

## 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:

“Evaluation of Phase I Demonstrations of the Pharmacy Quality Alliance.” 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.

This proposed information collection was previously published in the **Federal Register** on February 10th, 2009 and allowed 60 days for public comment. One comment was 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 May 26, 2009.

**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 e-mail at [OIRA\\_submissionomb.eop.gov](mailto:OIRA_submissionomb.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 e-mail at [doris.lefkowitz@ahrq.hhs.gov](mailto:doris.lefkowitz@ahrq.hhs.gov).

#### SUPPLEMENTARY INFORMATION:

##### “Evaluation of Phase I Demonstrations of the Pharmacy Quality Alliance”

AHRQ proposes to conduct an independent evaluation of five Phase I demonstrations undertaken by the Pharmacy Quality Alliance (PQA). The PQA launched the five demonstration projects to test the feasibility of implementing a pharmacy provider report card system, which will be used to provide feedback to pharmacies on

their performance. The goals of the demonstrations are to obtain feedback from pharmacists on the credibility of the performance reports and their utility in performance improvement, and to identify the most efficient and useful ways to implement a performance-based quality reporting system. The evaluation will be conducted for AHRQ by its contractor, the CNA Corporation and Thomas Jefferson Medical College.

The purpose of this evaluation is to identify problems associated with the implementation of a performance-based quality reporting system. The evaluation of the Phase I demonstrations will:

- Test the feasibility and utility of (1) using 15 PQA claims-based measures on pharmacy performance and (2) a survey of consumers about their experience with pharmacy services, which was developed by the PQA;
- Determine the resource (time and cost) requirements for collecting the data and generating the pharmacy performance reports; and
- Provide a base of knowledge that enables the PQA to improve the implementation process, increase operational efficiency, reduce operational costs, and enhance the utility and validity of the performance measures.

This project is being conducted pursuant to AHRQ’s statutory authority to conduct and support research and evaluations on health care and on systems for the delivery of such care, including activities with respect to (1) the quality, effectiveness, efficiency, appropriateness and value of health care services and (2) quality measurement and improvement. 42 U.S.C. 299a(a)(1) and (2).

#### Method of Collection

The evaluation will include the following two data collections: (1) On-site interviews with key staff from each demonstration project and (2) a survey of pharmacy staff. The data will be collected to obtain the following types of information necessary for the evaluation:

- Organizational background related to quality measurement, organizational resources for quality measurement;
- Measurement methodology;
- Opinions on the performance measures;
- The process for disseminating the performance measures;
- Incentives and penalties for participation in pharmacy quality improvement;

- Usability of the performance reports;
- Future directions for quality measurement in the organization; and
- Respondent characteristics.

On-site interviews with key demonstration participants.

On-site interviews will be conducted with up to six persons at each of the five demonstration sites. The study will try to interview representatives from the following job functions: (1) Pharmacy operations management; (2) clinical pharmacy staff; (3) quality improvement; (4) utilization management; (5) analytics management responsible for oversight of performance report analyses; (6) analytics staff assigned to complete the performance reports; (7) information technology (IT) staff responsible for developing and/or coordinating Internet components of the project; and (8) senior management (executive leadership, i.e., Vice President level and above).

#### Survey of Pharmacy Staff

A pharmacy staff survey will be developed to yield additional quantitative data about the demonstration projects. The sample will consist of practicing pharmacists who are participating in the demonstration sites and who received one or more of the performance reports. It will also include field managers and supervisors. At each of the five sites, up to 100 pharmacy staff members will be sampled, with an expected response rate of 75 percent, yielding 75 respondents per site.

#### Estimated Annual Respondent Burden

Exhibit 1 shows the estimated annualized burden hours for this one year data collection. On-site interviews will be conducted with 6 staff members from each of the 5 demonstration projects and will last about 1 hour and 15 minutes. The survey of pharmacists will be completed by about 75 staff members from each demonstration project and is estimated to take 30 minutes to complete. The total estimated annualized burden is 226 hours.

Exhibit 2 shows the estimated annualized cost burden associated with the respondents’ time to participate in this evaluation. The cost burden is estimated to be \$10,753.

EXHIBIT 1—ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of projects	Number of responses per project	Hours per response	Total burden hours
Demonstration Staff Interviews .....	5	6	1.25	38
Survey of Pharmacists .....	5	* 75	30/60	188
Total .....	10	na	na	226

\* We expect that some demonstration projects will have fewer than 75 responses, but we are indicating 75 responses here to avoid underestimating the response burden.

EXHIBIT 2—ESTIMATED ANNUALIZED COST BURDEN

Form name	Number of projects	Total burden hours	Average hourly wage rate *	Total cost burden
Demonstration Staff Interviews .....	5	38	\$47.58	\$1,808
Survey of Pharmacists .....	5	188	47.58	8,945
Total .....	10	226	na	10,753

\* Based on the national average wage for pharmacists (29–1051), National Compensation Survey: Occupational Wages in the United States May 2007, U.S. Department of Labor, Bureau of Labor Statistics.

**Estimated Annual Costs to the Federal Government**

The estimated total cost to the Federal government for this one year evaluation is \$208,874. Exhibit 3 shows a breakdown of the costs.

EXHIBIT 3—ESTIMATED ANNUAL COSTS TO THE FEDERAL GOVERNMENT

Component	Total
Developing the interview guide and survey instrument .....	\$33,905
Preparing OMB clearance submission .....	6,704
Site visits to each demonstration .....	73,368
Analyzing the data from each demonstration site. ....	54,835
Preparing a final report .....	40,062
Total .....	208,874

**Request for Comments**

In accordance with the above-cited Paperwork Reduction Act legislation, 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: April 15, 2009.  
**Carol M. Clancy,**  
*Director.*  
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**BILLING CODE 4160–90–M**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Health Resources and Services Administration**

**Agency Information Collection Activities: Submission for OMB Review; Comment Request**

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, e-mail [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or call the HRSA Reports Clearance Office on (301) 443–1129.

The following request has been submitted to the Office of Management

and Budget for review under the Paperwork Reduction Act of 1995:

**Proposed Project: Voluntary Partner Surveys in the Health Resources and Services Administration—(OMB No. 0915–0212): Extension**

In response to Executive Order 12862, the Health Resources and Services Administration (HRSA) conducts voluntary customer surveys of its partners to assess strengths and weaknesses in program services. To continue the periodic customer or partner satisfaction survey activities, HRSA is requesting an extension of approval from OMB. HRSA partners are, typically, State or local governments, health care facilities, health care consortia, and health care providers. Partner surveys to be conducted by HRSA might include, for example, brief surveys of grantees to determine satisfaction with a technical assistance contractor, or, in-class evaluation forms completed by providers who receive training from HRSA grantees, to measure satisfaction with the training experience. Results of these surveys will be used to plan and direct program efforts as needed to improve service. Focus groups may also be used as a potential method to obtain input on services and training. Focus groups, in-class evaluation surveys, and satisfaction surveys provide valuable input from HRSA partners and customers on agency services and materials.

The estimated annual burden is as follows: