1. Gregory Fred Bormann, Mitchell, South Dakota; to acquire voting shares of United Bancorporation, Osseo, Wisconsin, and thereby indirectly acquire voting shares of Farmers State Bank, Stckney, South Dakota; United Bank, Osseo, Wisconsin, Clarke County State Bank, Osceola, Iowa; Bank of Poynette, Poynette, Wisconsin; Cambridge State Bank, Cambridge, Wisconsin; and Lincoln Community Bank, Merrill, Wisconsin.


Michael J. Lewandowski,
Associate Secretary of the Board.

[FR Doc. 2015–17589 Filed 7–16–15; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meeting

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: In accordance with section 10(a)(2) of the Federal Advisory Committee Act (5 U.S.C. App. 2), announcement is made of an Agency for Healthcare Research and Quality (AHRQ) Special Emphasis Panel (SEP) meeting on “AHRQ RFA HS15–001 Patient Safety Learning Laboratories: Innovative Design and Development to Improve Healthcare Delivery Systems (P30).” Each SEP meeting will commence in open session before closing to the public for the duration of the meeting.

DATES: July 21–22, 2015 (Open on July 21 from 8:00 a.m. to 8:30 a.m. and closed for the remainder of the meeting).

ADDRESSES: Gaithersburg Marriott Washingtonian Center, 9751 Washingtonian Boulevard, Gaithersburg, Maryland 20878.

FOR FURTHER INFORMATION CONTACT: Anyone wishing to obtain a roster of members, agenda or minutes of the non-confidential portions of this meeting should contact: Mrs. Bonnie Campbell, Committee Management Officer, Office of Extramural Research, Education and Priority Populations, AHRQ, 540 Gaither Road, Room 2038, Rockville, Maryland 20850, Telephone: (301) 427–1554.

Agenda items for this meeting are subject to change as priorities dictate.

SUPPLEMENTARY INFORMATION: A Special Emphasis Panel is a group of experts in fields related to health care research who are invited by the Agency for Healthcare Research and Quality (AHRQ), and agree to be available, to conduct on an as needed basis, scientific reviews of applications for AHRQ support. Individual members of the Panel do not attend regularly scheduled meetings and do not serve for fixed terms or a long period of time. Rather, they are asked to participate in particular review meetings which require their type of expertise.

Each SEP meeting will commence in open session before closing to the public for the duration of the meeting. The SEP meeting referenced above will be closed to the public in accordance with the provisions set forth in 5 U.S.C. App. 2, section 10(d), 5 U.S.C. 552(b)(4), and 5 U.S.C. 552(b)(6). Grant applications for the “AHRQ RFA HS15–001 Patient Safety Learning Laboratories: Innovative Design and Development to Improve Healthcare Delivery Systems (P30)” are to be reviewed and discussed at this meeting. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Sharon B. Arnold, AHRQ Director.

[FR Doc. 2015–17633 Filed 7–16–15; 8:45 am]

BILLING CODE 4160–90–P

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: “Medical Office Survey on Patient Safety Culture Comparative Database.” In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501–3521, AHRQ invites the public to comment on this proposed information collection.

This proposed information collection was previously published in the Federal Register on March 23rd, 2014 and allowed 60 days for public comment. No comments were received. The purpose of this notice is to allow an additional 30 days for public comment.

DATES: Comments on this notice must be received by August 17, 2015.

ADDRESSES: Written comments should be submitted to: AHRQ’s OMB Desk Officer by fax at (202) 395–6974 (attention: AHRQ’s desk officer) or by email at OIRA_submission@omb.eop.gov (attention: AHRQ’s desk officer).

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

Medical Office Survey on Patient Safety Culture Comparative Database

Background on the Medical Office Survey on Patient Safety Culture (Medical Office SOPS). In 1999, the Institute of Medicine called for health care organizations to develop a “culture of safety” such that their workforce and processes focus on improving the reliability and safety of care for patients (IOM, 1999: To Err is Human: Building a Safer Health System). To respond to the need for tools to assess patient safety culture in health care, AHRQ developed and pilot tested the Medical Office SOPS with OMB approval (OMB NO.0935–0131; Approved July 5, 2007).

The survey is designed to enable medical offices to assess provider and staff opinions about patient safety issues, medical error, and error reporting. The survey includes 38 items that measure 10 composites of patient safety culture. In addition to the composite items, 14 items measure how often medical offices have problems exchanging information with other settings and other patient safety and quality issues. AHRQ made the survey publicly available along with a Survey User’s Guide and other toolkit materials in December 2008 on the AHRQ Web site (located at http://www.ahrq.gov/professionals/quality-patient-safety/patientsafetyculture/medical-office/index.html). Since its release, the survey has been voluntarily used by hundreds of medical offices in the U.S.

The Medical Office SOPS Comparative Database consists of data from the AHRQ Medical Office SOPS.
Medical offices in the U.S. are asked to submit data voluntarily from the survey to AHRQ, through its contractor Westat. The Medical Office SOPS Database (OMB No. 0935-0196, last approved on June 12, 2012) was developed by AHRQ in 2011 in response to requests from medical offices interested in knowing how their patient safety culture survey results compare to those of other medical offices in their efforts to improve patient safety.

**Rationale for the information collection.** The Medical Office SOPS and the Comparative Database support AHRQ’s goals of promoting improvements in the quality and safety of health care in medical office settings. The survey, toolkit materials, and comparative database results are all made publicly available on AHRQ’s Web site. Technical assistance is provided by AHRQ through its contractor at no charge to medical offices, to facilitate the use of these materials for medical office patient safety and quality improvement.

The goal of this project is to renew the Medical Office SOPS Comparative Database. This Database will: (1) Allow medical offices to compare their patient safety culture survey results with those of other medical offices, (2) Provide data to medical offices to facilitate internal assessment and learning in the patient safety improvement process, and (3) Provide supplemental information to help medical offices identify their strengths and areas with potential for improvement in patient safety culture. This study is being conducted by AHRQ through its contractor—Westat, 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 health care services; quality measurement and improvement; and database development. 42 U.S.C. 299a(a)(1), (2), and (8).

**Method of Collection**

To achieve the goal of this project the following activities and data collections will be implemented: (1) Eligibility and Registration Form—The medical office point-of-contact (POC) completes a number of data submission steps and forms, beginning with the completion of an online eligibility and registration form. The purpose of this form is to determine the eligibility status and initiate the registration process for medical offices seeking to voluntarily submit their Medical Office SOPS data to the Medical Office SOPS Comparative Database. (2) Data Use Agreement—The purpose of the data use agreement, completed by the medical office POC, is to state how data submitted by medical offices will be used and provides confidentiality assurances. (3) Medical Office Site Information Form—The purpose of the site information form is to obtain basic information about the characteristics of the medical offices submitting their Medical Office SOPS data to the Medical Office SOPS Comparative Database (e.g., number of providers and staff, ownership, and type of specialty). The medical office POC completes the form. (4) Data Files Submission—The number of submissions to the database is likely to vary each year because medical offices do not administer the survey and submit data every year. Data submission is typically handled by one POC who is either an office manager, nurse manager, or a survey vendor who contracts with a medical office to collect their data. POCs submit data on behalf of 10 medical offices, on average, because many medical offices are part of a health system that includes many medical office sites, or the POC is a vendor that is submitting data for multiple medical offices. After registering, if registrants are deemed eligible to submit data, an automated email is sent to authenticate the account and update the user password. Next the POC enters medical office information and uploads their survey questionnaire and submits a data use agreement. POCs then upload their data file(s), using the medical office data file specifications, to ensure that users submit standardized and consistent data in the way variables are named, coded, and formatted.

Survey data from the AHRQ Medical Office SOPS are used to produce three types of products: (1) A Medical Office SOPS Comparative Database Report that is produced periodically and made publicly available on the AHRQ Web site (see http://www.ahrq.gov/professionals/quality-patient-safety/patientsafetyculture/medical-office/2014/index.html); (2) Individual Medical Office Survey Feedback Reports that are confidential, customized reports produced for each medical office that submits data to the database (the number of reports produced is based on the number of medical offices submitting each year); and (3) Research data sets of individual-level and medical office-level de-identified data to enable researchers to conduct analyses.

Medical offices are asked to voluntarily submit their Medical Office SOPS survey data to the Comparative Database. The data are then cleaned and aggregated and used to produce a Comparative Database Report that displays averages, standard deviations, and percentile scores on the survey’s 38 items that measure 10 composites of patient safety culture, and 14 items measuring how often medical offices have problems exchanging information with other settings and other patient safety and quality issues. The report also displays these results by medical office characteristics (size of office, specialty, geographic region, etc.) and respondent characteristics (staff position). Data submitted by medical offices are used to give each medical office its own customized survey feedback report that presents the medical office’s results compared to the latest comparative database results.

Medical offices use the Medical Office SOPS, Comparative Database Reports and Individual Medical Office Survey Feedback Reports for a number of purposes, to:

- Raise staff awareness about patient safety.
- Diagnose and assess the current status of patient safety culture in their medical office.
- Identify strengths and areas for improvement in patient safety culture.
- Evaluate the cultural impact of patient safety initiatives and interventions.
- Compare patient safety culture survey results with other medical offices in their efforts to improve patient safety and health care quality.

**Estimated Annual Respondent Burden**

Exhibit 1 shows the estimated annualized burden hours for the respondents’ time to participate in the database. An estimated 150 POCs, each representing an average of 10 individual medical offices each, will complete the database submission steps and forms annually. Completing the registration form will take about 3 minutes. The Medical Office Information Form is completed by all POCs for each of their medical offices (150 × 10 = 1,500 forms in total) and is estimated to take 5 minutes to complete. Each POC will complete a data use agreement which takes 3 minutes to complete and submitting the data will take an hour on average. The total burden is estimated to be 291 hours.

Exhibit 2 shows the estimated annualized cost burden based on the respondents’ time to submit their data.
The cost burden is estimated to be $13,968 annually.

**EXHIBIT 1—ESTIMATED ANNUALIZED BURDEN HOURS**

<table>
<thead>
<tr>
<th>Form name</th>
<th>Number of respondents/POCs</th>
<th>Number of responses per POC</th>
<th>Hours per response</th>
<th>Total burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Eligibility/Registration Form</td>
<td>150</td>
<td>1</td>
<td>3/60</td>
<td>8</td>
</tr>
<tr>
<td>Data Use Agreement</td>
<td>150</td>
<td>1</td>
<td>3/60</td>
<td>8</td>
</tr>
<tr>
<td>Medical Office Information Form</td>
<td>150</td>
<td>10</td>
<td>5/60</td>
<td>125</td>
</tr>
<tr>
<td>Data Files Submission</td>
<td>150</td>
<td>1</td>
<td>1</td>
<td>150</td>
</tr>
<tr>
<td>Total</td>
<td>600</td>
<td>NA</td>
<td>NA</td>
<td>291</td>
</tr>
</tbody>
</table>

**EXHIBIT 2—ESTIMATED ANNUALIZED COST BURDEN**

<table>
<thead>
<tr>
<th>Form name</th>
<th>Number of respondents/POCs</th>
<th>Total burden hours</th>
<th>Average hourly wage rate *</th>
<th>Total cost burden</th>
</tr>
</thead>
<tbody>
<tr>
<td>Registration Form</td>
<td>150</td>
<td>8</td>
<td>$48.00</td>
<td>$384</td>
</tr>
<tr>
<td>Data Use Agreement</td>
<td>150</td>
<td>8</td>
<td>48.00</td>
<td>384</td>
</tr>
<tr>
<td>Medical Office Information Form</td>
<td>150</td>
<td>125</td>
<td>48.00</td>
<td>6,000</td>
</tr>
<tr>
<td>Data Files Submission</td>
<td>150</td>
<td>150</td>
<td>48.00</td>
<td>7,200</td>
</tr>
<tr>
<td>Total</td>
<td>600</td>
<td>816</td>
<td>NA</td>
<td>13,968</td>
</tr>
</tbody>
</table>

*Mean hourly wage rate of $48.00 for Medical and Health Services Managers (SOC code 11–9111) was obtained from the May 2013 National Industry-Specific Occupational Employment and Wage Estimates, NAICS 621100—Offices of Physicians located at [http://www.bls.gov/oes/2013/may/naics4_621100.htm](http://www.bls.gov/oes/2013/may/naics4_621100.htm).

**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.

Sharon B. Arnold,
Director.

[FR Doc. 2015–17635 Filed 7–16–15; 8:45 am]

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

**[30Day–15–15BM]**

**Agency Forms Undergoing Paperwork Reduction Act Review**

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to omb@cdc.gov. Direct written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

**Proposed Project**


**Background and Brief Description**

NIOSH, under Public Law 91–506, Sections 20 and 22 (Section 20–22, Occupational Safety and Health Act of 1977) has the responsibility to conduct research relating to innovative methods,