DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0554]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Manufactured Food Regulatory Program Standards

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 April 2, 2010.

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 0910–0601. Also include the FDA docket number found in brackets in the heading of this document.

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

Manufactured Food Regulatory Program Standards—(OMB Control Number 0910–0601)—Extension

I. Background

In the **Federal Register** of July 20, 2006 (71 FR 41221), FDA announced the availability of a draft document entitled "Manufactured Food Regulatory Program Standards." These draft program standards are the framework that States should use to design and manage its manufactured food program. The implementation of the program standards will be negotiated as an option for payment under the State food contract. States that are awarded this option will receive up to $25,000 over a period of 5 years to fully implement the program standards. In the first year of implementing the program standards, the State program conducts a baseline self-assessment to determine if they meet the elements of each standard. The State program should use the worksheets and forms contained herein; however it can use alternate forms that are equivalent. The State program maintains the documents and verifying records required for each standard. The information contained in the documents must be current and fit-for-use. If the State program fails to meet all program elements and documentation requirements of a standard, it develops a strategic plan to fully implement the program standards in 5 years. The strategic plan includes the following: (1) The individual element or documentation requirement of the standard that was not met, (2) improvements needed to meet the program element or documentation requirement of the standard, and (3) projected completion dates for each task.

II. Electronic Access

Persons with access to the Internet may obtain the draft program standards at *http://www.fda.gov/downloads/RegulatoryInformation/Guidances/UCM125448.pdf.*

In the **Federal Register** of December 2, 2009 (74 FR 63154), 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]

No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
44	1	44	40	1,760

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.

TABLE 2.—ESTIMATED FIRST-YEAR BASELINE SELF-ASSESSMENT BURDEN[1]

No. of Respondents	Five-Year Frequency per Response	Total First-Year Responses	Hours per Response	Total Hours
17	1	17	200	3,400

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.

Dated: February 25, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–4340 Filed 3–2–10; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Notice of Meeting; National Commission on Children and Disasters

AGENCY: Administration for Children and Families, Department of Health and Human Services.

ACTION: Notice of meeting.

DATE: The meeting will be held on Tuesday, March 23, 2010, from 9:30 a.m. to 3:30 p.m.

ADDRESSES: The meeting will be held at the Administration for Children and Families, 901 D Street SW., Washington, DC 20024. To attend either in person or via teleconference, please register by 5 p.m. Eastern Time, March 18, 2010. To register, please e-mail *jacqueline.haye@acf.hhs.gov* with "Meeting Registration" in the subject line, or call (202) 205–9560. Registration