Submission of Nominations for the Evaluation Set 22

Proposed Substances: Today's notice also invites voluntary public nominations for substances not listed in this notice. Nominations are most useful if they include the nominator, including full name, title, affiliation, email address, and telephone number.

ATSDR will evaluate all data and information associated with nominated substances and will determine the final list of substances that will be chosen for toxicological profile development. Substances will be chosen according to ATSDR's specific guidelines for selection, found in the *Selection Criteria* announced in the **Federal Register** on May 7th, 1993 (87 FR 27288).

Please submit nominations by one of the following methods:

- *E-mail: jxt1@cdc.gov*.
- *Fax:* 770.488.4178.
- *Mail:* CDR Jessilynn Taylor, 1600 Clifton Rd, NE., MS F32, Atlanta, GA, 30333.

Please ensure that your comments are submitted within the specified nomination period. Nominations received after the closing date will be marked as late and may be considered only if time permits.

Dated: October 19, 2007.

Ken Rose,

Director, Office of Policy, Planning and Evaluation, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry.

[FR Doc. E7–21018 Filed 10–24–07; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 1999N–2337 (formerly Docket No. 99N–2337)]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; CGMP for Blood and Blood Components; Notification of Consignees and Transfusion Recipients Receiving Blood and Blood Components at Increased Risk of Transmitting HCV Infection (''Lookback'')

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''CGMP for Blood and Blood Components; Notification of Consignees and Transfusion Recipients Receiving Blood and Blood Components at Increased Risk of Transmitting HCV Infection (''Lookback'')'' 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 August 24, 2007 (72 FR 48766), 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 0910–0610. The approval expires on October 31, 2010. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: October 19, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–21055 Filed 10–24–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0278]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry on Continuous Marketing Applications: Pilot—Scientific Feedback and Interactions During Development of Fast Track Products Under the Prescription Drug User Fee Act

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled '' Guidance for Industry on Continuous Marketing Applications: Pilot—Scientific Feedback and Interactions During Development of Fast Track Products Under the Prescription Drug User Fee Act'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4816.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of May 21, 2007 (72 FR 28495), 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 0910–0518. The approval expires on September 30, 2010. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: October 19, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–21056 Filed 10–24–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N-0390]

User Fee Program for Advisory Review of Direct-to-Consumer Television Advertisements for Prescription Drug and Biological Products; Request for Notification of Participation and Number of Advertisements for Review

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for notification of participation.

SUMMARY: The Food and Drug Administration (FDA) is issuing this notice to explain the new direct-to-consumer (DTC) user fee program (DTC user fee program) established by the Food and Drug Administration Amendments Act of 2007 (FDAAA) and, as required by the new law, to ask companies to notify FDA within 30 calendar days if they intend to participate in the DTC user fee program during fiscal year (FY) 2008 and, if they do plan to participate, to identify the number of DTC television advertisements for prescription drug and biological products they plan to