Dated: July 16, 2008.

Brendan C. Kelly,

OPRE Reports Clearance Officer.

[FR Doc. E8–17352 Filed 7–29–08; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Proposed Projects

Title: Generic Clearance To Conduct Qualitative Data Collections.

OMB No.: New Collection.

Description: The Office of Planning, Research and Evaluation (OPRE), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), intends to request approval from the Office of Management and Budget (OMB) for a generic clearance that will allow OPRE to conduct a variety of qualitative data collections. Over the next three years, OPRE anticipates undertaking a variety of new research projects in the fields of cash welfare, employment and self-sufficiency, Head Start, child care, healthy marriage and responsible fatherhood, and child welfare. In order to inform the development of OPRE research, to maintain a research agenda that is rigorous and relevant, and to ensure that research products are as current as possible, OPRE will engage in a variety of qualitative data collections in concert with researchers and practitioners throughout the field. OPRE envisions using a variety of techniques including semi-structured discussions, focus groups, telephone interviews, and in person observations and site visits, in order to integrate the perspectives of program operators, policy officials and members of the research community.

Following standard Office of Management and Budget (OMB) requirements, OPRE will submit a change request to OMB individually for every group of data collection activities undertaken under this generic clearance. OPRE will provide OMB with a copy of the individual instruments or questionnaires (if one is used), as well as other materials describing the project.

Respondents: Administrators or staff of State and local agencies or programs in the relevant fields; academic researchers; and policymakers at various levels of government.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Semi-Structured Discussion and Information-Gathering Protocol	600	1	.5	300

Estimated Total Annual Burden Hours: 300.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: *OPREInfoCollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

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 ACF.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–D–0379]

Draft Guidance for Industry: Nucleic Acid Testing to Reduce the Possible Risk of Parvovirus B19 Transmission by Plasma-Derived Products; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft document entitled ''Guidance for Industry: Nucleic Acid Testing (NAT) to Reduce the Possible Risk of Parvovirus B19 Transmission by Plasma-Derived Products,'' dated July 2008. The draft guidance document provides to manufacturers of plasma-derived products recommendations for performing parvovirus B19 NAT as an in-process test for Source Plasma and recovered plasma to identify and help to prevent the use of plasma units containing high levels of parvovirus B19. The draft guidance also recommends how to report to the FDA implementation of parvovirus B19 NAT.

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 comment on this draft guidance before it begins work on the final version of the guidance, submit written or electronic comments on the draft guidance by October 28, 2008.

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 (CBER), Food and Drug Administration, 1401 Rockville Pike, suite 200N, Rockville, MD 20852–1448. Send one self-addressed adhesive label to assist the office in processing your requests. The draft guidance may also be obtained by mail by calling CBER at 1–800–835–4709 or 301–827–1800. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

Submit written comments on the draft guidance to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.regulations.gov*.

FOR FURTHER INFORMATION CONTACT: Paul E. Levine, Jr., Center for Biologics Evaluation and Research (HFM–17), Food and Drug Administration, 1401 Rockville Pike, suite 200N, Rockville, MD 20852–1448, 301–827–6210.

SUPPLEMENTARY INFORMATION: