Board (PRB) membership is required by 5 U.S.C. 4314(c)(4). The PRB reviews and evaluates the initial appraisal of a senior executive's performance by the supervisor, and makes recommendations regarding performance ratings, performance awards, and pay-for-performance pay adjustments to the Chairman.

The following individuals have been designated to serve on the Commission's Performance Review Board:

Eileen Harrington, Executive Director, Chair.

Willard K. Tom, General Counsel.

Pauline M. Ippolito, Deputy Director, Bureau of Economics.

Richard A. Feinstein, Director, Bureau of Competition.

Jessica L. Rich, Deputy Director, Bureau of Consumer Protection.

By direction of the Commission.

Richard C. Donohue,

Acting Secretary.

[FR Doc. 2011–21021 Filed 8–17–11; 8:45 am]

BILLING CODE 6750–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the National Biodefense Science Board

AGENCY: Department of Health and Human Services, Office of the Secretary.

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Services is hereby giving notice that the National Biodefense Science Board (NBSB) will be holding a public meeting. The meeting is open to the public.

DATES: The NBSB will hold a public meeting on September 22, 2011 from 9 a.m. to 5 p.m. E.S.T. The agenda is subject to change as priorities dictate.

ADDRESSES: Almas Temple—Sphinx Grand Ballroom; 1315 K Street, NW., Washington, DC 20005 (adjacent to the Hamilton Crowne Plaza Washington). To attend by teleconference, please check the NBSB September meeting webpage, *http://www.phe.gov/Preparedness/legal/boards/nbsb/meetings/Pages/110922meeting.aspx.* Individuals who wish to attend the meeting in person should send an e-mail to *NBSB@HHS.GOV* with ''NBSB Registration'' in the subject line.

FOR FURTHER INFORMATION: *e-mail: NBSB@HHS.GOV*

SUPPLEMENTARY INFORMATION: Pursuant to section 319M of the Public Health Service Act (*42 U.S.C. 247d–7f*) and section 222 of the Public Health Service Act (*42 U.S.C. 217a*), the Department of Health and Human Services established the National Biodefense Science Board. The Board shall provide expert advice and guidance to the Secretary on scientific, technical, and other matters of special interest to the Department of Health and Human Services regarding current and future chemical, biological, nuclear, and radiological agents, whether naturally occurring, accidental, or deliberate. The Board may also provide advice and guidance to the Secretary and/or the Assistant Secretary for Preparedness and Response on other matters related to public health emergency preparedness and response.

Background: The majority of this public meeting will be dedicated to a discussion of the findings of the NBSB's Anthrax Vaccine Working Group. Subsequent agenda topics will be added as priorities dictate. Any additional agenda topics will be available on the Board's September meeting webpage prior to the public meeting.

Availability of Materials: The meeting agenda and materials will be posted prior to the meeting on the September meeting webpage at *http://www.phe.gov/Preparedness/legal/boards/nbsb/meetings/Pages/110922meeting.aspx.*

Procedures for Providing Public Input: Any member of the public providing oral comments at the meeting must sign-in at the registration desk and provide his/her name, address, and affiliation. All written comments must be received prior to September 21, 2011 and should be sent by e-mail to *NBSB@HHS.GOV* with ''NBSB Public Comment'' as the subject line. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should e-mail *nbsb@hhs.gov.*

Dated: August 8, 2011.

Nicole Lurie,

Assistant Secretary for Preparedness and Response.

[FR Doc. 2011–21163 Filed 8–17–11; 8:45 am]

BILLING CODE 4150–37–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Low Income Home Energy Assistance Program (LIHEAP) Household Report.

OMB No.: 0970–0060.

Description: This report is an annual activity required by statute (42 U.S.C. 8629) and Federal reguations (45 CFR 96.92) for the Low Income Home Energy Assistance Program (LIHEAP). Submission of the completed report is one requirement for LIHEAP grantees applying for Federal LIHEAP block grant funds. States, the District of Columbia, and the Commonwealth of Puerto Rico are required to report statistics for the previous Federal fiscal year on:

- Assisted and applicant households, by type of LIHEAP assistance;
- Assisted and applicant households, by type of LIHEAP assistance and poverty level;
- Assisted households, regardless of the type(s) of LIHEAAP assistance;
- Assisted households, by type of LIHEAP assistance, having at least one vulnerable member broken out; by a person at least 60 years or younger, disabled person, or a child five years older or younger;
- Assisted households, by type of LIHEAP assistance, with at least one member age 2 years or under;
- Assisted households, by type of LIHEAP assistance, with at least one member ages 3 years through 5 years; and
- Assisted households, regardless of the type(s) of LIHEAP assistance, having at least one member 60 years or older, disabled, or five years old or younger.

Insular areas (other than the Commonwealth of Puerto Rico) and Indian Tribal Grantees are required to submit data only on the number of households receiving heating, cooling, energy crisis, or weatherization benefits.

The information is being collected for the Department's annual *LIHEAP Report to Congress.* The data also provide information about the use of LIHEAP funds. Finally, the data are used in the calculation of LIHEAP performance measures under the Government Performance and Results Act of 1993. The data elements will allow the accuracy of measuring LIHEAP targeting performance and LIHEAP cost efficiency.

Respondents: State Governments, Tribal Governments, Insular Areas, the District of Columbia, and the Commonwealth of Puerto Rico.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Assisted Household Report—Long Form	52	1	25	1,300
Assisted Household Report—Short Form	164	1	1	164
Applicant Household Report	52	1	13	676

Estimated Total Annual Burden Estimates: 2,140.

Additional Information:

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, *Attn:* ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. *E-mail address: infocollection@acf.hhs.gov.*

OMB Comment:

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following:

Office of Management and Budget, Paperwork Reduction Project, *Fax:* 202–395–6974, *Attn:* Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2011–21107 Filed 8–17–11; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0424]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Temporary Marketing Permit Applications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by September 19, 2011.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, *Attn:* FDA Desk Officer, FAX: 202–395–7285, or e-mailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0133. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3793.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Temporary Marketing Permit Applications—21 CFR 130.17(c) and (i)—(OMB Control Number 0910–0133)—Extension

Section 401 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 341) directs FDA to issue regulations establishing definitions and standards of identity for food ''[w]henever * * * such action will promote honesty and fair dealing in the interest of consumers * * *.'' Under section 403(g) of the FD&C Act (21 U.S.C. 343(g)), a food that is subject to a definition and standard of identity prescribed by regulation is misbranded if it does not conform to such definition and standard of identity. Section 130.17 (§ 130.17) provides for the issuance by FDA of temporary marketing permits that enable the food industry to test consumer acceptance and measure the technological and commercial feasibility in interstate commerce of experimental packs of food that deviate from applicable definitions and standards of identity. Section 130.17(c) enables the Agency to monitor the manufacture, labeling, and distribution of experimental packs of food that deviate from applicable definitions and standards of identity. The information so obtained can be used in support of a petition to establish or amend the applicable definition or standard of identity to provide for the variations. Section 130.17(i) specifies the information that a firm must submit to FDA to obtain an extension of a temporary marketing permit.

In the **Federal Register** of June 10, 2011 (76 FR 34080), FDA published a 60-day notice requesting public comment on the proposed collection of information. No comments were received.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

21 CFR section	No. of respondents	No. of responses per respondent	Total annual responses	Average burden per response	Total hours
130.17(c)	13	2	26	25	650