytic Method *(Describe method of reliability testing & rationale):*

AAD Testing Project

Data abstracted from patient records were used to calculate inter-rater reliability for the measure.

Data analysis at the data element level included:

- Percent agreement; and
- Kappa statistic to adjust for chance agreement.

2a2.3 Testing Results *(Reliability statistics, assessment of adequacy in the context of norms for the test conducted):*

Reliability: N, % Agreement, Kappa (95% Confidence Interval)

Melanoma diagnosis

Overall: 466, 95.92%, 0.8603 (0.7988-0.9218)

Denominator: 466, 97.00%, 0.4012 (0.0923-0.7101)

Numerator: 466, 95.92%, 0.8603 (0.7988-0.9218)

Exceptions: 466, 100%, kappa statistic is noncalculable*

This measure demonstrates moderate reliability, as shown in results from the above analysis.

Follow-up with other physician as part of treatment plan

Overall: 79, 69.62%, 0.3291 (0.1051-0.5531)

Denominator: 79, 97.47%, kappa statistic is noncalculable*

Numerator: 79, 69.62%, 0.3291 (0.1051-0.5531)

Exceptions: 466, 100%, kappa statistic is noncalculable*

This measure demonstrates fair reliability, as shown in results from the above analysis.

Follow-up with other physician sent as part of treatment plan

Overall: 40, 72.50%, 0.1129 (0.000-0.5593)

Denominator: 40, 100%, kappa statistic is noncalculable*

Numerator: 40, 72.50%, 0.1129 (0.000-0.5593)

Exceptions: 40, 100%, kappa statistic is noncalculable*

This measure demonstrates fair reliability, as shown in results from the above analysis.

Follow-up scheduled in recall system as part of treatment plan

Overall: 465, 93.55%, 0.6606 (0.5431-0.7781)

Denominator: 465, 100%, kappa statistic is noncalculable*

Numerator: 465, 93.55%, 0.6606 (0.5431-0.7781)

Exceptions: 465, 100%, kappa statistic is noncalculable*

This measure demonstrates almost perfect reliability, as shown in results from the above analysis.

Follow-up recall information as part of treatment plan
 Overall: 402, 89.05%, kappa statistic is noncalculable*
 Denominator: 402, 100%, kappa statistic is noncalculable*
 Numerator: 402, 89.05%, kappa statistic is noncalculable*
 Exceptions: 402, 100%, kappa statistic is noncalculable*
 This measure demonstrates almost perfect reliability, as shown in results from the above analysis.

Follow-up self exam as part of treatment plan
 Overall: 273, 81.32%, 0.6225 (0.5291-0.7159)
 Denominator: 273, 100%, kappa statistic is noncalculable*
 Numerator: 273, 81.32%, 0.6225 (0.5291-0.7159)
 Exceptions: 273, 100%, kappa statistic is noncalculable*
 This measure demonstrates almost perfect reliability, as shown in results from the above analysis.

Follow-up treatment plan documented
 Overall: 79, 92.41%, 0.000 (0.000-0.7692)
 Denominator: 79, 100%, kappa statistic is noncalculable*
 Numerator: 79, 92.41%, 0.000 (0.000-0.7692)
 Exceptions: 79, 100%, kappa statistic is noncalculable*
 This measure demonstrates almost perfect reliability, as shown in results from the above analysis.

*Kappa Statistics cannot be calculated because of complete agreement. Confidence intervals cannot be calculated because to do so would involve dividing by zero which cannot be done.

2b. VALIDITY. Validity, Testing, including all Threats to Validity: H M L I

2b1.1 Describe how the measure specifications (measure focus, target population, and exclusions) are consistent with the evidence cited in support of the measure focus (criterion 1c) and identify any differences from the evidence:

The policy statement focuses on the communication between physicians. The measure specifications are consistent with the evidence cited in support of the measure focus, as the measure captures visits of all patients, regardless of age, with a new occurrence of melanoma.

2b2. Validity Testing. (Validity testing was conducted with appropriate method, scope, and adequate demonstration of validity.)

2b2.1 Data/Sample (Description of the data or sample including number of measured entities; number of patients; dates of data; if a sample, characteristics of the entities included):

An expert panel was used to assess face validity of the measure. This panel consisted of the following 16 members, with representation from the following specialties:

List of Work Group Members

Raj Behal, MD, MPH (Co-Chair)(methodology)
 Dirk Elston, MD (Co-Chair)(dermatology)
 Stephen Bines, MD (general surgery)
 Peter C. Dandalides, MD (health plan)
 Evan Farmer, MD (dermatology)
 Rutledge Forney, MD (dermatology)
 Andrea Gelzer, MD, MS FACP (health plan)
 Robert Gilson, MD (dermatology)
 Stephen Helms, MD (dermatology)
 Abrar Qureshi, MD (dermatology)

Todd Schlessinger, MD (dermatology)
John Schneider, MD, PhD (family medicine)
Janet (Jessie) Sullivan, MD(dermatology)
Arthur Sober, MD (dermatology)
Steven Strode, MD, Med, MPH (family medicine)
William Wooden, MD (plastic surgery)

2b2.2 Analytic Method *(Describe method of validity testing and rationale; if face validity, describe systematic assessment):*

All PCPI performance measures are assessed for content validity by a panel of expert work group members during the development process. Additional input on the content validity of draft measures is obtained through a 30-day public comment period and by also soliciting comments from a panel of consumer, purchaser, and patient representatives convened by the PCPI specifically for this purpose. All comments received are reviewed by the expert work group and the measures adjusted as needed. Other external review groups (eg, focus groups) may be convened if there are any remaining concerns related to the content validity of the measures.

The expert panel was used to assess face validity of the measure. This panel consisted of 16 members, with representation from the following specialties: general surgery, dermatology, family medicine, plastic surgery, and health plan representatives.

The aforementioned panel was asked to rate their agreement with the following statement:

The scores obtained from the measure as specified will accurately differentiate quality across providers.

Scale 1-5, where 1=Strongly Disagree; 3=Neither Disagree nor Agree; 5=Strongly Agree

2b2.3 Testing Results *(Statistical results, assessment of adequacy in the context of norms for the test conducted; if face validity, describe results of systematic assessment):*

The results of the expert panel rating of the validity statement were as follows: N = 11; Mean rating = 4.5.

Percentage in the top two categories (4 and 5): 100%

Frequency Distribution of Ratings

1 – 0
2 – 0
3 – 0
4 – 5
5 – 6

POTENTIAL THREATS TO VALIDITY. *(All potential threats to validity were appropriately tested with adequate results.)*

2b3. Measure Exclusions. *(Exclusions were supported by the clinical evidence in 1c or appropriately tested with results demonstrating the need to specify them.)*

2b3.1 Data/Sample for analysis of exclusions *(Description of the data or sample including number of measured entities; number of patients; dates of data; if a sample, characteristics of the entities included):*
PCPI Testing Project

- o All sites were instructed to select the number of charts that they had entered into the American Academy of Dermatology (AAD) 2011 PQRS Registry. For this measure, abstractors reviewed 476 patient visits.
- o The measurement period (data collected from patients seen) was January 2011 through July 2011
- o Chart auditing was performed mid-July 2011 through mid-August 2011 (1 month total)

2b3.2 Analytic Method *(Describe type of analysis and rationale for examining exclusions, including exclusion related to patient preference):*

Exceptions were analyzed for frequency and variability across providers.

2b3.3 Results *(Provide statistical results for analysis of exclusions, e.g., frequency, variability, sensitivity analyses):*

PCPI Testing Project

Reliability: N, % Agreement, Kappa (95% Confidence Interval)

Melanoma diagnosis

Exceptions: 466, 100%, kappa statistic is noncalculable*

This measure demonstrates perfect reliability, as shown in results from the above analysis.

Follow-up with other physician as part of treatment plan

Exceptions: 466, 100%, kappa statistic is noncalculable*

This measure demonstrates perfect reliability, as shown in results from the above analysis.

Follow-up with other physician sent as part of treatment plan

Exceptions: 40, 100%, kappa statistic is noncalculable*

This measure demonstrates perfect reliability, as shown in results from the above analysis.

Follow-up scheduled in recall system as part of treatment plan

Exceptions: 465, 100%, kappa statistic is noncalculable*

This measure demonstrates perfect reliability, as shown in results from the above analysis.

Follow-up recall information as part of treatment plan

Exceptions: 402, 100%, kappa statistic is noncalculable*

This measure demonstrates perfect reliability, as shown in results from the above analysis.

Follow-up self exam as part of treatment plan

Exceptions: 273, 100%, kappa statistic is noncalculable*

This measure demonstrates perfect reliability, as shown in results from the above analysis.

Follow-up treatment plan documented

Exceptions: 79, 100%, kappa statistic is noncalculable*

This measure demonstrates perfect reliability, as shown in results from the above analysis.

* Kappa statistics cannot be calculated because of complete agreement. Confidence intervals cannot be calculated because to do so would involve dividing by zero which cannot be done.

2b4. Risk Adjustment Strategy. *(For outcome measures, adjustment for differences in case mix (severity) across measured entities was appropriately tested with adequate results.)*

2b4.1 Data/Sample *(Description of the data or sample including number of measured entities; number of patients; dates of data; if a sample, characteristics of the entities included):*

This measure is not risk adjusted.

2b4.2 Analytic Method *(Describe methods and rationale for development and testing of risk model or risk stratification including selection of factors/variables):*

This measure is not risk adjusted.

2b4.3 Testing Results *(Statistical risk model: Provide quantitative assessment of relative contribution of*

model risk factors; risk model performance metrics including cross-validation discrimination and calibration statistics, calibration curve and risk decile plot, and assessment of adequacy in the context of norms for risk models. Risk stratification: Provide quantitative assessment of relationship of risk factors to the outcome and differences in outcomes among the strata):

Not applicable

2b4.4 If outcome or resource use measure is not risk adjusted, provide rationale and analyses to justify lack of adjustment: As a process measure, no risk adjustment is necessary.

2b5. Identification of Meaningful Differences in Performance. (The performance measure scores were appropriately analyzed and discriminated meaningful differences in quality.)

2b5.1 Data/Sample (Describe the data or sample including number of measured entities; number of patients; dates of data; if a sample, characteristics of the entities included):

AAD Testing Project

o All sites were instructed to select the number of charts that they had entered into the American Academy of Dermatology (AAD) 2011 PQRS Registry. For this measure, abstractors reviewed 476 patient visits.

o The measurement period (data collected from patients seen) was January 2011 through July 2011

o Chart auditing was performed mid-July 2011 through mid-August 2011 (1 month total)

2b5.2 Analytic Method (Describe methods and rationale to identify statistically significant and practically/meaningfully differences in performance):

AAD Testing Project

The variability in physician performance rates across the measure were calculated by decile

2b5.3 Results (Provide measure performance results/scores, e.g., distribution by quartile, mean, median, SD, etc.; identification of statistically significant and meaningfully differences in performance):

AAD Testing Project

N is the number of physicians with 10 or more quality events

Measure 138

N = 48

Performance Rate	Decile
0.8576	0.1
0.9786	0.2
1.0000	0.3
1.0000	0.4
1.0000	0.5
1.0000	0.6
1.0000	0.7
1.0000	0.8
1.0000	0.9
1.0000	1

2b6. Comparability of Multiple Data Sources/Methods. (If specified for more than one data source, the various approaches result in comparable scores.)

2b6.1 Data/Sample (Describe the data or sample including number of measured entities; number of patients; dates of data; if a sample, characteristics of the entities included):

PCPI Testing Project

o All sites were instructed to select the number of charts that they had entered into the American Academy of Dermatology (AAD) 2011 PQRS Registry. For this measure, abstractors reviewed 476 patient visits.

o The measurement period (data collected from patients seen) was January 2011 through July 2011

- o Chart auditing was performed mid-July 2011 through mid-August 2011 (1 month total)

2b6.2 Analytic Method *(Describe methods and rationale for testing comparability of scores produced by the different data sources specified in the measure):*

Reliability was measured as the ratio of signal to noise. The signal in this case is the proportion of the variability in measured performance that can be explained by real differences in physician performance. Reliability at the level of the specific physician is given by:

Reliability = Variance (physician-to-physician) / [Variance (physician-to-physician) + Variance (physician-specific-error)]

Reliability is the ratio of the physician-to-physician variance divided by the sum of the physician-to-physician variance plus the error variance specific to a physician. A reliability of zero implies that all the variability in a measure is attributable to measurement error. A reliability of one implies that all the variability is attributable to real differences in physician performance.

Reliability testing was performed by using a beta-binomial model. The beta-binomial model assumes the physician performance score is a binomial random variable conditional on the physician's true value that comes from the beta distribution. The beta distribution is usually defined by two parameters, alpha and beta. Alpha and beta can be thought of as intermediate calculations to get to the needed variance estimates.

Reliability is estimated at two different points, at the minimum number of quality reporting events for the measure and at the mean number of quality reporting events per physician. For this measure, the minimum number required to be included is 10 events. Given the structure of the PQRS program, a physician may choose to submit or not submit to PQRS on any given claim. Since these data contain results on a large number of physicians, limiting the reliability analysis to only those physicians who are participating in the program will eliminate the bias introduced by the inclusion of from physicians who are in the data but are not submitting claims to PQRS.

2b6.3 Testing Results *(Provide statistical results, e.g., correlation statistics, comparison of rankings; assessment of adequacy in the context of norms for the test conducted):*

The total number of physicians reporting on this measure is 402. Of those, 48 met the minimum number of quality reporting events for inclusion in the reliability analysis. For this measure 11.94 percent of physicians are included in the analysis, and the average number of quality reporting events for physicians included is 16.13 for a total of 774 events. The average number of quality reporting events for the remaining 88.06 percent of physicians who aren't included is 3.03 for a total of 1,073 events.

For this measure, the reliability at the minimum level of quality reporting events (ie 10 quality events) was .7681 . The reliability at the average number of quality reporting events was .8423

This measure has high reliability when evaluated at both the minimum level of quality reporting events and at the average number of quality events.

Data analyses were conducted by using SAS/STAT software, version 8.2 (SAS Institute, Cary, North Carolina).

2c. Disparities in Care: H M L I NA *(If applicable, the measure specifications allow identification of disparities.)*

2c.1 If measure is stratified for disparities, provide stratified results *(Scores by stratified categories/cohorts):* We encourage the results of this measure to be stratified by race, ethnicity, gender, and primary language, and have included these variables as recommended data elements to be collected.

2c.2 If disparities have been reported/identified (e.g., in 1b), but measure is not specified to detect disparities, please explain:

The PCPI advocates that performance measure data should, where possible, be stratified by race, ethnicity, and primary language to assess disparities and initiate subsequent quality improvement activities addressing identified disparities, consistent with recent national efforts to standardize the collection of race and ethnicity data. A 2008 NQF report endorsed 45 practices including stratification by the aforementioned variables.(1) A 2009 IOM report “recommends collection of the existing Office of Management and Budget (OMB) race and Hispanic ethnicity categories as well as more fine-grained categories of ethnicity(referred to as granular ethnicity and based on one’s ancestry) and language need (a rating of spoken English language proficiency of less than very well and one’s preferred language for health-related encounters).”(2)

References:

(1)National Quality Forum Issue Brief (No.10). Closing the Disparities Gap in Healthcare Quality with Performance Measurement and Public Reporting. Washington, DC: NQF, August 2008.

(2)Race, Ethnicity, and Language Data: Standardization for Health Care Quality Improvement. March 2010. AHRQ Publication No. 10-0058-EF. Agency for Healthcare Research and Quality, Rockville, MD. Available at: <http://www.ahrq.gov/research/iomracereport>. Accessed May 25, 2010.

2.1-2.3 Supplemental Testing Methodology Information:

Steering Committee: Overall, was the criterion, *Scientific Acceptability of Measure Properties*, met? (Reliability and Validity must be rated moderate or high) Yes ☒ No ☒

Provide rationale based on specific subcriteria:

If the Committee votes No, STOP

3. USABILITY

Extent to which intended audiences (e.g., consumers, purchasers, providers, policy makers) can understand the results of the measure and are likely to find them useful for decision making. (**evaluation criteria**)

C.1 Intended Actual/Planned Use (Check all the planned uses for which the measure is intended): Professional Certification or Recognition Program, Public Reporting, Quality Improvement (Internal to the specific organization)

3.1 Current Use (Check all that apply; for any that are checked, provide the specific program information in the following questions): Public Reporting, Professional Certification or Recognition Program, Quality Improvement (Internal to the specific organization)

3a. Usefulness for Public Reporting: H ☒ M ☒ L ☒ I ☒

(The measure is meaningful, understandable and useful for public reporting.)

3a.1. Use in Public Reporting - disclosure of performance results to the public at large (If used in a public reporting program, provide name of program(s), locations, Web page URL(s)). If not publicly reported in a national or community program, state the reason AND plans to achieve public reporting, potential reporting programs or commitments, and timeline, e.g., within 3 years of endorsement: **[For Maintenance** – If not publicly reported, describe progress made toward achieving disclosure of performance results to the public at large and expected date for public reporting; provide rationale why continued endorsement should

be considered.】

This measure was used in the Physician Quality Reporting System from 2009 through 2011. This measure is currently included in PQRS 2012.

<http://www.cms.gov/pqrs>

The PCPI believes that the reporting of participation information is a beneficial first step on a trajectory toward the public reporting of performance results, which is appropriate since the measure has been tested and the reliability of the performance data has been validated. Continued NQF endorsement will facilitate our ongoing progress toward this public reporting objective.

3a.2. Provide a rationale for why the measure performance results are meaningful, understandable, and useful for public reporting. If usefulness was demonstrated (e.g., focus group, cognitive testing), describe the data, method, and results: [The PCPI believes that the reporting of participation information is a beneficial first step on a trajectory toward the public reporting of performance results, which is appropriate since the measure has been tested and the reliability of the performance data has been validated.](#) Continued NQF endorsement will facilitate our ongoing progress toward this public reporting objective.

3.2 Use for other Accountability Functions (payment, certification, accreditation). If used in a public accountability program, provide name of program(s), locations, Web page URL(s): [This measure is used in a Maintenance of Certification program.](#)

3b. Usefulness for Quality Improvement: H● M● L● I●

(The measure is meaningful, understandable and useful for quality improvement.)

3b.1. Use in QI. If used in quality improvement program, provide name of program(s), locations, Web page URL(s):

[For Maintenance – *If not used for QI, indicate the reasons and describe progress toward using performance results for improvement】.*

All PCPI measures are suitable for use in quality improvement initiatives and are made freely available on the PCPI website and through the implementation efforts of medical specialty societies and other PCPI members. The PCPI strongly encourages the use of its measures in QI initiatives and seeks to provide information on such initiatives to PCPI members.

3b.2. Provide rationale for why the measure performance results are meaningful, understandable, and useful for quality improvement. If usefulness was demonstrated (e.g., QI initiative), describe the data, method and results:

The PCPI believes that the use of PCPI measures in quality improvement initiatives is a beneficial way to gather scientific data with which to improve physician performance. This is appropriate since the measure has been tested and the reliability of the performance data has been validated. NQF endorsement will facilitate our ongoing progress toward this quality improvement objective.

Overall, to what extent was the criterion, *Usability*, met? H● M● L● I●

Provide rationale based on specific subcriteria:

4. FEASIBILITY

Extent to which the required data are readily available, retrievable without undue burden, and can be implemented for performance measurement. (**evaluation criteria**)

4a. Data Generated as a Byproduct of Care Processes: H● M● L● I●

4a.1-2 How are the data elements needed to compute measure scores generated? *(Check all that apply).*

See Guidance for Definitions of Rating Scale: H=High; M=Moderate; L=Low; I=Insufficient; NA=Not Applicable

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Data used in the measure are:

generated by and used by healthcare personnel during the provision of care, e.g., blood pressure, lab value, medical condition

4b. Electronic Sources: H ☐ M ☐ L ☐ I ☐

4b.1 Are the data elements needed for the measure as specified available electronically (*Elements that are needed to compute measure scores are in defined, computer-readable fields*): ALL data elements in electronic health records (EHRs)

4b.2 If ALL data elements are not from electronic sources, specify a credible, near-term path to electronic capture, OR provide a rationale for using other than electronic sources:

4c. Susceptibility to Inaccuracies, Errors, or Unintended Consequences: H ☐ M ☐ L ☐ I ☐

4c.1 Identify susceptibility to inaccuracies, errors, or unintended consequences of the measurement identified during testing and/or operational use and strategies to prevent, minimize, or detect. If audited, provide results:

We are not aware of any unintended consequences related to this measurement.

4d. Data Collection Strategy/Implementation: H ☐ M ☐ L ☐ I ☐

A.2 Please check if either of the following apply (*regarding proprietary measures*):

4d.1 Describe what you have learned/modified as a result of testing and/or operational use of the measure regarding data collection, availability of data, missing data, timing and frequency of data collection, sampling, patient confidentiality, time and cost of data collection, other feasibility/implementation issues (e.g., fees for use of proprietary measures):

This measure was found to be reliable and feasible for implementation.

Overall, to what extent was the criterion, *Feasibility*, met? H ☐ M ☐ L ☐ I ☐

Provide rationale based on specific subcriteria:

OVERALL SUITABILITY FOR ENDORSEMENT

Does the measure meet all the NQF criteria for endorsement? Yes ☐ No ☐

Rationale:

If the Committee votes No, STOP.

If the Committee votes Yes, the final recommendation is contingent on comparison to related and competing measures.

5. COMPARISON TO RELATED AND COMPETING MEASURES

If a measure meets the above criteria and there are endorsed or new related measures (either the same measure focus or the same target population) or competing measures (both the same measure focus and the same target population), the measures are compared to address harmonization and/or selection of the best measure before a final recommendation is made.

5.1 If there are related measures (*either same measure focus or target population*) **or competing measures** (*both the same measure focus and same target population*), list the NQF # and title of all related and/or competing measures:

5a. Harmonization

5a.1 If this measure has EITHER the same measure focus OR the same target population as NQF-

endorsed measure(s): Are the measure specifications completely harmonized?

5a.2 If the measure specifications are not completely harmonized, identify the differences, rationale, and impact on interpretability and data collection burden:

5b. Competing Measure(s)

5b.1 If this measure has both the same measure focus and the same target population as NQF-endorsed measure(s):

Describe why this measure is superior to competing measures (e.g., a more valid or efficient way to measure quality); OR provide a rationale for the additive value of endorsing an additional measure. (Provide analyses when possible):

No competing measures have been identified.

CONTACT INFORMATION

Co.1 Measure Steward (Intellectual Property Owner): AMA-convened Physician Consortium for Performance Improvement, 330 N. Wabash Ave., Suite 39300, Chicago, Illinois, 60611

Co.2 Point of Contact: Samantha, Tierney, Samantha.Tierney@ama-assn.org, 312-464-5524-

Co.3 Measure Developer if different from Measure Steward: American Medical Association - Physician Consortium for Performance Improvement, 515 N. State St., Chicago, Illinois, 60654

Co.4 Point of Contact: Diedra, Joseph, MPH, diedra.joseph@ama-assn.org, 312-464-4904-

Co.5 Submitter: Diedra, Joseph, MPH, diedra.joseph@ama-assn.org, 312-464-4904-, American Medical Association - Physician Consortium for Performance Improvement

Co.6 Additional organizations that sponsored/participated in measure development: American Academy of Dermatology and National Committee for Quality Assurance

Co.7 Public Contact: Diedra, Joseph, MPH, diedra.joseph@ama-assn.org, 312-464-4904-, American Medical Association - Physician Consortium for Performance Improvement

ADDITIONAL INFORMATION

Workgroup/Expert Panel involved in measure development

Ad.1 Provide a list of sponsoring organizations and workgroup/panel members' names and organizations. Describe the members' role in measure development.

Raj Behal, MD, MPH (Co-Chair)

Dirk Elston, MD (Co-Chair)

Stephen Bines, MD

Peter C. Dandalides, MD

Evan Farmer, MD

Rutledge Forney, MD

Andrea Gelzer, MD, MS FACP

Robert Gilson, MD

Stephen Helms, MD

Abrar Qureshi, MD

Todd Schlessinger, MD

John Schneider, MD, PhD

Janet (Jessie) Sullivan, MD

Arthur Sober, MD
Steven Strode, MD, Med, MPH
William Wooden, MD

PCPI measures are developed through cross-specialty, multi-disciplinary work groups. All medical specialties and other health care professional disciplines participating in patient care for the clinical condition or topic under study must be equal contributors to the measure development process. In addition, the PCPI strives to include on its work groups individuals representing the perspectives of patients, consumers, private health plans, and employers. This broad-based approach to measure development ensures buy-in on the measures from all stakeholders and minimizes bias toward any individual specialty or stakeholder group. All work groups have at least two co-chairs who have relevant clinical and/or measure development expertise and who are responsible for ensuring that consensus is achieved and that all perspectives are voiced.

Ad.2 If adapted, provide title of original measure, NQF # if endorsed, and measure steward. Briefly describe the reasons for adapting the original measure and any work with the original measure steward: Not applicable

Measure Developer/Steward Updates and Ongoing Maintenance

Ad.3 Year the measure was first released: 2007

Ad.4 Month and Year of most recent revision: 09, 2010

Ad.5 What is your frequency for review/update of this measure? Please see Additional Information/Comments

Ad.6 When is the next scheduled review/update for this measure? 09, 2012

Ad.7 Copyright statement: Physician Performance Measures (Measures) and related data specifications, developed by the American Medical Association (AMA) in collaboration with the Physician Consortium for Performance Improvement (the Consortium) and the National

Committee for Quality Assurance (NCQA) pursuant to government sponsorship under subcontract 6205-05-054 with

Mathematica Policy Research, Inc. under contract 500-00-0033 with Centers for Medicare & Medicaid Services.

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obtain all necessary licenses from the owners of these code sets. The AMA, NCQA, the Consortium and its members disclaim all liability for use or accuracy of any Current Procedural Terminology (CPT®) or other coding contained in the specifications.
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Ad.8 Disclaimers:

Ad.9 Additional Information/Comments: Coding/Specifications updates occur annually. The PCPI has a formal measurement review process that stipulates regular (usually on a three-year cycle, when feasible) review of the measures. The process can also be activated if there is a major change in scientific evidence, results from testing or other issues are noted that materially affect the integrity of the measure.

Date of Submission (MM/DD/YY): 10/03/2011