Times and Dates: 10 a.m.–5 p.m., June 5, 2008. 8:30 a.m.–12:30 p.m., June 6, 2008.

Place: Auditorium A, Global Communications Center, Building 19, 1600 Clifton Road, NE., Atlanta, Georgia 30333.

Status: Open to the public, limited only by the space available.

Please Note: Due to current security measures, a valid government issued identification card with photo is required for admittance into the Roybal facility. Non-U.S. citizens wishing to attend should contact: Thomas G. Savel, M.D., telephone, (404) 498–2475. The deadline for notification of attendance is May 22, 2008.

Purpose: The board provides advice to the Secretary, HHS, and the Director, CDC, on strategies and goals for the programs and research within the national center; conducts peer-review of scientific programs; and monitors the overall strategic direction and focus of the national center. The board also performs second-level peer review of applications for grants-in-aid for research and research training activities, cooperative agreements, and research contract proposals relating to the broad areas within the national center.

Matters to be Discussed: The agenda will include an overview of the National Center for Public Health Informatics (NCPHI), including its mission, scope and goals. Participants will give detailed presentations of select intramural and extramural NCPHI activities. NCPHI intramural activity topics include biosurveillance, electronic laboratory reporting, and health information exchanges; extramural NCPHI activities include those involving its five Centers of Excellence in Public Health Informatics. Discussions focusing on future NCPHI program activities are also planned.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Thomas G. Savel, M.D., Designated Federal Official, National Center for Public Health Informatics, CDC, 1600 Clifton Road, NE., Mail Stop E–78, Atlanta, Georgia 30333; Telephone, (404) 498–2475.

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: May 9, 2008.

Elaine L. Baker,

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

[FR Doc. E8–11328 Filed 5–20–08; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB No.: 0970–0288]

Proposed Information Collection Activity; Comment Request

Proposed Projects: Title: Evaluation of the Improving Child Welfare Outcome through Systems of Care Grant Program.

Description: The 1994 Amendments to the Social Security Act (SSA) authorize the U.S. Department of Health and Human Services to review State child and family service programs to ensure conformance with the requirements in titles IV–B and IV–E of SSA. Under the Final Rule, which took effect March 25, 2000, States are assessed for substantial conformity with certain Federal requirements for child-welfare services. The Child and Family Service Reviews (CFSR), administered by the Children's Bureau, are designed to ensure conformity with Federal child-welfare requirements and, ultimately, to help States improve child-welfare services and outcomes, specifically safety, permanency and well-being outcomes for child-welfare-involved children and their families. States determined not to have achieved substantial conformity in any of the areas assessed are required to develop and implement Program Improvement Plans (PIP) addressing the areas of nonconformity.

The Systems of Care grant cluster, from which these data are proposed to be collected, is designed to encourage public child-welfare agencies to address the issues identified in their State's CFSR. The data collected from these demonstration sites will allow the Children's Bureau to test whether this approach can help States reach the goals stated in their PIP and explore how child welfare can benefit from being part of a system of care. Data will be collected via interviews, forms completed by project staff, surveys, focus groups and case-file reviews. Data also will be collected to determine the extent to which the Technical Assistance (TA) provided, brokered or contracted by the TA and Evaluation Center is meeting the needs of the grantees, and how.

Respondents: Systems of Care Project Directors (members of the Systems of Care collaborative may include representatives from mental health, juvenile justice, education, health, among others); child-welfare agency supervisors and caseworkers; partner agency caseworkers; and families who have been involved with the child-welfare system.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Stakeholder Survey	270	1	.5	135
Child-Welfare Agency Survey	600	1	1	600
Supervisor Interviews	90	1	1	90
Stakeholder Interviews	90	1	1	90
Project Director Interviews	23	1	1	23
Case Study Interviews	25	1	1.5	37.5
Focus Group with Family Members	102	1	1.5	153
Parent Partner Interviews	24	1	1	24
Child-Welfare Agency and Partner Agency Focus Groups	280	1	1.5	420
Community Description Form	9	1	2.5	22.5
Organizational Structure Form for Case Study Sites	3	1	2	6
Organizational Structure Form for Non-Case Study Sites	20	1	1	20
Collaborative Membership Form	23	1	1.5	34.5
Major Activities Form	23	1	1.5	34.5
Training and Technical Assistance Quality Assurance Assessment	23	1	1	23
Training and Technical Assistance Conference Call Feedback Forms	10	12	.25	30

Estimated Total Annual Burden Hours: 1,743.

In compliance with the requirements of Section 506(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 Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, *Attn:* ACF Reports Clearance Officer. E-mail address: *infocollection@acf.hhs.gov*. 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: May 14, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E8–11188 Filed 5–20–08; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB No.: 0970–0177]

Submission for OMB Review; Comment Request

Title: OCSE–157 Child Support Enforcement Program Annual Data Report.

Description: The information obtained from this form will be used to: (1) Report Child Support Enforcement activities to the Congress as required by law; (2) calculate incentive measures performance and performance indicators utilized in the program; and (3) assist the Office of Child Support Enforcement in monitoring and evaluating State Child Support programs.

Respondents: State, Local or Tribal Government.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
OCSE–157	54	1	7	378.0

Estimated Total Annual Burden Hours: 378.0

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, 378.0 370 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 following: Office of Management and Budget, Paperwork Reduction Project, Fax: 202–395–6974, Attn: Desk Officer for the Administration for Children and Families.

Dated: May 14, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E8–11190 Filed 5–20–08; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–D–0263]

Draft Guidance for Industry: Requalification Method for Reentry of Blood Donors Deferred Because of Reactive Test Results for Antibody to Hepatitis B Core Antigen (Anti-HBc); 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: Requalification Method for Reentry of Blood Donors Deferred Because of Reactive Test Results for Antibody to Hepatitis B Core Antigen (Anti-HBc)'' dated May 2008. The draft guidance document provides recommendations to establishments that collect human blood or blood components for a requalification method or process to reenter deferred donors into a donor pool based on a determination that the previous tests that were repeatedly reactive for anti-HBc were falsely positive and that there is no evidence of infection with Hepatitis B virus (HBV).

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 August 19, 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