DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Extended Work Schedules in the New Economy: Health and Safety Risks to Workers; RFA OH–01–006

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:

Name: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Extended Work Schedules in the New Economy: Health and Safety Risks to Workers; RFA OH–01–006.

Times and Dates: 8 a.m.–8:30 a.m., June 11, 2001. (Open) 8:30 a.m.–5 a.m., June 11, 2001. (Closed) 8:30 a.m.–5 a.m., June 12, 2001. (Closed)

Place: Sheraton Station Square, 7 Station Square Drive, Pittsburgh, PA 15219

Status: Portions of 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 Deputy Director for Program Management, CDC, pursuant to Public Law 92–463.

Matters to be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to Program Announcement: RFA OH–01–006.

For Further Information Contact: Pervis C. Major, Ph.D., Scientific Review Administrator, National Institute for Occupational Safety and Health, CDC, 1095 Willowdale Road, M/S B228, Morgantown, West Virginia 26505, telephone 304–285–5979.

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: May 21, 2001.

Carolyn J. Russell,

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

[FR Doc. 01–13239 Filed 5–24–01; 8:45 am]

BILLING CODE 4163–19–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee on Immunization Practices: 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: Advisory Committee on Immunization Practices (ACIP).

Times and Dates: 8:30 a.m.–6:45 p.m., June 20, 2001.

8 a.m.–5:15 p.m., June 21, 2001.

Place: Atlanta Marriott Century Center, 2000 Century

Boulevard, N.E., Atlanta, Georgia 30345–3377.

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

Purpose: The Committee is charged with advising the Director, CDC, on the appropriate uses of immunizing agents. In addition, under 42 U.S.C. 1396s, the Committee is mandated to establish and periodically review and, as appropriate, revise the list of vaccines for administration to vaccine-eligible children through the Vaccines for Children (VFC) program, along with schedules regarding the appropriate periodicity, dosage, and contraindications applicable to the vaccines.

Matters to be Discussed: The agenda will include a discussion on vaccine safety issues for yellow fever vaccine: Is a yellow fever vaccine booster needed every 10 years; adult high-risk hepatitis B immunization; current epidemiology of HBV infection in the US; status of immunization of high risk persons in STD clinics, prisons, and non-traditional settings; update on tetanus toxoid vaccine shortage; what should be CDC's role if the influenza vaccine supply remains unclear for the 2001–2002 season; summary of the live-attenuated influenza vaccine working group meeting; vaccine safety updates: the Brighton collaboration, Institute of Medicine Report on measles, mumps and rubella vaccine and autism; update on thimerosal; update from the National Center for Infectious Diseases; update from the National Immunization Program; update from the Food and Drug Administration; update from the National Institutes of Health; update from the Vaccine Injury Compensation Program; update from the National Vaccine Program; final decision on general recommendations for immunization; discontinuation of human rabies vaccine for intradermal pre-exposure use; update on current phase III HIV vaccine efficacy trials; update on risk of meningococcal disease among microbiology laboratory workers; use of economic evaluation for setting health policy; recommended childhood immunization schedule, 2002; and should there be an immunization schedule for adult immunization.

Agenda items are subject to change as priorities dictate.

For Further Information Contact: Gloria A. Kovach, Program Analyst, Epidemiology and Surveillance Division, National Immunization Program, CDC, 1600 Clifton Road, NE, m/s E61, Atlanta, Georgia 30333. Telephone 404/639–8096.

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: May 21, 2001.

Carolyn J. Russell,

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

[FR Doc. 01–13238 Filed 5–24–01; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of 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: Safety and Occupational Health Study Section (SOHSS), National Institute for Occupational Safety and Health (NIOSH).

Times and Dates: 8:30 a.m.–5:30 p.m., June 28, 2001; 8 a.m.–5 p.m., June 29, 2001.

Place: Embassy Suites, 1900 Diagonal Road, Alexandria, VA, 22314.

Status: Open 8:30 a.m.–9:30 a.m., June 28, 2001; Closed 9:30 a.m.–5:30 p.m., June 28, 2001; Closed 8 a.m.–5 p.m., June 29, 2001.

Purpose: The Safety and Occupational Health Study Section will review, discuss, and evaluate grant application(s) received in response to the Institute's standard grants review and funding cycles pertaining to research issues in occupational safety and health and allied areas.

It is the intent of NIOSH to support broad-based research endeavors in keeping with the Institute's program goals which will lead to improved understanding and appreciation for the magnitude of the aggregate health burden associated with occupational injuries and illnesses, as well as to support more focused research projects which will lead to improvements in the delivery of occupational safety and health services and the prevention of work-related injury and illness. It is anticipated that research funded will promote these program goals.

Matters to be Discussed: The meeting will convene in open session from 8:30–9:30 a.m. on June 28, 2001, to address matters related to the conduct of Study Section business. The remainder of the meeting will proceed in closed session. The purpose of the closed sessions is for the Safety and Occupational

Health Study Section to consider safety and occupational health related grant applications. These portions of 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 Associate Director for Management and Operations, CDC, pursuant to Pub. L. 92–463.

Agenda items are subject to change as priorities dictate.

For Further Information Contact: Charles N. Rafferty, Ph.D., NIOSH Scientific Review Administrator, Bethesda, Maryland. Telephone (301)435–3562, E-mail *raffertc@csr.nih.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: May 21, 2001.

Carolyn J. Russell,

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

[FR Doc. 01–13237 Filed 5–24–01; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Statewide Automated Child Welfare Information System (SACWIS) Assessment Review GuidE (SARGE).

OMB No. 0970–0159.

Description: HHS cannot fulfill its obligation to effectively serve the nation's Adoption and Foster Care populations, nor report meaningful and reliable information to Congress about the extent of problems facing these children or the effectiveness of assistance provided to this population, without access to timely and accurate information. Currently, SACWIS systems support State efforts to meet the following Federal reporting requirements: the Adoption and Foster Care Analysis and Reporting System (AFCARS) required by section 479(b)(2) of the Social Security Act; the National Child Abuse and Neglect Data System (NCANDS); Child Abuse Prevention and Treatment Act (CAPTA); and the new Chafee Independent Living Program.

Forty-eight States and the District of Columbia have developed or have committed to develop a SACWIS system with Federal financial participation. The purpose of these reviews is to ensure that all aspects of the project, as described in the approved Advance Planning Document, have been adequately completed, and conform to applicable regulations and policies.

To initiate a review, States will submit the completed SACWIS Assessment Review GuidE (SARGE) and other documentation at the point that they have completed system development and the system is operational statewide. The additional documents submitted as part of this process should all be readily available to the State as a result of good project management.

The information collected in the SACWIS Assessment Review Guide will allow State and Federal officials to determine if the State's SACWIS system meets the requirements for title IV–E Federal financial participation defined at 45 CFR 1355.50. Additionally, other States will be able to use the documentation provided as part of this review process in their own system development efforts.

Respondents: State Title IV–E Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Review	6	1	200	1200
Estimated Total Annual Burden Hours				1200

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: May 22, 2001.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 01–13257 Filed 5–24–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Oncologic Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Oncologic Drugs Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.