

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Food and Drug Administration****[Docket No. FDA-2011-N-0002]****Risk Communication Advisory Committee; Notice of Meeting****AGENCY:** Food and Drug Administration, HHS.**ACTION:** Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Risk Communication Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the Agency on FDA's regulatory issues.

Date and Time: The meeting will be held on May 5, 2011, from 8 a.m. to 5 p.m.

Location: Food and Drug Administration, 5630 Fishers Lane, Conference Room, rm. 1066, Rockville, MD 20857.

Contact Person: Lee L. Zwanziger, Office of Policy, Planning and Budget, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 3278, Silver Spring, MD 20993-0002, 301-796-9151, FAX: 301-847-8611, e-mail: RCAC@fda.hhs.gov, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), and follow the prompts to the desired center or product area. Please call the Information Line for up-to-date information on this meeting. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site and call the appropriate advisory committee hot line/phone line to learn about possible modifications before coming to the meeting.

Agenda: On May 5, 2011, the committee will hear and discuss developments in FDA's ongoing communications programs. The discussion will focus on the use of different channels for information dissemination, tracking how information is gathered and spread, and thoughts on reaching less accessible target audiences.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background

material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at <http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm>. Scroll down to the appropriate advisory committee link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before April 29, 2011. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. on May 5, 2011. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before April 21, 2011. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by April 22, 2011.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Lee L. Zwanziger at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at <http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm> for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: March 17, 2011.

Leslie Kux,*Acting Assistant Commissioner for Policy.*

[FR Doc. 2011-6788 Filed 3-22-11; 8:45 am]

BILLING CODE 4160-01-P**DEPARTMENT OF HEALTH AND HUMAN SERVICES****Indian Health Service****Memorandum of Agreement Between the Indian Health Service and the Department of Interior; Bureau of Indian Affairs and Bureau of Indian Education****AGENCY:** Indian Health Service, HHS.**ACTION:** Notice.

SUMMARY: The Indian Health Service (IHS) is providing notice of a Memorandum of Agreement (MOA) between the IHS and the Department of the Interior (DOI), signed in 2009, and has developed an amendment to that MOA that includes language consistent with Section 703 of the Indian Health Care Improvement Act (IHCIA), Public Law 94-437, as amended. The purpose of the MOA and the amendment is to advance our partnership with Tribes and Federal stakeholders on alcohol and substance abuse prevention and treatment. The Patient Protection and Affordable Care Act's, Public Law 111-148, permanent authorization of the Indian Health Care Improvement Act (IHCIA) establishes timelines and requirements for coordinated actions by the Department of Interior (DOI), the Department of Health and Human Services (HHS), Tribes and Tribal organizations. Specifically, Section 703 of the IHCIA provides new authorities that permit the DOI and HHS, acting through the Indian Health Service (IHS), to develop and enter into a Memorandum of Agreement (MOA), or review and update any existing memoranda of agreement, as required by Section 4205 of the Indian Alcohol and Substance Abuse Prevention and Treatment Act of 1986 (25 U.S.C.2411). DOI and IHS signed an MOA on this topic in 2009, and have developed an amendment to that MOA that includes language consistent with the new IHCIA provision. In accordance with Section 703 of the IHCIA, which states that the MOA between the IHS and DOI shall be published in the **Federal Register**, the agency is publishing notice of this MOA and the amendment to this MOA.

DATES: The original MOA was effective on December 12, 2009. The amendment is effective March 1, 2011.