

## EXHIBIT 2—ESTIMATED ANNUALIZED COST BURDEN—Continued

Data collection method	Number of respondents	Total burden hours	Average hourly wage rate*	Total cost burden
Focus Groups .....	99	149	33.51	4,993
Key Informant Interviews .....	30	30	33.51	1,005
Total .....	40,349	5542	NA	185,712

\*Based upon the mean of the average wages for healthcare practitioner and technical occupations (29–0000) presented in the National Compensation Survey: Occupational wages in the United States, May 2009, U.S. Department of Labor, Bureau of Labor Statistics.

**Estimated Annual Costs to the Federal Government**

Exhibit 3 shows the estimated total and annualized cost to the government for this one year project. The total cost is estimated to be \$350,000 to conduct the one-time survey, 11 focus groups,

and 30 key informant interviews and to analyze and present their results. This amount is the contract total for AFYA's contract with AHRQ to evaluate the NGC. This amount, includes the costs for project development and management (\$70,000 or 20% of the entire contract amount); data collection

activities (\$105,000 or 30% of the entire contract amount); data processing and analysis (\$70,000 or 20% of the entire contract amount); and administrative support activities and reporting (\$105,000 or 30% of the entire contract amount).

## EXHIBIT 3—ESTIMATED TOTAL AND ANNUALIZED COST

Cost component	Total cost	Annualized cost
Project Development and Management .....	\$70,000	\$70,000
Data Collection Activities .....	105,000	105,000
Data Processing and Analysis .....	70,000	70,000
Administrative Support and Reporting .....	105,000	105,000
Total .....	350,000	350,000

**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 healthcare research and healthcare 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: November 10, 2010.

**Carolyn M. Clancy,**  
Director.

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES****Food and Drug Administration**

[Docket No. FDA–2009–N–0554]

**Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Manufactured Food Regulatory Program Standards**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled “Manufactured Food Regulatory Program Standards” has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** Daniel Gittleston, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–5156, [Daniel.Gittleston@fda.hhs.gov](mailto:Daniel.Gittleston@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of March 3, 2010 (75 FR 9605), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An

agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0601. The approval expires on September 30, 2013. A copy of the supporting statement for this information collection is available on the Internet at <http://www.reginfo.gov/public/do/PRAMain>.

Dated: November 12, 2010.

**Leslie Kux,**

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–29055 Filed 11–17–10; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES****National Institutes of Health**

**National Institute of Arthritis and Musculoskeletal and Skin Diseases; Notice of Closed Meetings**

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C.,