I. Background

In the Federal Register of August 2, 2010 (75 FR 45130), FDA published a notice of availability of a draft CPG with a 90-day comment period to request comments on its proposal that certain criteria should be considered in recommending enforcement action against animal feed or feed ingredients that are adulterated due to the presence of Salmonella. The Agency has received a request for a 60-day extension of the comment period for the draft CPG. The request conveyed concern that the current 90-day comment period does not allow sufficient time to develop a meaningful or thoughtful response to the draft CPG. FDA has considered the request and is extending the comment period for the draft CPG for 60 days, until December 31, 2010.

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: October 26, 2010.
Dara Corrigan,
Associate Commissioner for Regulatory Affairs.

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[DOcket No. FDA–2010–D–0378]

Draft Compliance Policy Guide Sec. 690.800 Salmonella in Animal Feed: Availability; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA) is extending to December 31, 2010, the comment period for a notice of availability of a draft compliance policy guide (CPG) that appeared in the Federal Register of August 2, 2010 (75 FR 45130). In the document, FDA requested comments on its proposal that certain criteria should be considered in recommending enforcement action against animal feed or feed ingredients that are adulterated due to the presence of Salmonella. The Agency is taking this action in response to requests for an extension to allow interested persons additional time to submit comments.

DATES: Submit either electronic or written comments by December 31, 2010.

ADDRESSES: Submit electronic comments on the draft CPG to http://www.regulations.gov. Submit written comments on the draft CPG to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Kim Young, Center for Veterinary Medicine (HFV–230), Food and Drug Administration, 7519 Standish Pl., MPN–4, rm. 106, Rockville, MD 20855, 240–276–9200, e-mail: Kim.young@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:
reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advise the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class.

**Matters To Be Discussed:** The agenda for the Advisory Board meeting includes: NIOSH Program Update and Program Evaluation; Department of Labor (DOL) Program Update; Department of Energy (DOE) Program Update; Los Alamos National Laboratory Work Group Update; Board Session to Discuss Evaluating Exposure Potential for Radiological Materials in Minor Quantities or Uses; SEC petitions for: Linde Ceramics Plant (Towanda, New York), General Electric Company (Evdendale, Ohio), Dow Chemical (Madi son, Illinois), Simonds Saw and Steel Company (Lockport, New York), Hangar 481 of Kirkland Airforce Base (Albuquerque, New Mexico), BWX Technologies (Lynchburg, Virginia), and Texas City Chemicals Inc. (Texas City, Texas); SEC Petition Status Updates; SEC Class Definition Assessment Report; Subcommittee and Work Group Reports; and Board Work Sessions.

The agenda is subject to change as priorities dictate.

In the event an individual cannot attend, written comments may be submitted in accordance with the redaction policy provided below. Any written comments received will be provided at the meeting and should be submitted to the contact person below well in advance of the meeting.

**Policy on Redaction of Board Meeting Transcripts (Public Comment):** (1) If a person making a comment gives his or her name, no attempt will be made to redact that name; (2) NIOSH will take reasonable steps to ensure that individuals making public comment are aware of the fact that their comments (including their name, if provided) will appear in a transcript of the meeting posted on a public Web site. Such reasonable steps include: (a) A statement read at the start of each public comment period stating that transcripts will be posted and names of speakers will not be redacted; (b) A printed copy of the statement mentioned in (a) above will be displayed on the table where individuals sign up to make public comments; (c) A statement such as outlined in (a) above will also appear with the agenda for the Board Meeting when it is posted on the NIOSH Web site; (d) A statement such as in (a) above will appear in the Federal Register Notice that announces Board and Subcommittee meetings; (3) If an individual in making a statement reveals personal information (e.g., medical information) about themselves that information will not usually be redacted. The NIOSH FOIA coordinator will, however, review such revelations in accordance with the Freedom of Information Act and the Federal Advisory Committee Act and if deemed appropriate, will redact such information; (4) All disclosures of information concerning third parties will be redacted; and (5) If it comes to the attention of the DFO that an individual wishes to share information with the Board but objects to doing so in a public forum, the DFO will work with that individual, in accordance with the Federal Advisory Committee Act, to find a way that the Board can hear such comments.

**Contact Person for More Information:** Theodore Katz, M.P.A., Designated Federal Officer, NIOSH, CDC, 1600 Clifton Road, NE., Mailstop E–20, Atlanta, Georgia 30333; telephone: (513) 533–6800, toll free: 1 (800) CDC–INFO, e-mail: ocas@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register Notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.


Elaine L. Baker, Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

National Institutes of Health

**National Cancer Institute; Notice of Closed Meeting**

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 meeting.

The meeting 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 Cancer Institute Special Emphasis Panel; Clinical Proteomic Technologies for Cancer Initiative Research.

**Date:** December 13–15, 2010.

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

**Agenda:** To review and evaluate grant applications.

**Place:** Bethesda North Marriott Hotel & Conference Center, 5701 Marinelli Road, Bethesda, MD 20852.

**Contact Person:** Adriana Stoica, PhD, Scientific Review Officer, Special Review &