

**DEPARTMENT OF HEALTH AND HUMAN SERVICES****Office of the National Coordinator for Health Information Technology; American Health Information Community Chronic Care Workgroup Meeting****ACTION:** Announcement of meeting.**SUMMARY:** This notice announces the 24th meeting of the American Health Information Community Chronic Care Workgroup in accordance with the Federal Advisory Committee Act (Pub. L. No. 92-463, 5 U.S.C., App.)**DATES:** April 9, 2008, from 1 p.m. to 4 p.m., Eastern Time.**ADDRESSES:** Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 4090. Please bring photo ID for entry to a Federal building.**FOR FURTHER INFORMATION CONTACT:** <http://www.hhs.gov/healthit/ahic/chroniccare/>**SUPPLEMENTARY INFORMATION:** The workgroup will hear testimony on ways to use information technology to better coordinate care for patients with chronic conditions and will discuss this information in light of opportunities to better facilitate patient care coordination.The meeting will be available via Web cast. For additional information, go to: <http://www.hhs.gov/healthit/ahic/chroniccare/cc-instruct.html>.

Dated: March 6, 2008.

**Judith Sparrow,***Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.*

[FR Doc. E8-5160 Filed 3-17-08; 8:45 am]

**BILLING CODE 4150-45-M****DEPARTMENT OF HEALTH AND HUMAN SERVICES****Office of the National Coordinator for Health Information Technology; American Health Information Community Personalized Healthcare Workgroup Meeting****ACTION:** Announcement of meeting.**SUMMARY:** This notice announces the 14th meeting of the American Health Information Community Personalized Healthcare Workgroup in accordance with the Federal Advisory Committee Act (Pub. L. No. 92-463, 5 U.S.C., App.).**DATES:** April 7, 2008, from 1 p.m. to 4 p.m. [Eastern Time].**ADDRESSES:** Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 4090. Please bring photo ID for entry to a Federal building.**FOR FURTHER INFORMATION CONTACT:** <http://www.hhs.gov/healthit/ahic/healthcare/>**SUPPLEMENTARY INFORMATION:** The Workgroup will discuss possible common data standards to incorporate interoperable, clinically useful genetic/genomic information and analytical tools into Electronic Health Records (EHRs) to support clinical decision-making for the clinician and consumer.The meeting will be available via Web cast. For additional information, go to: [http://www.hhs.gov/healthit/ahic/healthcare/phc\\_instruct.html](http://www.hhs.gov/healthit/ahic/healthcare/phc_instruct.html).

Dated: March 6, 2008.

**Judith Sparrow,***Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.*

[FR Doc. E8-5161 Filed 3-17-08; 8:45 am]

**BILLING CODE 4150-45-M****DEPARTMENT OF HEALTH AND HUMAN SERVICES****Office of the National Coordinator for Health Information Technology; American Health Information Community Population Health and Clinical Care Connections Workgroup Meeting****ACTION:** Announcement of meeting.**SUMMARY:** This notice announces the 25th meeting of the American Health Information Community Population Health and Clinical Care Connections Workgroup in accordance with the Federal Advisory Committee Act (Pub. L. No. 92-463, 5 U.S.C., App.)**DATES:** April 3, 2008, from 1 p.m. to 4 p.m. (Eastern Time).**ADDRESSES:** Mary C. Switzer Building (330 C Street, SW., Washington, DC 20201), Conference Room 4090 (please bring photo ID for entry to a Federal building).**FOR FURTHER INFORMATION CONTACT:** <http://www.hhs.gov/healthit/ahic/population/>.**SUPPLEMENTARY INFORMATION:** The Workgroup will continue its discussion on how to facilitate the flow of reliable health information among population health and clinical care systems necessary to protect and improve the public's health.The meeting will be available via Web cast. For additional information, go to: [http://www.hhs.gov/healthit/ahic/population/pop\\_instruct.html](http://www.hhs.gov/healthit/ahic/population/pop_instruct.html).

Dated: March 6, 2008.

**Judith Sparrow,***Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.*

[FR Doc. E8-5162 Filed 3-17-08; 8:45 am]

**BILLING CODE 4150-45-M****DEPARTMENT OF HEALTH AND HUMAN SERVICES****Centers for Disease Control and Prevention****Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): National Institute for Occupational Safety and Health (NIOSH): Occupational Safety and Health Training Project Grants Announcement for Research (PAR) 06-484**

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC) announces the aforementioned meeting.

*Time and Date:* 1 p.m.-3:30 p.m., April 1, 2008 (Closed).*Place:* NIOSH, 2400 Century Parkway, Conference Room 4211-NIOSH-2, Atlanta, GA 30345.*Status:* The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c) (4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92-463.*Matters to be Discussed:* The meeting will include the review, discussion, and evaluation of "NIOSH Occupational Safety and Health Training Project Grants, PAR 06-484."*Contact Person for More Information:* Charles N. Rafferty, Ph.D., Assistant Director for Review and Policy Office of Extramural Program, Office of Extramural Coordination and Special Projects, NIOSH, CDC, 2400 Century Parkway, NE., Atlanta, GA 30345, Telephone (404) 498-2500.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.

Dated: March 12, 2008.

**Diane Allen,**

*Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.*

[FR Doc. E8-5376 Filed 3-17-08; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2008-N-0157 (formerly 2007N-0105)]

#### Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Mental Models Study of Food Terrorism Risk Awareness

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Mental Models Study of Food Terrorism Risk Awareness" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:**

Jonna Capezzuto, Office of the Chief Information Officer (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-4659.

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of July 24, 2007 (72 FR 40309), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0618. The approval expires on February 28, 2011. A copy of the supporting statement for this information collection is available on the Internet at <http://www.reginfo.gov/public/do/PRAMain>.

Dated: March 10, 2008.

**Jeffrey Shuren,**

*Assistant Commissioner for Policy.*

[FR Doc. E8-5361 Filed 3-17-08; 8:45 am]

BILLING CODE 4160-01-S

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2008-N-0162]

#### Agency Information Collection Activities; Proposed Collection; Comment Request; Prescription Drug Product Labeling; Medication Guide Requirements

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on regulations requiring the distribution of patient labeling, called Medication Guides, for certain products that pose a serious and significant public health concern requiring distribution of FDA-approved patient medication.

**DATES:** Submit written or electronic comments on the collection of information by May 19, 2008.

**ADDRESSES:** Submit electronic comments on the collection of information to <http://www.regulations.gov>. Submit written comments on the collection of information to Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:**

Elizabeth Berbakos, Office of the Chief Information Officer (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1482.

**SUPPLEMENTARY INFORMATION:** Under the PRA (44 U.S.C. 3501-3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or

provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumption used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

#### Prescription Drug Product Labeling; Medication Guide Requirements (OMB Control Number 0910-0393)—Extension

FDA regulations require the distribution of patient labeling, called Medication Guides, for certain prescription human drug and biological products used primarily on an outpatient basis that pose a serious and significant public health concern requiring distribution of FDA-approved patient medication information. These Medication Guides inform patients about the most important information they should know about these products in order to use them safely and effectively. Included is information such as the drug's approved uses, contraindications, adverse drug reactions, and cautions for specific populations, with a focus on why the particular product requires a Medication Guide. These regulations are intended to improve the public health by providing information necessary for patients to use certain medications safely and effectively.

The regulations contain the following reporting requirements that are subject to the PRA. The estimates for the burden hours imposed by the following regulations are listed in table 1 of this document:

- 21 CFR 208.20—Applicants must submit draft Medication Guides for FDA