information collections must be mailed within 30 days of this notice directly to the OMB desk officer:

OMB Human Resources and Housing Branch, Attention: Brenda Aguilar, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: May 22, 2002.

John P. Burke, III,

Paperwork Reduction Act Team Leader, CMS Reports Clearance Officer, CMS Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards.

[FR Doc. 02–13866 Filed 6–3–02; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB review; comment request

Title: Child and Family Services Plan, Annual Progress and Services Report, and Budget Request

OMB No.: 0980–0047

Description: Under title IV–B, subparts 1 and 2, of the Social Security Act, States and Indian Tribes are to submit a five year Child and Family Services Plan, an Annual Progress and Services Report (APSR), and an annual budget request and estimated expenditure report (CFS–101). The plan is used by States and Indian Tribes to develop and implement services, and describe coordination efforts with other Federal, state and local programs. The APSR is used to provide updates and changes in the goals and services under the five year plan. The CFS–101 will be submitted annually with the APSR to apply for appropriated funds for the next fiscal year. The CFSP also includes the required State plans under Section 106 of the Child Abuse Prevention and Treatment Act and section 477 of title IV–E, the Chafee Foster Care Independence Program.

Respondents: 300

Annual Burden Estimates:

Instrument	Number of respondent	Number of responses per respondent	Average burden hours per response	Total burden hours
CFSP	300	1	500	150,000/5 = 30,000
APSR	300	1	270	81,000
CFS101	300	1	5	1,500
Estimated Total Annual Burden Hours:				112,500

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Information Services, 470 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer.

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, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF.

Dated: May 29, 2002.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 02–13880 Filed 6–3–02; 8:45 am]

BILLING CODE 4104–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02D–0124]

Draft Guidance for Industry: Notifying FDA of Fatalities Related to Blood Collection or Transfusion; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance document entitled ''Guidance for Industry: Notifying FDA of Fatalities Related to Blood Collection or Transfusion'' dated June 2002. The draft guidance document, when finalized, is intended to provide recommendations to blood collection or transfusion facilities on reporting fatalities related to blood or blood component collection or blood transfusion to FDA's Center for Biologics Evaluation and Research (CBER).

DATES: Submit written or electronic comments on the draft guidance to ensure their adequate consideration in preparation of the final document by September 3, 2002. General comments on agency guidance documents are welcome at any time.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Office of Communication, Training, and Manufacturers Assistance (HFM–40), Center for Biologics Evaluation and Research, Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448. Send one self-addressed adhesive label to assist the office in processing your requests. The document may also be obtained by mail by calling the CBER Voice Information System at 1–800–835–4709 or 301–827–1800, or by fax by calling the FAX Information System at 1–888–CBER–FAX or 301–827–3844. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

Submit written or electronic comments on the document 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.

FOR FURTHER INFORMATION CONTACT: Valerie A. Butler, Center for Biologics Evaluation and Research (HFM–17), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448, 301–827–6210.