Procurement and Grants Office, Program Announcement #99055, Centers for Disease Control and Prevention (CDC), 2920 Brandywine Road, Suite 3000, Atlanta, Georgia 30341, Telephone (770) 488–2724, Internet address: anf3@cdc.gov.

For program technical assistance, contact: Ted Jones, Program Manager, Office of Research Grants, National Center for Injury Prevention and Control, Centers for Disease Control and Prevention (CDC), 4770 Buford Highway, NE, Mailstop K–5, Atlanta, GA 30341–3724, Telephone (770) 488–4824, Internet address: tmj1@cdc.gov.

This and other CDC announcements are available through the CDC homepage on the Internet. The address for the CDC homepage is <http://www.cdc.gov>.

Interested applicants may receive a draft copy of the ''Policy for Solicitation and Selection of Injury Research Grant Proposals'' by calling 770/488–4265.

Dated: March 10, 1999.

John L. Williams,

Director, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 99–6311 Filed 3–15–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee on Childhood Lead Poisoning Prevention: Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the National Center for Environmental Health (NCEH) of the Centers for Disease Control and Prevention (CDC) announces the following committee meeting.

Name: Advisory Committee on Childhood Lead Poisoning Prevention.

Times and Dates: 8:30 a.m.–5 p.m., April 12, 1999; 8:30 a.m.–12 p.m., April 13, 1999.

Place: Atlanta Marriott Gwinnett Place, 1775 Pleasant Hill Hill Road, Duluth, Georgia 30136, telephone 770/925–2340.

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

Purpose: The Committee shall provide advice and guidance to the Secretary; the Assistant Secretary for Health; and the Director, CDC, regarding new scientific knowledge and technological developments and their practical implications for childhood lead poisoning prevention efforts. The Committee shall also review and report regularly on childhood lead poisoning prevention practices and recommend improvements in national childhood lead poisoning prevention efforts.

Matters to Be Discussed: Agenda items include: Childhood Lead Poisoning Prevention activities update, HCFA's Medicaid lead screening policy, Screening and Case Management Working Group updates, HUD lead program update, global dimensions of the lead problem, USAID update, and emerging issues involving foreign-born children.

Agenda items are subject to change as priorities dictate.

Opportunities will be provided during the meeting for oral comments. Depending on the time available and the number of requests, it may be necessary to limit the time of each presenter.

Contact Person for More Information: Barbara Nelson, Program Analyst, Lead Poisoning Prevention Branch, Division of Environmental Hazards and Health Effects, NCEH, CDC, 4770 Buford Highway, NE, M/S F–42, Atlanta, Georgia 30341–3724, telephone 770/488–7272, fax 770/488–7335.

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.

John C. Burckhardt,

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

[FR Doc. 99–6310 Filed 3–15–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Project

Title: Developmental Disabilities Council State Plan.

OMB No.: 0980–0162.

Description: Developmental Disabilities Councils (DD Councils) in each State are required under the Developmental Disabilities Assistance and Bill of Rights Act (42 U.S.C. 6000 et seq.) to develop plans on a triennial basis and to review those plans at least annually. Each Council develops its plan as a basis for promoting systems change and capacity building in service systems for persons with developmental disabilities in the State. The State plan must be made available for public comment in the State and must be approved by the Governor of the State. After that it is submitted to the Department of Health and Human Services, which will use the information to ensure compliance of the State with requirements in the Act. The information in the State plan is also used as one basis for providing technical assistance, such as during site visits. The burden statement of 130 hours per State for preparing this plan is annualized over the three-year period of the plan. This requirement was recently redesigned with significant involvement of State Developmental Disabilities Councils.

Respondents: State, Local or Tribal Government.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Three Year State Plan	55	1	130	7,150

Estimated Total Annual Burden Hours: 7,150.

In compliance with the requirements of Section 3506(c)(2)(A) 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: March 9, 1999.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 99–6291 Filed 3–15–99; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 93N–0445]

Agency Information Collection Activities; Announcement of OMB Approval; Financial Disclosure by Clinical Investigators

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Financial Disclosure by Clinical Investigators'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezzuto, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of December 31, 1998 (63 FR 72171), 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 0910–0396. The approval expires on March 31, 2002. A copy of the supporting statement for this information collection is available on the Internet at ''http://www.fda.gov/ohrms/dockets''.

Dated: March 9, 1999.

William K. Hubbard,

Acting Deputy Commissioner for Policy.

[FR Doc. 99–6339 Filed 3–15–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98D–0362]

Site Specific Stability Data for Drug and Biologic Applications; Public Meeting; Request for Comment

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comment.

SUMMARY: The Food and Drug Administration (FDA) is announcing a public meeting on site specific stability data for drug and biologic applications. The agency has scheduled the public meeting to discuss scientific issues related to a section of the draft guidance entitled ''Draft Guidance for Industry—Stability Testing of Drug Substances and Drug Products.'' Specifically, the agency will discuss the section of the draft guidance entitled ''Site-Specific Stability Data for Drug and Biologic Applications.'' The agency invites comments on issues related to the meeting.

DATES: The public meeting will be held on March 31, 1999, from 9 a.m. to 2 p.m. Submit written notices of participation by March 24, 1999. Submit written comments on the specific issues of the meeting by June 14, 1999.

ADDRESSES: The public meeting will be held at the Holiday Inn Bethesda, 8120 Wisconsin Ave., Bethesda, MD 20814. Submit written notices of participation to Kimberly L. Topper or Angie Whitacre (addresses below). Submit written comments on the draft guidance to the Dockets Management Branch (HFA–305), 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Kimberly L. Topper or Angie Whitacre, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7001, or e-mail topperk@cder.fda.gov.

Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by March 24, 1999. Oral presentations from the public will be scheduled. Time allotted for each presentation may be limited. Those persons desiring to make formal oral presentations should notify the contact person before March 24, 1999 (providing name, firm name, address, and telephone number), and submit a brief statement of the general nature of the evidence or arguments they wish to present, and an indication of the approximate time requested to make their presentation.

Dated: March 9, 1999.

William K. Hubbard,

Acting Deputy Commissioner for Policy.

[FR Doc. 99–6301 Filed 3–15–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer on (301) 443–7978.

Comments are invited on: (a) whether the proposed collections of information are 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) ways to enhance 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.