Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: October 3, 2001.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 01–25419 Filed 10–9–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information collection Activity; Comment Request

Proposed Projects

Title: Application and Program Reporting Requirements for children's Justice Act Grants.

OMB No.: 0980–0196 Expired (5/31/01).

Description: The Program Instruction, prepared in response to the Children's Justice Act and authorized by Title I of the Child Abuse Prevention and Treatment Act (CAPTA) (as amended) and in the process of reauthorization, provides direction to States and Territories to accomplish the purposes of assisting States in developing, establishing, and operating programs designed to improve: (1) The handling of child abuse and neglect cases, particularly child sexual abuse and exploitation, in a manner which limits additional trauma to the child victim; (2) the handling of cases of suspected child abuse or neglect related fatalities; and (3) the investigation and prosecution of cases of child abuse and neglect, particularly child sexual abuse and exploitation.

This Program Instruction contains information collection requirements that are found in P.L. 104–235 at Sections 107(b), 107(d), and pursuant to receiving a grant award. The information being collected is required by statute to be submitted pursuant to receiving a grant award. The information submitted will be used by the agency to ensure compliance with the statute; to monitor, evaluate, and measure grantee achievements in addressing the investigation and prosecution of child abuse and neglect; and to report to Congress.

Respondents: State Governments.

Annual Burden Estimates:

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Application	52	1	40	2080
Annual Performance Report	52	1	20	1040
Estimated Total Annual Burden Hours				3,120

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: October 3, 2001.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 01–25420 Filed 10–9–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Community-Based Family Resource and Support Program.

OMB No.: 0970–0155 expired 05/31/2001.

Description: The Program Instruction, prepared in response to the enactment of the Community-Based Family Resource and Support Grants (CBFRS), as set forth in Title II of Pub. L. 104–235, Child Abuse Prevention and Treatment Act Amendments of 1996, and in the process of reauthorization, provides direction to the States and Territories to accomplish the purposes of (1) supporting State efforts to develop, operate, expand and enhance a network of community-based, prevention focused, family resource and support programs that coordinate resources among existing human service organizations within the State; and (2) fostering an understanding, appreciation, and knowledge of diverse populations in order to be effective in preventing and treating child abuse and neglect. This Program Instruction contains information collection requirements that are found in Pub. L. 104–235 at Sections 202(1)(A); 202(1)(B); 203(b)(1)(B); 205; 207; and pursuant to receiving a grant award. The information submitted will be used by the agency to ensure compliance with the statute, complete the calculation of the grant award entitlement, and provide training and technical assistance to the grantee.

Respondents: State Government.

Annual Burden Estimates:

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Application	52	1	40	2080
Annual Report	52	1	24	1248
Estimated Total Annual Burden Hours				3328

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, S.W., Washington, D.C. 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: October 2, 2001.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 01–25421 Filed 10–9–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Science Board to the Food and Drug Administration 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). The meeting will be open to the public.

Name of Committee: Science Board to the Food and Drug Administration.

General Function of the Committee: The board shall provide advice primarily to the agency's Senior Advisor for Science, and as needed, to the Commissioner of Food and Drugs and other appropriate officials on specific complex and technical issues as well as emerging issues within the scientific community in industry and academia. Additionally, the board will provide advice to the agency on keeping pace with technical and scientific evolutions in the fields of regulatory science, formulating an appropriate research agenda, and upgrading its scientific and research facilities to keep pace with these changes. It will also provide the means for critical review of agency sponsored intramural and extramural scientific research programs.

Date and Time: The meeting will be held on November 16, 2001, from 9 a.m. to 4:30 p.m.

Location: 5630 Fishers Lane, rm. 1066, Rockville, MD.

Contact: Susan Mackie Bond, Office of Science Coordination and Communication (HF–33), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–6687, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12603. Please call the Information Line for up-to-date information on this meeting.

Agenda: The board will hear and discuss external science review for FDA's Center for Devices and Radiological Health, emerging issues in FDA's oversight of clinical research, and emerging issues in pharmaceutical manufacturing.

Procedure: 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 November 1, 2001. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before November 1, 2001, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: October 1, 2001.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 01–25370 Filed 10–9–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Public Law 104–13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to OMB under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, call the HRSA Reports Clearance Officer on (301) 443–1129.

Comments are invited 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) 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