Status: Open to the public, teleconference access limited only by availability of telephone ports.

Purpose: Under the charge of the Board of Scientific counselors (BSC), NCEH/ATSDR the Program Peer Review Subcommittee establishes and monitors working groups of technical experts that perform program peer reviews of National Center for Environmental Health and the Agency for Toxic Substances and Disease Registry. The Subcommittee, working with the NCEH/ATSDR, Office of Science (OS), will establish a schedule and process for program peer reviews, nominate working group members, review summary reports and recommendations, and report back to the BSC. The OS will establish agency policy for program peer review and directly support each working group by collating program documents, and organizing the working groups review and site visit. Each NCEH/ATSDR program eligible for review will be reviewed every 5 years according to CDC/ATSDR policy.

Matters to be Discussed: The teleconference agenda will include a review of action items from the previous meeting, discussion and updates on the program peer review process and an update on the Hazards Substances Emergency Events Surveillance System.

SUPPLEMENTARY INFORMATION: This conference call is scheduled to begin at 12:30 p.m. eastern standard time. To participate in the teleconference, please dial (877) 315–6535 and enter conference code 383520.

FOR FURTHER INFORMATION CONTACT: Drue Barrett, Ph.D., Executive Secretary, PRRS, NCEH/ATSDR, M/S E–28, 1600 Clifton Road, NE., Atlanta, Georgia 30333, telephone (404) 498–0003.

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: January 10, 2005.

Alvin Hall,

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

[FR Doc. 05–782 Filed 1–13–05; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Institute for Occupational Safety and Health Advisory Board on Radiation and Worker Health

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: Advisory Board on Radiation and Worker Health (ABRWH), and Subcommittee for Dose Reconstruction and Site Profile Reviews, National Institute for Occupational Safety and Health (NIOSH).

Subcommittee Meeting Time and Date: 8:30 a.m.–12 p.m., February 7, 2005.

Committee Meeting Times and Dates: 1 p.m.–5 p.m., February 7, 2005. 8 a.m.–4:45 p.m., February 8, 2005. 7 p.m.–8:30 p.m., February 8, 2005. 8:30 a.m.–4:30 p.m., February 9, 2005.

Place: Adam's Mark St. Louis, 4th and Chestnut Street, St. Louis, Missouri 63102, telephone 314–241–7400, fax 314–241–9839.

Status: Open to the public, limited only by the space available. The meeting space accommodates approximately 500 people.

Background: The ABRWH was established under the Energy Employees Occupational Illness Compensation Program Act (EEOICPA) of 2000 to advise the President, delegated to the Secretary of Health and Human Services (HHS), on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Board include providing advice on the development of probability of causation guidelines which have been promulgated by HHS, as a final rule, advice on methods of dose reconstruction which have also been promulgated by HHS, as a final rule, advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation program, and advice on petitions to add classes of workers to the Special Exposure Cohort (SEC).

In December 2000 the President delegated responsibility for funding, staffing, and operating the Board to HHS, which subsequently delegated this authority to the CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, and renewed on August 3, 2003.

Purpose: This board is charged with (a) providing advice to the Secretary, HHS on the development of guidelines under Executive Orders 13179; (b) providing advice to the Secretary, HHS on the scientific validity and quality of dose reconstruction efforts performed for this Program; and (c) upon request by the Secretary, HHS, advise the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class.

Matters To Be Discussed: Agenda for this meeting will focus on Program Status Reports from NIOSH and the Department of Labor; Site Profile Review of Bethlehem Steel; Task 3 Procedures Review; Site Profile Review of Mallinckrodt (Destrehan Street Facility); Travel Policy; Status Report of SC&A Task Orders and Costs; SEC Petition Evaluation Report—Mallinckrodt to include NIOSH Reports and Recommendations and Petitioners Comments on Report; Subcommittee Report & Board Discussion on First Set of Case Reviews; SEC Petition Evaluation Report—Iowa Army Ammunition Plant (IAAP) to include NIOSH Reports and Recommendations and Petitioners Comments on Report; and Board working sessions. There will be an evening public comment period scheduled for February 8, 2005, and public comment periods on all meeting days.

The Subcommittee for Dose Reconstruction and Site Profile Reviews will convene on February 7, 2005, from 8:30 a.m.–12 p.m. and will focus on review of draft minutes; discussion of Case Sampling Matrix, Summary of First Set of Case Reviews/ Preparation of Recommendation for Full Board and selection of Third Set of Individual Dose Reconstruction Cases for Board Review.

The agenda is subject to change as priorities dictate. In the event an individual cannot attend, written comments may be submitted. Any written comments received will be provided at the meeting and should be submitted to the contact person below well in advance of the meeting.

FOR FURTHER INFORMATION CONTACT: Larry Elliott, Executive Secretary, ABRWH, NIOSH, CDC, 4676 Columbia Parkway, Cincinnati, Ohio 45226, telephone 513/533–6825, fax 513/533–6826.

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: January 10, 2005.

Alvin Hall,

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

[FR Doc. 05–781 Filed 1–13–05; 8:45 am]

BILLING CODE 4163–19–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Citizens Advisory Committee on Public Health Service Activities and Research at Department of Energy (DOE) Sites: Savannah River Site Health Effects Subcommittee (SRSHES)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Agency for Toxic Substances and Disease Registry (ATSDR) and the Centers for Disease Control and Prevention (CDC) announce the following meeting.

Name: Citizens Advisory Committee on Public Health Service Activities and

Research at Department of Energy (DOE) Sites: Savannah River Site Health Effects Subcommittee (SRSHES).

Time and Date: 8 a.m.–12:30 p.m., January 25, 2005.

Place: Augusta Towers Hotel & Convention Center, 2651, Perimeter Parkway, Augusta, GA 30909, telephone 706–855–8100, fax 706–860–7334.

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

Background: Under a Memorandum of Understanding (MOU) signed in December 1990 with DOE, and replaced by MOUs signed in 1996 and 2000, the Department of Health and Human Services (HHS) was given the responsibility and resources for conducting analytic epidemiologic investigations of residents of communities in the vicinity of DOE facilities, workers at DOE facilities, and other persons potentially exposed to radiation or to potential hazards from non-nuclear energy production use. HHS delegated program responsibility to CDC.

In addition, a memo was signed in October 1990 and renewed in November 1992, 1996, and in 2000, between ATSDR and DOE. The MOU delineates the responsibilities and procedures for ATSDR's public health activities at DOE sites required under sections 104, 105, 107, and 120 of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA or ''Superfund''). These activities include health consultations and public health assessments at DOE sites listed on, or proposed for, the Superfund National Priorities List and at sites that are the subject of petitions from the public; and other health-related activities such as epidemiologic studies, health surveillance, exposure and disease registries, health education, substance-specific applied research, emergency response, and preparation of toxicological profiles.

Purpose: This subcommittee is charged with providing advice and recommendations to the Director of CDC and the Administrator of ATSDR pertaining to CDC's and ATSDR's public health activities and research at this DOE site. The purpose of this meeting is to provide a forum for community, American Indian Tribal, and labor interaction, and to serve as a vehicle for communities, American Indian Tribes, and labor to express concerns and provide advice and recommendations to CDC and ATSDR.

Matters To Be Discussed: Agenda items include a presentation on Radiation Epidemiology from the National Center for Environmental Health (NCEH), CDC, and a Subcommittee discussion on the Advanced Technologies and Laboratories International, Inc., final report.

Agenda items are subject to change as priorities dictate.

Inability to confirm attendance of quorum prevented publication 15 days prior to the meeting.

FOR FURTHER INFORMATION CONTACT: Mr. Phillip Green, Executive Secretary, SRSHES, Radiation Studies Branch, Division of Environmental Hazards and Health Effects, National Center for Environmental Health, CDC, 1600 Clifton Road, NE. (E–39), Atlanta, Georgia 30333, telephone (404) 498–1800, fax (404) 498–1811.

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 ATSDR.

Dated: January 10, 2004.

Alvin Hall,

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

[FR Doc. 05–784 Filed 1–13–05; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

Privacy Act of 1974; Report of New System

AGENCY: Department of Health and Human Services (HHS), Centers for Medicare & Medicaid Services (CMS).

ACTION: Notice of New System of Records (SOR).

SUMMARY: In accordance with the requirements of the Privacy Act of 1974, we are proposing to establish a new system of records, called the ''Cytology Personnel Record System (CYPERS), HHS/CMS/CMSO, 09–70–0543.'' The primary purpose of CYPERS is to assure CMS of the accuracy and reliability of gynecologic cytology testing by compliance with the CLIA statutory requirements. This will be accomplished by tracking and monitoring the enrollment, participation, and performance of individual cytotechnologists and physicians participating in CMS approved gynecologic cytology proficiency testing programs.

Information retrieved from this system of records will be used to support regulatory, reimbursement, and policy functions performed within the agency or by a contractor or consultant; support constituent requests made to a Congressional representative; and support litigation involving the agency.

We have provided background information about the proposed system in the **SUPPLEMENTARY INFORMATION** section, below. Although the Privacy Act requires only that the ''routine use'' portion of the system be published for comment, CMS invites comments on all portions of this notice. See **EFFECTIVE DATES** section for comment period.

EFFECTIVE DATES: CMS filed a new system report with the Chair of the House Committee on Government Reform and Oversight, the Chair of the Senate Committee on Governmental Affairs, and the Administrator, Office of Information and Regulatory Affairs, Office of Management and Budget (OMB) on December 23, 2004. In any event, we will not disclose any information under a routine use until forty (40) calendar days after publication. We may defer implementation of this system of records or one or more of the routine use statements listed below if we receive comments that persuade us to defer implementation.

ADDRESSES: The public should address comments to: Director, Division of Privacy Compliance Data Development (DPCDD), CMS, Room N2–04–27, 7500 Security Boulevard, Baltimore, Maryland 21244–1850. Comments received will be available for review at this location, by appointment, during regular business hours, Monday through Friday from 9 a.m.–3 p.m., eastern time zone.

FOR FURTHER INFORMATION CONTACT:

David Escobedo, Finance, Systems and Budget Group, Center for Medicaid and State Operations, Centers for Medicare and Medicaid Services, 7500 Security Boulevard, Room S3–18–11, Baltimore, Maryland 21244–1850, Telephone Number: (410) 786–5401.

Thomas Hamilton, Survey and Certification Group, Center for Medicaid and State Operations, Centers for Medicare and Medicaid Services, 7500 Security Boulevard, Room S2–12–25, Baltimore, Maryland 21244–1850, Telephone Number: (410) 786–9493.

SUPPLEMENTARY INFORMATION:

I. Description of the New System of Records

A. Statutory and Regulatory Basis for System of Records

Section 353(f)(4)(A) of the Public Health Service Act (42 U.S.C. 263a) mandates that the Secretary establish national standards for quality assurance in cytology services designed to assure consistent, valid, and reliable test performance by cytology laboratories. Section 353(f)(4)(B)(iv) requires, ''* * * the periodic confirmation and evaluation of the proficiency of individuals involved in screening or interpreting cytological preparations, including announced and unannounced