U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters To Be Discussed: The meeting will include the review, discussion, and evaluation of ''CDC Grants for Public Health Research Dissertation (Panel A–3), PAR07–231.''

Contact Person for More Information: Sheree Marshall Williams, Ph.D., M.Sc., Scientific Review Administrator, CDC, 1600 Clifton Road, NE., Mailstop D72, Atlanta, GA 30333, Telephone (404) 639–4896.

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: June 6, 2008.

Elaine L. Baker,

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

[FR Doc. E8–13185 Filed 6–11–08; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: State Plan for Foster Care, Independent Living Services and Adoption Assistance under Title IV–E of the Social Security Act.

OMB No.: 0980–0141.

Description: A State plan is required by sections 471 and 477(b)(2), part IV–E of the Social Security Act (the Act) for each public child welfare agency requesting Federal funding for foster care, independent living services and adoption assistance under the Act. The State plan is a comprehensive narrative description of the nature and scope of a State's programs and provides assurances the programs will be administered in conformity with the specific requirements stipulated in title IV–E. The plan must include all applicable State statutory, regulatory, or policy references and citation for each requirement as well as supporting documentation. A State may use the pre-print format prepared by the Children's Bureau of the Administration for Children and Families or a different format on the condition that the format used includes all of the title IV–E State plan requirements of the Act.

Respondents: State and Territorial Agencies (State Agencies) administering or supervising the administration of the title IV–E program.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Title IV–E State Plan	13	1	15	195

Estimated Total Annual Burden Hours: 195.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, 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. E-mail address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project.

Fax: 202–395–6974,

Attn: Desk Officer for the Administration for Children and Families.

Dated: June 5, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E8–13089 Filed 6–11–08; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2001–D–0067] (formerly Docket No. 2001D–0185)

Draft Guidance for Industry on Providing Regulatory Submissions in Electronic Format—Postmarketing Individual Case Safety Reports; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry entitled ''Providing Regulatory Submissions in Electronic Format—Postmarketing Individual Case Safety Reports.'' This draft guidance consolidates and revises information in two existing draft guidances pertaining to electronic submission of postmarketing individual case safety reports (ICSRs) and attachments to ICSRs (ICSR attachments). The submission of ICSRs and ICSR attachments in an electronic format significantly improves the agency's efficiency in processing, archiving, and reviewing the reports.

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the agency considers your comments on this draft guidance before it begins work on the final version of the guidance, submit written or electronic comments on the draft guidance, including comments regarding proposed collection of information, by August 11, 2008.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, rm. 2201, Silver Spring, MD 20993–0002, or to the Office of Communication, Training, and Manufacturers Assistance (HFM–40), Center for Biologics Evaluation and Research (CBER), Food and Drug Administration, 1401 Rockville Pike, suite 200N, Rockville, MD 20852–1448. Send one self-addressed adhesive label to assist that office in processing your