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: PA #01066

Contact Person for More Information: Marsha Jones, Health Scientist, Centers for Disease Control and Prevention, National Center for Infectious Diseases, 1600 Clifton Road, m/s C19, Atlanta, GA., 30333. Telephone (404)639–2603, email: *maj4@cdc.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: July 25, 2001.

Carolyn J. Russell,

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

[FR Doc. 01–18970 Filed 7–30–01; 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 Projects

Title: Project 1099.

OMB No.: 0970–0183.

Description: A voluntary program which provides States' Child Support Enforcement agencies, upon their request, access to the earned and unearned income information reported to IRS by employers and financial institutions. The IRS 1099 information is used to locate noncustodial parents and to verify income and employment.

Respondents: State IV–D programs.

TABLE OF BURDEN ESTIMATES FOR INFORMING PARENTS OF THEIR RIGHTS AND RESPONSIBILITIES AND FOR PROVIDING TRAINING

Reporting	Number of respondents	Number of responses per respondent per year	Average burden hours per response	Total burden hours
States	12	12	2	288

Estimated Total Annual Burden Hours: 288.

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: July 24, 2001.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 01–18950 Filed 7–30–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. 01D–0294 and 01D–0295]

Draft Guidances for Industry on Providing Regulatory Submissions to Office of Food Additive Safety in Electronic Format: General Considerations and for Food Additive and Color Additive Petitions; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of two draft guidances for industry entitled ''Providing Regulatory Submissions to Office of Food Additive Safety in Electronic Format—General Considerations'' and ''Providing Regulatory Submissions to Office of Food Additive Safety in Electronic Format for Food Additive and Color Additive Petitions.'' These documents are the first in a series of guidance documents intended to provide guidance for industry regarding the preparation of regulatory submissions in electronic format to the Office of Food Additive Safety (OFAS), Center for Food Safety and Applied Nutrition (CFSAN). OFAS is providing these draft guidances as part of its implementation of 21 CFR part 11 and the Food Additives Regulatory Management (FARM) Project.

DATES: Submit written or electronic comments concerning these draft guidances by October 1, 2001, to ensure adequate consideration in the preparation of revised guidances, if warranted. However, you may submit written or electronic comments at any time. Submit written comments concerning the collection of information by October 1, 2001.

ADDRESSES: Submit written comments concerning these draft guidances and the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to http://www.fda.gov/dockets/ecomments. All comments should be identified with the corresponding docket number found in brackets in the heading of this document. Submit written requests for single copies of the draft guidances for industry entitled ''Providing Regulatory Submissions to