

available. The meeting will also be webcast at www.bioethics.gov.

Under authority of Executive Order 13521, dated November 24, 2009, the President established the Commission. The Commission is an expert panel of not more than 13 members who are drawn from the fields of bioethics, science, medicine, technology, engineering, law, philosophy, theology, or other areas of the humanities or social sciences. The Commission advises the President on bioethical issues arising from advances in biomedicine and related areas of science and technology. The Commission seeks to identify and promote policies and practices that ensure scientific research, health care delivery, and technological innovation are conducted in a socially and ethically responsible manner.

The main agenda item for the Commission's seventeenth meeting is to discuss the BRAIN Initiative and ongoing work in neuroscience.

The draft meeting agenda and other information about the Commission, including information about access to the webcast, will be available at www.bioethics.gov.

The Commission welcomes input from anyone wishing to provide public comment on any issue before it. Respectful debate of opposing views and active participation by citizens in public exchange of ideas enhances overall public understanding of the issues at hand and conclusions reached by the Commission. The Commission is particularly interested in receiving comments and questions during the meeting that are responsive to specific sessions. Written comments will be accepted at the registration desk and comment forms will be provided to members of the public in order to write down questions and comments for the Commission as they arise. To accommodate as many individuals as possible, the time for each question or comment may be limited. If the number of individuals wishing to pose a question or make a comment is greater than can reasonably be accommodated during the scheduled meeting, the Commission may make a random selection.

Written comments will also be accepted in advance of the meeting and are especially welcome. Please address written comments by email to info@bioethics.gov, or by mail to the following address: Public Commentary, Presidential Commission for the Study of Bioethical Issues, 1425 New York Avenue NW., Suite C-100, Washington, DC 20005. Comments will be publicly available, including any personally identifiable or confidential business

information that they contain. Trade secrets should not be submitted.

Anyone planning to attend the meeting who needs special assistance, such as sign language interpretation or other reasonable accommodations, should notify Esther Yoo by telephone at (202) 233-3960, or email at Esther.Yoo@bioethics.gov in advance of the meeting. The Commission will make every effort to accommodate persons who need special assistance.

Dated: April 29, 2014.

Lisa M. Lee,

Executive Director, Presidential Commission for the Study of Bioethical Issues.

[FR Doc. 2014-10761 Filed 5-9-14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed information collection project: "*Updating and Expanding the AHRQ QI Toolkit for Hospitals.*" In accordance with the Paperwork Reduction Act of 1995, Public Law 104-13 (44 U.S.C. 3506(c)(2)(A)), AHRQ invites the public to comment on this proposed information collection.

DATES: Comments on this notice must be received by July 11, 2014.

ADDRESSES: Written comments should be submitted to: Doris Lefkowitz, Reports Clearance Officer, AHRQ, by email at doris.lefkowitz@ahrq.hhs.gov.

Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427-1477, or by email at doris.lefkowitz@ahrq.hhs.gov.

SUPPLEMENTARY INFORMATION:

Proposed Project

Updating and Expanding the AHRQ QI Toolkit for Hospitals

AHRQ has developed sets of Quality Indicators (QIs) that can be used to document quality and safety conditions at U.S. hospitals. Three sets of QIs are particularly relevant for hospitals and include: The Inpatient Quality Indicators (IQIs), the Patient Safety Indicators (PSIs), and the Pediatric Quality Indicators (PDIs). The IQIs contain measures of volume, mortality, and utilization for common medical conditions and major surgical procedures. The PSIs are a set of measures to screen for potentially preventable adverse events that patients may experience during hospitalization. The PDIs measure the quality of pediatric health care, mainly focusing on preventable complications that occur as a consequence of hospitalization among pediatric patients. These QIs have been previously developed and evaluated by AHRQ, and are in use at a number of hospitals throughout the country. The QIs and supportive documentation on how to work with them are posted on AHRQ's Web site at www.qualityindicators.ahrq.gov.

Despite the availability of the QIs as tools to help hospitals assess their performance, many U.S. hospitals have limited experience with the use of such measurement tools, or in using quality improvement methods to improve their performance as assessed by these measures. To this end, RAND has previously contracted with AHRQ to develop an AHRQ Quality Indicators Toolkit for Hospitals (Toolkit). This Toolkit is publicly available and is posted on AHRQ's Web site at <http://www.ahrq.gov/professionals/systems/hospital/qitoolkit/index.html>. The Toolkit assists hospitals in both using the QIs and improving the quality and safety of the care they provide, as measured by those indicators. As such, the Toolkit includes: (1) Instruction on how a hospital can apply the QIs to its inpatient data to estimate rates for each indicator; (2) methods the hospital can use to evaluate these QI rates for identifying opportunities for improvement; (3) strategies for implementing interventions (or evidence-based best practices); (4) methods to measure progress and performance on the QIs; (5) tools for evaluating the cost-effectiveness of these changes; and (6) discussion of the value of using the QIs for quality improvement as well as potential challenges and barriers to quality improvement efforts that incorporate the QIs and how to help overcome them.

OMB approval was obtained for the development and evaluation of the original Toolkit in 2012, Development and Evaluation of AHRQ's Quality Indicators Improvement Toolkit (OMB # 0935-0164), which consisted of a protocol very similar to the one described in this statement.

Since the release of the Toolkit in 2012, the QIs have been updated and expanded, best practices have advanced, and many hospitals have improved their understanding of their quality improvement needs as well as increased their familiarity with the use of the Toolkit. These factors all point to the critical need to update the Toolkit. AHRQ has funded RAND, which partners with the University HealthSystem Consortium (UHC), to update and expand the Toolkit, and field test the updated Toolkit with hospitals as they carry out initiatives designed to improve performance on the QIs.

This research has the following goals:
 (1) To assess the usability of the updated Toolkit for hospitals—with an emphasis on the Pediatric Quality Indicators (PDI)—in order to improve the Toolkit, and

(2) To examine hospitals' experiences in implementing interventions to improve their performance on the AHRQ QIs, the results of which will be used to guide successful future applications of the Toolkit.

This study is being conducted by AHRQ through its contractor, the RAND Corporation, under contract number HHS290201000017I, pursuant to AHRQ's statutory authority to conduct and support research on healthcare and on systems for the delivery of such care, including activities with respect to the quality, effectiveness, efficiency, appropriateness and value of healthcare services and with respect to quality measurement and improvement. 42 U.S.C. 299a(a)(1) and (2).

Method of Collection

To achieve the goals of this project the following data collections will be implemented:

(1) Pre/post-test interview protocol—consisting of both open and closed ended questions will be administered prior to implementation of the Toolkit and again post implementation. The purpose of this data collection is to

obtain data on the steps the hospitals took to implement actions to improve performance on the QIs; their plans for making process changes; and their experiences in achieving changes and perceptions regarding lessons learned that could be shared with other hospitals.

(2) Update protocol—consisting of both open and closed ended questions will be administered three times during the study (quarterly during the implementation year). The purpose of this data collection is to capture longitudinal data regarding hospitals' progress in implementing changes, successes and challenges, and plans for subsequent actions. These data will include descriptive information on changes over time in the hospitals' implementation actions and how they are using the Toolkit, as well as experiential information on the perceptions of participants regarding the improvement implementation process and its effects. It also ensures the collection of information close to pertinent events, which avoids the recall bias associated with retrospective reporting of experiences.

(3) Usability testing protocol—also consisting of both open and closed ended questions will be administered once at the end of the evaluation period. The purpose of this data collection is to gather information from the hospitals on how they used each tool in the updated Toolkit, the ease of use of each tool, which tools were most helpful, suggested changes to improve each tool, and suggestions for other tools to add to the updated Toolkit. This information will be used in the revisions of the updated Toolkit following the end of the field test.

All the information obtained from the proposed data collection will be used to strengthen the updated Toolkit before finalizing and disseminating it to hospitals for their use. First, information will be collected from the six hospitals participating in the Toolkit field test about their experiences in implementing performance improvements related to the AHRQ QIs, which will be used to prepare experiential case examples for inclusion in the Toolkit as a resource for other hospitals. Second, feedback will be elicited from them about the usability of the Toolkit, which will be applied to modify and refine the Toolkit so that it

is as responsive as possible to the needs and priorities of the hospitals for which it is intended.

Estimated Annual Respondent Burden

Exhibit 1 shows the estimated annualized burden hours for the respondents' time to participate in this information collection. Three protocols will be used to collect data from respondents in interviews that will take one hour each. The pre/post-test interview protocol will be administered twice—at the beginning and end of the field-test year. The pre-test interviews will be performed as one-hour group interviews with the six hospitals' implementation teams at the start of the year. Each hospital's implementation team is expected to consist of about 5 people. At the end of the year, post-test interviews that last one hour each and use the same protocol as the pre-test interviews will be conducted during site visits at the six hospitals with the implementation team. Thus these 5 people of the implementation team at each hospital will be interviewed twice, both pre- and post-field test. At the post-test site visits, data will also be collected through one-hour interviews performed separately with 4 key stakeholder groups—physicians, nurses, clerks, and others—that are not on the implementation team. Each stakeholder group is expected to consist of about 5 people. Thus these 20 people from the 4 stakeholder groups at each hospital will be interviewed once for one hour post-field test. Interviewing these additional stakeholder groups will ensure that we gather information on stakeholder variations in perceptions and experiences, of which the implementation teams might not be aware.

The quarterly update protocol will be administered quarterly to 2 hospital staff members from each hospital during the year (in months 3, 6, and 9). The usability testing protocol will be administered to 4 staff members once at the end of the evaluation period. The total burden is estimated to be 240 hours.

Exhibit 2 shows the estimated annualized cost burden associated with the respondents' time to participate in the evaluation. The total cost burden is estimated to be \$7,179.

EXHIBIT 1—ESTIMATED ANNUALIZED BURDEN HOURS

Data collection	Number of respondents	Number of responses per respondent	Hours per response	Total burden hours
Pre/Post-Test Interview Protocol with Implementation Team	30	2	1	60

EXHIBIT 1—ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Data collection	Number of respondents	Number of responses per respondent	Hours per response	Total burden hours
Pre/Post-Test Interview Protocol with Stakeholder Groups	120	1	1	120
Quarterly Update Protocol	12	3	1	36
Usability Testing Protocol	24	1	1	24
Total	186	NA	NA	240

EXHIBIT 2—ESTIMATED ANNUALIZED COST BURDEN

Data collection	Number of respondents	Total burden hours	Average hourly wage rate *	Total cost burden
Pre/Post-Test Interview Protocol (Implementation Team and Stakeholder Groups)	150	180	\$29.91	\$5,384
Quarterly Update Protocol	12	36	29.91	1,077
Usability Testing Protocol	24	24	29.91	718
Total	186	240	NA	7,179

* Based upon the mean of the average wages taken from an average of hourly rates for occupations likely to be involved in the QI process (registered nurses, nurse practitioners, medical records and health information technicians, statisticians, and health technologists and technicians). Statistics are taken from the General Medical and Surgical Hospitals industry category in the May 2012 National Industry-Specific Occupational Employment and Wage Estimates from the Bureau of Labor Statistics, U.S. Department of Labor, accessed on January 22, 2014 [www.bls.gov/oes].

Request for Comments

In accordance with the Paperwork Reduction Act, comments on AHRQ's information collection are requested with regard to any of the following: (a) Whether the proposed collection of information is necessary for the proper performance of AHRQ health care research and health care information dissemination functions, including whether the information will have practical utility; (b) the accuracy of AHRQ's estimate of burden (including hours and costs) of the proposed collection(s) of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information upon the respondents, including the use of automated collection techniques or other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the Agency's subsequent request for OMB approval of the proposed information collection. All comments will become a matter of public record.

Dated: May 1, 2014.

Richard Kronick,
AHRQ Director.

[FR Doc. 2014-10752 Filed 5-9-14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meetings

AGENCY: Agency for Healthcare Research and Quality (AHRQ), HHS.

ACTION: Notice of Five AHRQ Subcommittee Meetings.

SUMMARY: The subcommittees listed below are part of AHRQ's Health Services Research Initial Review Group Committee. Grant applications are to be reviewed and discussed at these meetings. Each subcommittee meeting will commence in open session before closing to the public for the duration of the meeting. These meetings will be closed to the public in accordance with 5 U.S.C. App. 2 section 10(d), 5 U.S.C. 552b(c)(4), and 5 U.S.C. 552b(c)(6).

DATES: See below for dates of meetings:

1. Healthcare Safety and Quality Improvement Research (HSQR)

Date: June 17-18, 2014 (Open from 8:00 a.m. to 8:30 a.m. on June 17 and closed for remainder of the meeting).

2. Healthcare Effectiveness and Outcomes Research (HEOR)

Date: June 18, 2014 (Open from 8:00 a.m. to 8:30 a.m. on June 18 and closed for remainder of the meeting).

3. Health Care Research and Training (HCRT)

Date: June 19-20, 2014 (Open from 8:00 a.m. to 8:30 a.m. on June 19 and closed for remainder of the meeting).

4. Healthcare Information Technology Research (HITR)

Date: June 25-27, 2014 (Open from 5:30 p.m. to 6:00 p.m. on June 25 and closed for remainder of the meeting).

5. Health System and Value Research (HSVR)

Date: June 26, 2014 (Open from 8:30 a.m. to 9:00 a.m. on June 26 and closed for remainder of the meeting).

ADDRESSES: Hilton Washington DC/ Rockville Hotel & Executive Meeting Center, 1750 Rockville Pike, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: (To obtain a roster of members, agenda or minutes of the non-confidential portions of the meetings.)

Mrs. Bonnie Campbell, Committee Management Officer, Office of Extramural Research Education and Priority Populations, AHRQ, 540 Gaither Road, Suite 2000, Rockville, Maryland 20850, Telephone (301) 427-1554.

SUPPLEMENTARY INFORMATION: In accordance with section 10(a)(2) of the Federal Advisory Committee Act (5 U.S.C. App. 2), AHRQ announces meetings of the scientific peer review groups listed above, which are subcommittees of AHRQ's Health