

FDA's burden estimate is based on prior experience with Internet panel experiments similar to the study proposed here. Sixty panel members will take part in a pre-test of the study, estimated to last 30 minutes (0.5 hours), for a total of 30 hours. Approximately 15,000 respondents will complete a screener to determine eligibility for participation in the study, estimated to take 1 minute (0.016 hours), for a total of 125 hours. Fifty-four hundred (5,400) respondents will complete the full study, estimated to last 30 minutes (0.5 hours), for a total of 2,700 hours. The total estimated burden is 2,970 hours.

Dated: February 16, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010-3320 Filed 2-19-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention (CDC): Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92-463) of October 6, 1972, that the Advisory Committee to the Director, Centers for Disease Control and Prevention of the Department of Health and Human Services, has been renewed for a 2-year period extending through February 1, 2012.

Contact Person for More Information: Anne C. Haddix, PhD, Designated Federal Officer, ACD, CDC, 1600 Clifton Road, NE., M/S D14, Atlanta, Georgia 30333. Telephone 404-639-0663.

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 the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: February 11, 2010.

Andre Tyler,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control

Special Emphasis Panel (SEP): Healthy Passages Longitudinal Study of Youth, Funding Opportunity Announcement (FOA) DP 10-007, Initial Review.

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 p.m., April 20, 2010 (Closed).

Place: Teleconference.

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 initial review, discussion, and evaluation of applications received in response to "Healthy Passages Longitudinal Study of Youth, FOA DP 10-007."

Contact Person for More Information: Michael Dalmat, DRPH., Scientific Review Officer, National Center for Chronic Disease and Health Promotion, Office of the Director, Extramural Research Program Office, 4770 Buford Highway, NE., Mailstop K-92, Atlanta, GA 30341, Telephone: (770) 488-6423, E-mail: MED1@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.

Dated: February 10, 2010.

Elaine L. Baker,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-1996-N-0006] (formerly Docket No. 1996N-0277)

Safety and Efficacy Review for Additional Ingredients in Over-the-Counter Drug Products for Human Use; Request for Environmental Impact Data and Information

AGENCY: Food and Drug Administration, HHS.

ACTION: Request for data and information.

SUMMARY: We (Food and Drug Administration (FDA)) are requesting data and information regarding the potential environmental impact of amending over-the-counter (OTC) drug monographs to include certain active ingredients not previously marketed in the United States or marketed in the United States under approved applications after the OTC drug review began in 1972. Thirteen active ingredients have been found eligible for potential inclusion in OTC drug monographs based on time and extent applications (TEAs). We are currently evaluating the safety and effectiveness of these ingredients.

DATES: Submit data, information, and general comments by May 24, 2010.

ADDRESSES: Submit electronic or written data, information, and general comments in response to this document. Submit electronic comments to <http://regulations.gov>. Submit written comments to the Division of Dockets Management (HFA-305), 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Michael L. Koenig, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, rm. 5411, Silver Spring, MD 20993-0002, 301-796-2090.

SUPPLEMENTARY INFORMATION:

I. Ingredients Affected by This Notice

We are currently evaluating the safety and effectiveness of 13 active ingredients found eligible for possible addition to an OTC drug monograph via the TEA process described in 21 CFR 330.14. The ingredients under review are shown in table 1 of this document: