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.