other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: New collection;

Title of Information Collection: External Evaluation of National Centers of Excellence in Women's Health Program;

Form/OMB No.: 0990–New;

Use: This evaluation will assess the effectiveness of the National Centers of Excellence in Women's Health Program contracts. Specifically, the outcomes-based research question will ask: ''Overall, do women seen at CoEs perceive their clinical care to be better than those who receive healthcare elsewhere?'' Outreach surveys will assess community impacts.

Frequency: One-time on Occasion;

Affected Public: Individuals or Households;

Annual Number of Respondents: 6800;

Total Annual Responses: 6800;

Average Burden per Response: 12.65 minutes;

Total Annual Hours: 1433;

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, e-mail your request, including your address, phone number, OMB number, and OS document identifier, to *Sherette.funncoleman@hhs.gov,* or call the Reports Clearance Office on (202) 690–6162. Written comments and recommendations for the proposed information collections must be received within 30 days of this notice directly to the Desk Officer at the address below: OMB Desk Officer: John Kraemer, OMB Human Resources and Housing Branch, Attention: (OMB #0990–New), New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: February 9, 2007.

Alice Bettencourt,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

[FR Doc. E7–2769 Filed 2–15–07; 8:45 am]

BILLING CODE 4150–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Council for the Elimination of Tuberculosis Meeting (ACET)

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 following council meeting of the aforementioned committee.

Times and Dates: 8:30 a.m.–5 p.m., March 20, 2007. 8:30 a.m.–2 p.m., March 21, 2007.

Place: Corporate Square, Building 8, 1st Floor Conference Room, Atlanta, Georgia 30333, telephone (404) 639–8317.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 100 people.

Purpose: This council advises and makes recommendations to the Secretary, Health and Human Services, the Assistant Secretary for Health, and the Director, CDC, regarding the elimination of tuberculosis. Specifically, the Council makes recommendations regarding policies, strategies, objectives, and priorities; addresses the development and application of new technologies; and reviews the extent to which progress has been made toward eliminating tuberculosis.

Matters to be Discussed: Agenda items include issues pertaining to TB Disparities in African Americans; Response to Control Extensively Drug Resistant Tuberculosis in the U.S.; and U.S.-Mexico Border Issues and other related tuberculosis issues.

Agenda items are subject to change as priorities dictate.

FOR FURTHER INFORMATION CONTACT: Margie Scott-Cseh, National Center for HIV, STD, and TB Prevention, 1600 Clifton Road, NE., M/S E–07, Atlanta, Georgia 30333, telephone (404) 639–8317.

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.

Elaine L. Baker,

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

[FR Doc. E7–2766 Filed 2–15–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Infectious Diseases (NCID): Meeting

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 following meeting of the aforementioned committee.

Times and Dates: 9 a.m.–5:30 p.m., March 15, 2007; 8:30 a.m.–2:30 p.m., March 16, 2007.

Place: CDC, Building 19, 1600 Clifton Road, N.E., Atlanta, Georgia 30333.

Status: Open to the public, limited only by the space available.

Purpose: The Board of Scientific Counselors, NCID, provides advice and guidance to the Director, CDC, and Director, NCID, in the following areas: program goals and objectives; strategies; program organization and resources for infectious disease prevention and control; and program priorities.

Matters To Be Discussed: Agenda items will include:

1. Breakout Group Discussions:

Vaccine Preventable Disease Surveillance

Respiratory Outbreaks

Ecology of Emerging Zoonoses and Other Infectious Diseases

Food-Borne Diseases: The New Way Forward

Working with Academic Partners

Chlamydia Screening Programs in the U.S.

Antimicrobial Resistance

Infectivity Component of Infection Control

Exotic Animal Importation and Trade

2. Coordinating Center for Infectious Diseases Updates

3. Strategies for Identifying New Pathogens

Other agenda items include announcements/introductions; follow-up on actions recommended by the Board in May 2006; consideration of future directions, goals, and recommendations.

Agenda items are subject to change as priorities dictate.

Written comments are welcome and should be received by the contact person listed below prior to the opening of the meeting.

FOR FURTHER INFORMATION CONTACT: Tony Johnson, Office of the Director, NCID, CDC, Mailstop A–45, 1600 Clifton Road, NE, Atlanta, Georgia 30333, e-mail *tjohnson3@cdc.gov*; telephone 404/639–3856.

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.

Elaine L. Baker,

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

[FR Doc. E7–2753 Filed 2–15–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0274]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Establishing and Maintaining a List of United States Dairy Product Manufacturers/Processors With Interest in Exporting to Chile

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Establishing and Maintaining a List of U.S. Dairy Product Manufacturers/Processors with Interest in Exporting to Chile'' 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 December 7, 2006 (71 FR 70972), 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 0910–0509. The approval expires on January 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: February 9, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–2708 Filed 2–15–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0435]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on How to Use E-Mail to Submit a Notice of Intent to Slaughter for Human Food Purposes

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by March 19, 2007.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance:

Guidance for Industry on ''How to Use E-mail to Submit a Notice of Intent to Slaughter for Human Food Purposes,'' Section 512j, Federal Food, Drug, and Cosmetic Act; (OMB Control Number 0910–0450)—Extension

Section 512(j) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(j)) gives FDA the authority to set conditions under which animals treated with investigational new animal drugs may be marketed for food use. Under this authority, the Center for Veterinary Medicine (CVM), issues to a new animal drug sponsor (sponsors) a slaughter authorization letter that sets the terms under which investigational animals may be slaughtered. The United States Department of Agriculture (USDA) also monitors the slaughter of animals treated with investigational new animal drugs under the authority of the Meat Inspection Act (21 USC 601–95). Sponsors must submit slaughter notices each time investigational animals are presented for slaughter, unless this requirement is waived by an authorization letter (21 CFR 511.1(b)(5), 9 CFR 309.17). These notifications assist CVM and USDA in monitoring the safety of the food supply. Slaughter notices were previously submitted to CVM and USDA on paper (OMB No. 0910–0450). CVM's guidance on ''How to Use E-Mail to Submit a Notice of Intent to Slaughter for Human Food Purposes'' provides sponsors with the option to submit a slaughter notice as an e-mail attachment to CVM and USDA via the Internet. The electronic submission of slaughter notices is part of CVM's ongoing initiative to provide a method for paperless submissions.

In the **Federal Register** of November 8, 2006 (71 FR 65532), FDA published a 60-day notice soliciting comments on the information collection provisions of this collection. In response to this notice, no comments were received.

The likely respondents for this collection of information are new animal drug sponsors.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

Form No.	No. of Respondents	Annual Frequency per Response	Total Annual Responses[2]	Hours per Response	Total Hours
FDA Form #3488	25	.08	2	0.41	.82

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.