Secretary, ACIPC, NCIPC, CDC, 4770 Buford Highway, NE., M/S K02, Atlanta, Georgia 30341–3724, telephone 770/ 488–4694.

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: April 3, 2003.

Alvin Hall,

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

[FR Doc. 03–8744 Filed 4–9–03; 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: 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 committee meeting.

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

Times and Dates: 9 a.m.–5 p.m., May 1, 2003. 8:30 a.m.–3:30 p.m., May 2, 2003.

Place: CDC, Auditorium B, Building 1, 1600 Clifton Road, NE., 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 Secretary, the Assistant Secretary for Health, 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. Opening Session: NCID Update

a. Severe Acute Respiratory Syndrome b. Methicillin-Resistant Staphylococcus

Aureus

c. Malaria

2. Bioterrorism Update

3. Institute of Medicine Emerging

Infections Report

4. Global Health Activities

5. Emerging Infectious Diseases Journal

6. Updates

a. Pneumococcal Disease

c. Hepatitis

7. Infections and Chronic Diseases

8. Board meets with Director, CDC9. Discussions and Recommendations

Other agenda items include

announcements/introductions; follow-up on actions recommended by the Board in December 2002; 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.

Contact Person for More Information: Tony Johnson, Office of the Director, NCID, CDC, Mailstop E–51, 1600 Clifton Road, NE., Atlanta, Georgia 30333, e-mail

tjohnson3@cdc.gov; telephone 404/498–3249. 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: April 3, 2003.

Alvin Hall,

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

[FR Doc. 03–8737 Filed 4–9–03; 8:45 am] BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Institute for Occupational Safety and Health; Meeting Notice

The National Institute for Occupational Safety and Health (NIOSH) at the Centers for Disease Control and Prevention announces the following meeting:

Name: Continue Conceptual Discussions for Escape Respirator Standards Development Efforts Used for Respiratory Protection Against Chemical, Biological, Radiological, and Nuclear (CBRN) Agents.

Time and Date: 9 a.m.—5 p.m., April 29, 2003.

Place: Radisson Hotel Pittsburgh Green Tree, 101 Radisson Drive, Pittsburgh, Pennsylvania.

Status: This meeting is hosted by NIOSH and will be open to the public, limited only by the space available. The meeting room will accommodate approximately 175 people. Interested parties should make hotel reservations directly with the Radisson Hotel Pittsburgh Green Tree (412–922–8400 or 800–333–3333) before the cut off date of April 21, 2003, referencing the NIOSH/ NPPTL Public Meeting. Interested parties should confirm their attendance to this meeting by completing a registration form and forwarding it by e-mail (confserv@netl.doe.gov) or fax (304–285–

(*confserv@net1.doe.gov*) or fax (304–285– 4459) to the Event Management Office. A registration form may be obtained from the NIOSH Homepage (*http://www.cdc.gov/niosh*) by selecting "Conferences," and then the event.

Requests to make presentations at the public meeting should be mailed to the NIOSH Docket Officer, Robert A. Taft Laboratories, M/S C34, 4676 Columbia Parkway, Cincinnati, Ohio 45226, Telephone 513–533–8303, Fax 513–533–8285, F-mail *niocindocket@cdc.gov*. All requests to present should contain the name, address, telephone number, relevant business affiliations of the presenter, a brief summary of the presentation, and the approximate time requested for the presentation. Oral presentations should be limited to 15 minutes.

After reviewing the requests for presentations, NIOSH will notify each presenter of the approximate time that their presentation is scheduled to begin. If a participant is not present when his or her presentation is scheduled to begin, the remaining participants will be heard in order. At the conclusion of the meeting, an attempt will be made to allow presentations by any scheduled participants who missed their assigned times. Attendees who wish to speak but did not submit a request for the opportunity to make a presentation may be given this opportunity at the conclusion of the meeting, at the discretion of the presiding officer. Comments on the topics presented in this notice and at the meeting should be mailed to the NIOSH Docket Office, Robert A. Taft Laboratories, M/S C34, 4676 Columbia Parkway, Cincinnati, Ohio 45226, Telephone 513-533-8303, Fax 513-533-8285. Comments may also be submitted by e-mail to niocindocket@cdc.gov. E-mail attachments should be formatted as WordPerfect 6/7/8/9 or Microsoft Word. Comments should be submitted to NIOSH no later than May 30, 2003, and should reference docket number, NIOSH-002, in the subject heading.

Purpose: NIOSH will continue conceptual discussions of standards and testing processes for an Escape Respirator standard suitable for respiratory protection against CBRN Agents and review ongoing research to identify simulata materials for use as CBRN test surrogates for respirator research. NIOSH, along with the U.S. Army Soldier and Biological Chemical Command (SBCCOM) and the National Institute for Standards and Technology (NIST), will present information to attendees concerning the concept development for the Escape Respirator CBRN standard. Participants will be given an opportunity to ask questions and to present individual comments for consideration. Interested participants may obtain the latest copy of the Escape Respirator CBRN concept paper, as well as earlier versions of the concept papers used during the standard development effort, from the NIOSH contact identified below, or from the NIOSH National Personal Protective Technology Laboratory (NPPTL) Web site, address: http://www.cdc.gov/niosh/npptl. The April 15, 2003, concept paper will be used as the basis for discussion at the public meeting, as well as forming the basis for the new Escape Respirator CBRN statement of standard.

Recent acts of terrorism have created an urgent awareness of domestic security and preparedness issues. Municipal, state, and federal responder groups, particularly those in locations considered potential targets, have been developing and modifying response and consequence management plans. Since the World Trade Center and anthrax incidents, most emergency response agencies have operated with a heightened appreciation of the potential scope and sustained resource requirements for coping with such events. The Federal Interagency Board for Equipment Standardization and Interoperability (IAB) has worked to identify personal protective equipment that is already available on the market for responders' use. The IAB has identified the development of standards or guidelines for respiratory protection equipment as a top priority. NIOSH, NIST, National Fire Protection Association, and the Occupational Safety and Health Administration have entered into a Memorandum of Understanding defining each agency's or organization's role in developing, establishing, and enforcing standards or guidelines for responders' respiratory protective devices. NIST has initiated Interagency Agreements with NIOSH and SBCCOM to aid in the development of appropriate protection standards or guidelines. NIOSH has the lead in developing standards or guidelines to test, evaluate, and approve respirators.

NIOSH, SBCCOM, and NIST have hosted public meetings on April 17 and 18, 2001; June 18 and 19, 2002; and October 16 and 17, 2002, presenting their progress in assessing respiratory protection needs of responders to CBRN incidents. The methods or models for developing hazard and exposure estimates, and the status of evaluating test methods and performance standards that may be applicable in future CBRN respirator standards or guidelines, were discussed at these meetings.

Contact for Additional Information: Event Management, P.O. Box 880, 3610 Collins Ferry Road, Morgantown, WV 26507, Telephone 304–285–4750, Fax 304–285– 4459, E-mail confserv@netl.doe.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 the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: April 3, 2003.

Alvin Hall,

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

[FR Doc. 03–8738 Filed 4–9–03; 8:45 am] BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Institute for Occupational Safety and Health: 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 committee meeting:

Name: Board of Scientific Counselors, National Institute for Occupational Safety and Health (BSC, NIOSH).

Time and Date: 9 a.m.–2:45 p.m., April 30, 2003.

Place: The Washington Court, 525 New Jersey Avenue, NW., Washington, DC 20001– 1527, telephone 202/879–7918, fax 202/879– 7918.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people.

Purpose: The BSC, NIOSH, is charged with providing advice to the Director, NIOSH, on NIOSH research programs. Specifically, the Board provides guidance on the Institute's research activities related to developing and evaluating hypotheses, systematically documenting findings, and disseminating results.

Matters to be Discussed: Agenda items include a report from the Director of NIOSH; Report on NIOSH International Activities; update on the National Exposure at Work Survey; briefing on Outreach and Information for Small Businesses; update on NIOSH Occupational Asthma Research; closing remarks.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Roger Rosa, Ph.D., Executive Secretary, BSC, NIOSH, Centers for Disease Control and Prevention, 200 Independence Avenue, SW., Room 715H,Washington, DC 20201, telephone: 202/205–7856, fax: 202/260–4464, e-mail: *rrosa@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 the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: April 3, 2003.

Alvin Hall,

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

[FR Doc. 03–8752 Filed 4–9–03; 8:45 am] BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 03N-0135]

Agency Emergency Processing Under OMB Review; Guidance: Establishing and Maintaining a List of U.S. Dairy Product Manufacturers 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 proposed collection of information has been submitted to the Office of Management and Budget (OMB) for emergency processing under the Paperwork Reduction Act of 1995 (the PRA). FDA is preparing a guidance document intended to notify the public of procedures being implemented by the agency to assist U.S. firms that wish to export dairy products to Chile. FDA is taking this action in response to trade discussions with Chile that have been adjunct to the negotiations of the United States-Chile Free Trade Agreement. FDA is requesting this emergency processing under the PRA because a normal clearance is likely to impede completion of the United States-Chile Free Trade Agreement.

DATES: Fax or electronically mail written comments on the collection of information by May 12, 2003. **ADDRESSES:** OMB is still experiencing significant delays in the regular mail, including first class and express mail, and messenger deliveries are not being accepted. To ensure that comments on the information collection are received, OMB recommends that written comments be electronically mailed to sshapiro@omb.eop.gov or faxed to the Office of Information and Regulatory Affairs, OMB, Attn: Stuart Shapiro, Desk Officer for FDA, FAX 202-395-6974. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Peggy Robbins, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: FDA is preparing a guidance document intended to notify the public of procedures being implemented by the agency to assist U.S. firms that wish to export dairy products to Chile. FDA is taking this action in response to trade