NATIONAL QUALITY FORUM

Measure Evaluation 4.1 December 2009

This form contains the measure information submitted by stewards. Blank fields indicate no information was provided. Attachments also may have been submitted and are provided to reviewers. The subcriteria and most of the footnotes from the <u>evaluation criteria</u> are provided in Word comments within the form and will appear if your cursor is over the highlighted area. Hyperlinks to the evaluation criteria and ratings are provided in each section.

TAP/Workgroup (if utilized): Complete all yellow highlighted areas of the form. Evaluate the extent to which each subcriterion is met. Based on your evaluation, summarize the strengths and weaknesses in each section.

<u>Note</u>: If there is no TAP or workgroup, the SC also evaluates the subcriteria (yellow highlighted areas).

Steering Committee: Complete all **pink** highlighted areas of the form. Review the workgroup/TAP assessment of the subcriteria, noting any areas of disagreement; then evaluate the extent to which each major criterion is met; and finally, indicate your recommendation for the endorsement. Provide the rationale for your ratings.

Evaluation ratings of the extent to which the criteria are met

C = Completely (unquestionably demonstrated to meet the criterion)

P = Partially (demonstrated to partially meet the criterion)

M = Minimally (addressed BUT demonstrated to only minimally meet the criterion)

N = Not at all (NOT addressed; OR incorrectly addressed; OR demonstrated to NOT meet the criterion)

NA = Not applicable (only an option for a few subcriteria as indicated)

(for NQF staff use) NQF Review #: 1346	NQF Project: Child Health Quality Measures 2010
MEA	SURE DESCRIPTIVE INFORMATION
De.1 Measure Title: Children Who Are Expo	osed To Secondhand Smoke Inside Home
De.2 Brief description of measure: Deterr smoker smokes inside the child's house	mines the perentage of children who live with a smoker and if that
1.1-2 Type of Measure: Outcome De.3 If included in a composite or paired	with another measure, please identify composite or paired measure
De.4 National Priority Partners Priority Ar	rea: Population health

De.5 IOM Quality Domain:

De.6 Consumer Care Need: Staying healthy

CONDITIONS FOR CONSIDERATION BY NQF

Four conditions must be met before proposed measures may be considered and evaluated for suitability as voluntary consensus standards:	NQF Staff
 A. The measure is in the public domain or an intellectual property (measure steward agreement) is signed. Public domain only applies to governmental organizations. All non-government organizations must sign a measure steward agreement even if measures are made publicly and freely available. A.1 Do you attest that the measure steward holds intellectual property rights to the measure and the right to use aspects of the measure owned by another entity (e.g., risk model, code set)? Yes A.2 Indicate if Proprietary Measure (as defined in measure steward agreement): Proprietary measure A.3 Measure Steward Agreement: Agreement will be signed and submitted prior to or at the time of measure submission A.4 Measure Steward Agreement attached: 	A Y N
B. The measure owner/steward verifies there is an identified responsible entity and process to maintain and update the measure on a schedule that is commensurate with the rate of clinical innovation, but at least	B Y

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every 3 years. Yes, information provided in contact section	N
 C. The intended use of the measure includes <u>both</u> public reporting <u>and</u> quality improvement. ▶ Purpose: Public reporting, Internal quality improvement 	C Y N
 D. The requested measure submission information is complete. Generally, measures should be fully developed and tested so that all the evaluation criteria have been addressed and information needed to evaluate the measure is provided. Measures that have not been tested are only potentially eligible for a time-limited endorsement and in that case, measure owners must verify that testing will be completed within 12 months of endorsement. D.1Testing: Yes, fully developed and tested D.2 Have NQF-endorsed measures been reviewed to identify if there are similar or related measures? Yes 	D Y N
(for NQF staff use) Have all conditions for consideration been met? Staff Notes to Steward (<i>if submission returned</i>):	Met Y N
Staff Notes to Reviewers (issues or questions regarding any criteria):	
Staff Reviewer Name(s):	

TAP/Workgroup Reviewer Name:	
Steering Committee Reviewer Name:	
1. IMPORTANCE TO MEASURE AND REPORT	
Extent to which the specific measure focus is important to making significant gains in health care quality (safety, timeliness, effectiveness, efficiency, equity, patient-centeredness) and improving health outcomes for a specific high impact aspect of healthcare where there is variation in or overall poor performance. <i>Measures must be judged to be important to measure and report in order to be evaluated against the remaining criteria</i> . (evaluation criteria) 1a. High Impact	Eval Ratin g
(for NQF staff use) Specific NPP goal:	
 1a.1 Demonstrated High Impact Aspect of Healthcare: Affects large numbers, Leading cause of morbidity/mortality, Severity of illness 1a.2 1a.3 Summary of Evidence of High Impact: The U.S. Department of Health and Human Services' Healthy People 2020 has prioritized the need to decrease nonsmokers' exposure to second hand smoke (TU HP2020-11). 1a.4 Citations for Evidence of High Impact: U.S. Department of Health and Human Services. Healthy People 	1a C P M
2020. http://www.healthypeople.gov/HP2020/.	N
 1b. Opportunity for Improvement 1b.1 Benefits (improvements in quality) envisioned by use of this measure: The effects of exposure to secondhand smoke can be nearly as large as chronic smoking. Additionally, use of tobacco products by household members has an adverse impact on the health of the children. Reducing the proportion of children exposed to secondhand smoke will drastically improve their short and long term health outcomes. 1b.2 Summary of data demonstrating performance gap (variation or overall poor performance) across providers: Nationally, 7.6% of children age 0-17 years are exposed to second hand smoke inside their home. 	1b C□□ M□□ N□

1b.3 Citations for data on performance gap:

Child and Adolescent Health Measurement Initiative. 2007 National Survey of Children's Health, Data Resource Center for Child and Adolescent Health website.

1b.4 Summary of Data on disparities by population group:

Exposure to secondhand smoke inside the home varies by socioeconomic status, race/ethnicity, geographic location and child's age. Compared to children living at 400% FPL or greater, children living below 100% FPL, have 3.23 times the odds of exposure to secondhand smoke inside the come. Fifty percent of African American children and more than one-third of children from low-income families in smoking households were exposed to secondhand smoke inside the home.

Children living in rural locations are more likely to be exposed to secondhand smoke inside the home than children living in urban areas (12.4 vs. 6.5%).

2.6% of Hispanic children, 8.0% of white children and 13.6% of black children are exposed to secondhand smoke inside the home.

Prevalence of exposure to secondhand smoke inside the home increases as children get older. 4.8% of children age 0-5 years, 7.4% of children age 6-11 years and 10.4% of children age 12-17 years are exposed to secondhand smoke inside the home.

1b.5 Citations for data on Disparities:

Barnoya J, Glantz, SA. Cardiovascular effects of second hand smoke: Nearly as large as smoking. American Heart Association Special Report. 24 May, 2005.

Child and Adolescent Health Measurement Initiative. 2007 National Survey of Children's Health, Data Resource Center for Child and Adolescent Health website.

Lam TH. Leung GM. Ho LM. (2001). The effects of environmental tobacco smoke on health services utilization in the first eighteen months of life. Pediatrics, 107(6), 91-97.

Singh GK, Siahpush M, Kogan MD. Disparities in children's exposure to environmental tobacco smoke in the United States, 2007. Pediatrics. 2010;126(1):4-13.

1c. Outcome or Evidence to Support Measure Focus

1c.1 Relationship to Outcomes (For non-outcome measures, briefly describe the relationship to desired outcome. For outcomes, describe why it is relevant to the target population): Reducing the proportion of children exposed to secondhand smoke will drastically improve their short and long term health outcomes.

1c.2-3. Type of Evidence: Other Population-Based Research

1c.4 Summary of Evidence (as described in the criteria; for outcomes, summarize any evidence that healthcare services/care processes influence the outcome): Children who are exposed to secondhand smoke inside the home are less likely to be in very good or excellent overall health than children who live in a home where no one uses tobacco (78.9% vs. 85.5%).

1c.5 Rating of strength/quality of evidence (also provide narrative description of the rating and by whom):

1c.6 Method for rating evidence:

1c.7 Summary of Controversy/Contradictory Evidence:

1c.8 Citations for Evidence (other than guidelines):

1c.9 Quote the Specific guideline recommendation (including guideline number and/or page number):

1c.10 Clinical Practice Guideline Citation: 1c.11 National Guideline Clearinghouse or other URL: 1c C___ P___

M___ N___

1c.12 Rating of strength of recommendation (also provide narrative description of the rating and by whom):	
1c.13 Method for rating strength of recommendation (<i>If different from <u>USPSTF system</u>, also describe rating and how it relates to USPSTF</i>):	
1c.14 Rationale for using this guideline over others:	
TAP/Workgroup: What are the strengths and weaknesses in relation to the subcriteria for <i>Importance to Measure and Report</i> ?	1
Steering Committee: Was the threshold criterion, <i>Importance to Measure and Report</i> , met? Rationale:	1 Y N
2. SCIENTIFIC ACCEPTABILITY OF MEASURE PROPERTIES	
Extent to which the measure, <u>as specified</u> , produces consistent (reliable) and credible (valid) results about the quality of care when implemented. (<u>evaluation criteria</u>)	Eval Ratin g
2a. MEASURE SPECIFICATIONS	
S.1 Do you have a web page where current detailed measure specifications can be obtained? S.2 If yes, provide web page URL:	
2a. Precisely Specified	
2a.1 Numerator Statement (<i>Brief, text description of the numerator - what is being measured about the target population, e.g. target condition, event, or outcome</i>): Percentage of children who live in a household with someone who smokes and smoking occurs inside home	
2a.2 Numerator Time Window (<i>The time period in which cases are eligible for inclusion in the numerator</i>): Encounter or point in time.	
2a.3 Numerator Details (<i>All information required to collect/calculate the numerator, including all codes, logic, and definitions</i>): Children who live in a household with someone who smokes (K9Q40=Yes) and smoking occurs inside home	
(K9Q41=Yes) 2a.4 Denominator Statement (Brief, text description of the denominator - target population being measured): Children age 0-17 years	
2a.5 Target population gender: Female, Male 2a.6 Target population age range: Children age 0-17 years	
2a.7 Denominator Time Window (<i>The time period in which cases are eligible for inclusion in the denominator</i>): Denominator window is a fixed point in time anchored to "current".	
2a.8 Denominator Details (All information required to collect/calculate the denominator - the target population being measured - including all codes, logic, and definitions): Children age 0-17 years	2a- spec s
2a.9 Denominator Exclusions (<i>Brief text description of exclusions from the target population</i>): Excluded from denominator if child does not fall in target population age range of 0-17 years.	P

2a.10 Denominator Exclusion Details (*All information required to collect exclusions to the denominator, including all codes, logic, and definitions***):**

If child is older than 17 years of age, excluded from denominator.

2a.11 Stratification Details/Variables (*All information required to stratify the measure including the stratification variables, all codes, logic, and definitions***):** No stratification is required.

When the Exposure to Secondhand Smoke in Home measure was administered in its most recent form, in the 2007 National Survey of Children's Health, the survey included a number of child demographic variables that allow for stratification of the findings by possible vulnerability:

- Age
- Gender
- Geographic location- State, HRSA Region, National level Rural Urban Commuter Areas (RUCA)
- Race/ethnicity
- Health insurance- type, consistency
- Primary household language
- Household income

• Special Health Care Needs- status and type

2a.12-13 Risk Adjustment Type: No risk adjustment necessary

2a.14 Risk Adjustment Methodology/Variables (*List risk adjustment variables and describe conceptual models, statistical models, or other aspects of model or method***):**

2a.15-17 Detailed risk model available Web page URL or attachment:

2a.18-19 Type of Score: Rate/proportion

2a.20 Interpretation of Score: Better quality = Lower score

2a.21 Calculation Algorithm (*Describe the calculation of the measure as a flowchart or series of steps***):** To receive numerator of child living in a household with someone who smokes and smoking occurs inside home:

-Child lives in household with someone who smokes (K9Q40= Yes) AND -Smoking occurs within the child's home (K9Q41=Yes)

2a.22 Describe the method for discriminating performance (e.g., significance testing):

2a.23 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):* Best guideline to follow is the survey methodology used in the 2007 National Survey of Children's Health.

The goal of the NSCH sample design was to generate samples representative of populations of children within each state. An additional goal of the NSCH was to obtain state-specific sample sizes that were sufficiently large to permit reasonably precise estimates of the health characteristics of children in each state.

To achieve these goals, state samples were designed to obtain a minimum of 1,700 completed interviews. The number of children to be selected in each National Immunization Survey (NIS) estimation area was determined by allocating the total of 1,700 children in the state to each National Immunization Survey (NIS) estimation area within the state in proportion to the total estimated number of households with children in the NIS estimation area. Given this allocation, the number of households that needed to be screened in each NIS estimation area was calculated using the expected proportion of households with children under 18 years of age in the area. Then, the number of telephone numbers that needed to be called was computed using the expected working residential number rate, adjusted for expected nonresponse.

A total of 91,642 interviews were completed from April 2007 to July 2008 for the 2007 National Survey of Children's Health. A random-digit-dialed sample of households with children less than 18 years of age was selected from each of the 50 states and the District of Columbia. One child was randomly selected from all children in each identified household to be the subject of the survey. The respondent was a parent or guardian who knew about the child's health and health care.

2a.24 Data Source (Check the source(s) for which the measure is specified and tested) Survey: Patient	
2a.25 Data source/data collection instrument (<i>Identify the specific data source/data collection instrument, e.g. name of database, clinical registry, collection instrument, etc.</i>): 2007 National Survey of Children's Health	
2a.26-28 Data source/data collection instrument reference web page URL or attachment: URL ftp://ftp.cdc.gov/pub/Health_Statistics/NCHS/slaits/nsch07/1a_Survey_Instrument_English/NSCH_Questionn aire_052109.pdf	
2a.29-31 Data dictionary/code table web page URL or attachment: URL http://nschdata.org/Viewdocument.aspx?item=519	
2a.32-35 Level of Measurement/Analysis (<i>Check the level(s) for which the measure is specified and tested</i>) Population: national, Population: regional/network, Population: states	
2a.36-37 Care Settings (<i>Check the setting(s) for which the measure is specified and tested)</i> Other Applies to any care setting in which child receives care. Can stratify by usual source of care.	
2a.38-41 Clinical Services (Healthcare services being measured, check all that apply) Other Patient Experience	
TESTING/ANALYSIS	
2b. Reliability testing	
2b.1 Data/sample (description of data/sample and size): Qualitative testing of the entire 2007 National Survey of Children's Health was conducted by the National Center for Health Statistics. They conducted cognitive interviews with the 2007 NSCH Computer-Assisted Telephone Interview (CATI) to make sure the entire survey instrument was functioning properly. N=640 interviews were completed over 3 days in December 2006. The questionnaire was then revised and finalized based on feedback from participants in these interviews.	
2b.2 Analytic Method (type of reliability & rationale, method for testing): Cognitive testing was conducted to test reliability and interpretability of questions across population.	
2b.3 Testing Results (reliability statistics, assessment of adequacy in the context of norms for the test conducted): The Maternal and Child Health Bureau leads the development of the NSCH and NS-CSHCN survey and indicators, in collaboration with the National Center for Health Statistics (NCHS) and a national technical expert panel. The expert panel includes representatives from other federal agencies, state Title V leaders, family organizations, and child health researchers, and experts in all fields related to the surveys (adolescent health, family and neighborhoods, early childhood and development etc.). Previously validated questions and scales are used when available. Extensive literature reviewing and expert reviewing of items is conducted for all aspects of the survey. Respondents' cognitive understanding of the survey questions is assessed during the pretest phase and revisions made as required. All final data components are verified by NCHS and DRC/CAHMI staff prior to public release. Face validity is conducted in comparing results with prior years of the survey and/or results from other implementations of items. No specific reliability results are available for this measure. Please contact the CAHMI if quantitative measures are needed.	2b C P M N
2c. Validity testing	
2c.1 Data/sample (<i>description of data/sample and size</i>): 640 interviews were completed over 3 days in December 2006	2c C□
2c.2 Analytic Method (type of validity & rationale, method for testing): Cognitive testing was conducted with parents of children ages 0-17 years (interviews conducted over the phone with residential households).	P M

2c.3 Testing Results (statistical results, assessment of adequacy in the context of norms for the test conducted):	
Please see the references section for peer-reviewed articles which have used these items. Peer-reviewed papers generally undertake their own validity testing in order to meet strict peer review standards. See also Reliability Testing Results above.	
2d. Exclusions Justified	
2d.1 Summary of Evidence supporting exclusion(s):	
2d.2 Citations for Evidence:	
2d.3 Data/sample (description of data/sample and size):	2d
2d.4 Analytic Method (type analysis & rationale):	C P M
2d.5 Testing Results (e.g., frequency, variability, sensitivity analyses):	N NA
2e. Risk Adjustment for Outcomes/ Resource Use Measures	
2e.1 Data/sample (description of data/sample and size):	
2e.2 Analytic Method (type of risk adjustment, analysis, & rationale):	2e
2e.3 Testing Results (risk model performance metrics):	C P M N N
2e.4 If outcome or resource use measure is not risk adjusted, provide rationale:	
2f. Identification of Meaningful Differences in Performance	
2f.1 Data/sample from Testing or Current Use (description of data/sample and size):	
2f.2 Methods to identify statistically significant and practically/meaningfully differences in performance <i>(type of analysis & rationale)</i> :	
2f.3 Provide Measure Scores from Testing or Current Use (description of scores, e.g., distribution by quartile, mean, median, SD, etc.; identification of statistically significant and meaningfully differences in performance):	2f C P M N
2g. Comparability of Multiple Data Sources/Methods	
2g.1 Data/sample (description of data/sample and size):	2g
2g.2 Analytic Method (type of analysis & rationale):	C P M
2g.3 Testing Results (e.g., correlation statistics, comparison of rankings):	N NA
2h. Disparities in Care	2h
2h.1 If measure is stratified, provide stratified results (scores by stratified categories/cohorts):	C P

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2h.2 If disparities have been reported/identified, but measure is not specified to detect disparities, provide follow-up plans:	M N NA
TAP/Workgroup: What are the strengths and weaknesses in relation to the subcriteria for Scientific Acceptability of Measure Properties?	2
Steering Committee: Overall, to what extent was the criterion, <i>Scientific Acceptability of Measure</i> <i>Properties</i> , met? Rationale:	2 C P M N
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. (<u>evaluation criteria</u>)	Eval Ratin g
3a. Meaningful, Understandable, and Useful Information	
3a.1 Current Use: In use	
3a.2 Use in a public reporting initiative (disclosure of performance results to the public at large) (<i>If used in a public reporting initiative, provide name of initiative(s), locations, Web page URL(s).</i> <u>If not publicly reported</u> , state the plans to achieve public reporting within 3 years): U.S. Department of Health and Human Services, Health Resources and Services Administration, Maternal and Child Health Bureau. The Health and Well-Being of Children: A Portrait of States and the Nation 2007. Chartbook based on data from the 2007 National Survey of Children's Health. http://mchb.hrsa.gov/nsch07/index.html.	
3a.3 If used in other programs/initiatives (<i>If used in quality improvement or other programs/initiatives, name of initiative(s), locations, Web page URL(s). <u>If not used for QI</u>, state the plans to achieve use for QI within 3 years): The Data Resource Center websites have been accessed more than 18 million times since 2006. Thousands of state and national researchers, MCH providers and analysts use the data to report valid children's health data.</i> Healthy People 2010 uses items from the national surveys, and several more are slated to be added into Healthy People 2020.	
Testing of Interpretability (Testing that demonstrates the results are understood by the potential users for public reporting and quality improvement) 3a.4 Data/sample (description of data/sample and size): Focus groups were held with numerous stakeholder groups—family advocates, clinicians, Title V leaders, researchers—to obtain feedback on report formats. The Child and Adolescent Health Measurement Initiative led the focus groups and developed reports in accordance with a general consumer information framework. Additional focus groups were held when preparing data and reports for display on the Data Resource Center website. The Data Resource Center executive committee also reviewed report formats for interpretability and applicability.	
3a.5 Methods (e.g., focus group, survey, QI project): Focus groups	3a C 🗌
3a.6 Results (qualitative and/or quantitative results and conclusions):	P M N
3b/3c. Relation to other NQF-endorsed measures	
3b.1 NQF # and Title of similar or related measures:	
(for NQF staff use) Notes on similar/related <u>endorsed</u> or submitted measures:	

 3b. Harmonization If this measure is related to measure(s) already <u>endorsed by NQF</u> (e.g., same topic, but different target population/setting/data source <u>or</u> different topic but same target population): 3b.2 Are the measure specifications harmonized? If not, why? 	3b C P M N N NA
 3c. Distinctive or Additive Value 3c.1 Describe the distinctive, improved, or additive value this measure provides to existing NQF-endorsed measures: 5.1 If this measure is similar to measure(s) already endorsed by NQF (i.e., on the same topic and the same target population), Describe why it is a more valid or efficient way to measure quality: 	3c C P M N N NA
TAP/Workgroup: What are the strengths and weaknesses in relation to the subcriteria for Usability?	3
Steering Committee: Overall, to what extent was the criterion, <i>Usability</i> , met? Rationale:	3 C P M N
4. FEASIBILITY	
Extent to which the required data are readily available, retrievable without undue burden, and can be implemented for performance measurement. (<u>evaluation criteria</u>)	Eval Ratin g
4a. Data Generated as a Byproduct of Care Processes	4a
4a.1-2 How are the data elements that are needed to compute measure scores generated? Survey	C P M N
4b. Electronic Sources	
 4b.1 Are all the data elements available electronically? (elements that are needed to compute measure scores are in defined, computer-readable fields, e.g., electronic health record, electronic claims) Yes 4b.2 If not, specify the near-term path to achieve electronic capture by most providers. 	4b C□ P□
40.2 If not, specify the hear-term path to achieve electronic capture by most providers.	
4c. Exclusions	4c
4c.1 Do the specified exclusions require additional data sources beyond what is required for the numerator and denominator specifications? No	C P
4c.2 If yes, provide justification.	
4d. Susceptibility to Inaccuracies, Errors, or Unintended Consequences	
4d.1 Identify susceptibility to inaccuracies, errors, or unintended consequences of the measure and describe how these potential problems could be audited. If audited, provide results.	4d C P M N
4e. Data Collection Strategy/Implementation	4e
	C

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	<i>#</i> 1J 1 0
4e.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/frequency of data collection, patient confidentiality, time/cost of data collection, other feasibility/ implementation issues: Items are well understood and easy to implement. Items yield very low levels of missing values, don't know or refused answers.	P M N
4e.2 Costs to implement the measure (<i>costs of data collection, fees associated with proprietary measures</i>): Item is public domain and there is no cost associated with its use. Costs associated with administering surveys is not included here.	
4e.3 Evidence for costs:	
4e.4 Business case documentation:	
TAP/Workgroup: What are the strengths and weaknesses in relation to the subcriteria for <i>Feasibility</i> ?	4
Steering Committee: Overall, to what extent was the criterion, <i>Feasibility</i> , met? Rationale:	4 C P M N
RECOMMENDATION	
(for NQF staff use) Check if measure is untested and only eligible for time-limited endorsement.	Time- limite d
Steering Committee: Do you recommend for endorsement? Comments:	Y N A
CONTACT INFORMATION	
Co.1 Measure Steward (Intellectual Property Owner) Co.1 <u>Organization</u> Child and Adolescent Health Measurement Initiative on behalf of the Maternal and Child Health Bureau, Orego Health & Science University, 707 SW Gaines Street, Portland, Oregon, 97239	n
Co.2 <u>Point of Contact</u> Christina, Bethell, Ph.D., MPH, MBA, bethellc@ohsu.edu, 503-494-1892-	
Measure Developer If different from Measure Steward Co.3 <u>Organization</u> Maternal and Child Health Bureau, Parklawn Building Room 18-05, 5600 Fishers Lane, Rockville, Maryland, 208	57
Co.4 <u>Point of Contact</u> Christina, Bethell, Ph.D., MPH, MBA, bethellc@ohsu.edu, 503-494-1892-	
Co.5 Submitter If different from Measure Steward POC Christina, Bethell, Ph.D., MPH, MBA, bethellc@ohsu.edu, 503-494-1892-, Child and Adolescent Health Measure Initiative on behalf of the Maternal and Child Health Bureau	ment
Co.6 Additional organizations that sponsored/participated in measure development	
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.	

The Maternal and Child Health Bureau convenes a Technical Expert Panel (TEP) comprised of dozens of health services researchers, survey methodology experts, and clinical health experts on children's health to develop items for the National Survey of Children's Health. In addition, members of the National Center for Health Statistics are included in item construction and measure development. The TEP participates in all aspects of measure development.

Ad.2 If adapted, provide name of original measure: Ad.3-5 If adapted, provide original specifications URL or attachment

Measure Developer/Steward Updates and Ongoing Maintenance

Ad.6 Year the measure was first released: 2007

Ad.7 Month and Year of most recent revision: 04, 2007

Ad.8 What is your frequency for review/update of this measure? Updated every 4 years when a new National Survey of Children's Health is developed

Ad.9 When is the next scheduled review/update for this measure? 01, 2011

Ad.10 Copyright statement/disclaimers:

Ad.11 -13 Additional Information web page URL or attachment:

Date of Submission (MM/DD/YY): 08/30/2010