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 AHRQs 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.

[FR Doc. 2010–29010 Filed 11–17–10; 8:45 am]

BILLING CODE 4160–90–M

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 Gittleson, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–5156, *Daniel.Gittleson@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]

BILLING CODE 4160–01–P

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.,

as amended. 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.

Name of Committee: National Institute of Arthritis and Musculoskeletal and Skin Diseases Special Emphasis Panel, Clinical Trials Review.

Date: December 2, 2010.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Courtyard by Marriott, 5520 Wisconsin Avenue, Chevy Chase, MD 20815.

Contact Person: Charles H Washabaugh, Scientific Review Officer, Scientific Review Branch, National Institute of Arthritis, Musculoskeletal and Skin Diseases, National Institutes of Health, 6701 Democracy Blvd, Room 824, MSC 4872, Bethesda, MD 20817. 301–594–4952. *washabac@mail.nih.gov.*

Name of Committee: National Institute of Arthritis and Musculoskeletal and Skin Diseases Special Emphasis Panel, Small Grants Research Review.

Date: December 8, 2010.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, One Democracy Plaza, 6701 Democracy Boulevard, Bethesda, MD 20892. (Virtual Meeting.)

Contact Person: Eric H. Brown, Scientific Review Officer, Scientific Review Branch, National Institute of Arthritis, Musculoskeletal and Skin Diseases, National Institutes of Health, 6701 Democracy Blvd, Room 824, MSC 4872, Bethesda, MD 20817. (301) 594–4955. *browneri@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.846, Arthritis, Musculoskeletal and Skin Diseases Research, National Institutes of Health, HHS)

Dated: November 10, 2010.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–29091 Filed 11–17–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Agency Information Collection Activities: Request for Information

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security.

ACTION: 60-Day Notice and request for comments; Extension of an existing collection of information: 1651–0023.

SUMMARY: As part of its continuing effort to reduce paperwork and respondent burden, CBP invites the general public and other Federal agencies to comment on an information collection requirement concerning: Request for Information (CBP Form 28). This request for comment is being made pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104–13; 44 U.S.C. 3505(c)(2)).

DATES: Written comments should be received on or before January 18, 2011, to be assured of consideration.

ADDRESSES: Direct all written comments to U.S. Customs and Border Protection, *Attn:* Tracey Denning, Regulations and Rulings, Office of International Trade, 799 9th Street, NW., 5th Floor, Washington, DC 20229–1177.

FOR FURTHER INFORMATION CONTACT: Requests for additional information should be directed to Tracey Denning, U.S. Customs and Border Protection, Regulations and Rulings, Office of International Trade, 799 9th Street, NW., 5th Floor, Washington, DC 20229–1177, at 202–325–0265.

SUPPLEMENTARY INFORMATION: CBP invites the general public and other Federal agencies to comment on proposed and/or continuing information collections pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104–13; 44 U.S.C. 3505(c)(2)). The comments should address: (a) Whether the collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimates of the burden of the collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden including the use of automated collection techniques or the use of other forms of information technology; and (e) the annual costs burden to respondents or record keepers from the collection of information (a total capital/startup costs and operations and maintenance costs). The comments that are submitted will be summarized and included in the CBP request for Office of Management and Budget (OMB) approval. All comments will become a matter of public record. In this document CBP is soliciting comments concerning the following information collection:

Title: Request for Information.

OMB Number: 1651–0023.

Form Number: CBP Form 28.

Abstract: Under 19 U.S.C. 1500 and 1401a, Customs and Border Protection (CBP) is responsible for appraising imported merchandise by ascertaining its value, classifying merchandise under the tariff schedule, and assessing a rate and amount of duty to be paid. On occasions when the invoice or other documentation does not provide sufficient information for appraisement or classification, the CBP Officer requests additional information through the use of CBP Form 28, "Request for Information". This form is completed by CBP personnel requesting additional information and the importers, or their agents, respond in the format of their choice. CBP Form 28 is provided for by 19 CFR 151.11. A copy of this form and instructions are available at *http://forms.cbp.gov/pdf/CBP_Form_28.pdf.*

Current Actions: This submission is being made to extend the expiration date with no change to the burden hours or to CBP Form 28.

Type of Review: Extension (without change).

Affected Public: Businesses.

Estimated Number of Respondents: 60,000.

Estimated Time per Respondent: 1 hour.

Estimated Total Annual Burden Hours: 60,000.

Dated: November 15, 2010.

Tracey Denning,

Agency Clearance Officer, U.S. Customs and Border Protection.

[FR Doc. 2010–29085 Filed 11–17–10; 8:45 am]

BILLING CODE 9111–14–P

NATIONAL INDIAN GAMING COMMISSION

Notice of Inquiry and Request for Information; Notice of Consultation

AGENCY: National Indian Gaming Commission.

ACTION: Notice of inquiry; notice of Tribal consultations.

Authority: 25 U.S.C. 2706(b)(10); E.O. 13175.

SUMMARY: This Notice of Inquiry and Notice of Consultation advises the public that the National Indian Gaming Commission (NIGC) is conducting a comprehensive review of all regulations promulgated to implement the Indian Gaming Regulatory Act (IGRA), 25 U.S.C. 2701 *et seq.* The Commission is taking a fresh look at its rules in order to determine whether amendments are necessary to more effectively implement IGRA's policies of protecting Indian gaming as a means of generating Tribal revenue, ensuring that gaming is conducted fairly and honestly by both the operator and players, and ensuring that Tribes are the primary beneficiaries of gaming operations. The Commission's