DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0017]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Voluntary National Retail Food Regulatory Program Standards

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is extending the comment period for the notice of public meeting, that appeared in the Federal Register of August 9, 2010 (75 FR 47820). In the notice, FDA requested comments to gather stakeholder input on the development of a generic drug user fee program. The Agency is taking this action to allow interested persons additional time to submit comments.

DATES: Submit either electronic or written comments by August 1, 2011.

ADDRESSES: You may submit comments, identified by Docket No. FDA–2010–N–0381, by any of the following methods:

Electronic Submissions
Submit electronic comments in the following ways:

• Federal eRulemaking Portal: http://www.regulations.gov. Follow the instructions for submitting comments.

Written Submissions
Submit written submissions in the following ways:

• Fax: 301–827–6870.
• Mail/Hand delivery/Courier (for paper, disk, or CD–ROM submissions): Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Lan, 1061, Rockville, MD 20852.

Instructions: All submissions received must include the Agency name and docket number for this rulemaking. All comments received may be posted without change to http://www.regulations.gov, including any personal information provided. For additional information on submitting comments, see the “Request for Comments” heading of the SUPPLEMENTARY INFORMATION section of this document.

Docket: For access to the docket to read background documents or comments received, go to http://www.regulations.gov and insert the docket number, found in brackets in the heading of this document, into the “Search” box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Lan, 1061, Rockville, MD 20852.

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

II. Request for Comments
Interested persons may submit to the Division of Dockets Management (see ADDRESSES) either electronic or written comments on this document. It is only necessary to send one set of comments. It is no longer necessary to send two copies of mailed comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: June 3, 2011.

Leslie Kux,
Acting Assistant Commissioner for Policy.

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BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0012]

Strengthen and Promote the Role of Local Health Departments in Retail Food Safety Regulation (U–50)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of grant funds for the support of a cooperative agreement between the Center for Food Safety and Applied Nutrition (CFSAN) and the National Association of County and City Health Officials (NACCHO). The goal of the cooperative agreement for CFSAN is to have NACCHO conduct work that will strengthen the role of local health departments and help FDA/CFSAN