Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2017–17699 Filed 8–21–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Application Requirements for the Low Income Home Energy Assistance Program (LIHEAP) Plan.

OMB No.: 0970–0075.

Description: States, including the District of Columbia, tribes, tribal organizations, and U.S. territories applying for LIHEAP block grant funds must, prior to receiving federal funds, submit an annual application (Model Plan, ACF–122) that meets the LIHEAP statutory and regulatory requirements. In addition to the Model Plan, grantees are also required to complete the Mandatory Grant Application SF–424–Mandatory, which is the first section of the Model Plan.

The LIHEAP Model Plan is an electronic form and is submitted to the Administration for Children and Families (ACF), Office of Community Services (OCS) through the On-line Data Collection (OLDC) system within GrantSolutions, which is currently being used by all LIHEAP grantees to submit other required LIHEAP reporting forms. In order to reduce the reporting burden, all data entries from each grantee's prior year's submission of the Model Plan in OLDC is saved and re-populated (cloned) into the form for the following fiscal year's application.

OCS seeks renewal of this form without any changes. A sample model plan showing these proposed changes can be found on the U.S. Department of Health and Human Services, ACF/OCS LIHEAP Program Resources page at: *https://www.acf.hhs.gov/ocs/resource/funding-applications.*

On April 3, 2017, ACF published a **Federal Register** Notice seeking 60 days of public comment on this proposed information collection. One state grantee provided comments. ACF revised the Plan to address the comments by ensuring that open field boxes and attachment capability are available if the answer choices are insufficient to address the questions.

The revised model plan can be viewed on the OCS Web site at: *http://www.acf.hhs.gov/programs/ocs/programs/liheap.*

Respondents: State, the District of Columbia, U.S. Territories and Tribal governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
LIHEAP Detailed Model Plan	210	1	0.50	105

Estimated Total Annual Burden Hours (all respondents): 105.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201. Attention Reports Clearance Officer. All requests should be identified by the title of the information collection. Email 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, Email: *OIRA_SUBMISSION@OMB.EOP.GOV*, Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2017–17681 Filed 8–21–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–4885]

Pediatric Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA or the Agency) announces a forthcoming public advisory committee meeting of the Pediatric Advisory Committee (PAC). The general function of the committee is to provide advice and recommendations to the Agency on FDA's regulatory issues. The meeting will be open to the public. FDA is establishing a docket for public comments.

DATES: The meeting will be held on September 11, 2017, from 8:30 a.m. to 5:30 p.m. and September 12, 2017, from 8:30 a.m. to 1 p.m.

ADDRESSES: Hilton Washington DC/Rockville Hotel & Executive Meeting Center, 1750 Rockville Pike, Rockville, MD 20852. The hotel's telephone number is 301–468–1100. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at *http://www3.hilton.com/en/hotels/maryland/hilton-washington-dc-rockville-hotel-and-executive-meeting-ctr-IADMRHF/index.html.*

FDA is establishing a docket for public comment on this document. The docket number is FDA–2017–N–4885. The docket will close on September 13, 2017. Submit either electronic or written comments on this public meeting by that date. Late, untimely comments will not be considered. Electronic comments must be submitted on or before September 13, 2017. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of September 13, 2017. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Comments received on or before August 28, 2017, will be provided to the committee. Comments received after that date will be taken into consideration by FDA.