Compliance With Rules and Contacting Contest Winners

Finalists and Champions must comply with all terms and conditions of these official rules, and winning is contingent upon fulfilling all requirements herein. The initial finalists will be notified by email, telephone, or mail after the date of the judging.

Privacy

Personal information provided by entrants on the nomination form through the challenge Web site will be used to contact selected finalists. Information is not collected for commercial marketing. Winners are permitted to cite that they won this challenge.

The names, cities, and states of selected Champions will be made available in promotional materials and at recognition events.

General Conditions

The HHS/CDC reserves the right to cancel, suspend, and/or modify the challenge, or any part of it, for any reason, at HHS/CDC's sole discretion.

Authority: 15 U.S.C. 3719

Dated: August 3, 2015.

Pamela J. Cox,

Director, Division of the Executive Secretariat, Office of the Chief of Staff, Centers for Disease Control and Prevention.

[FR Doc. 2015–20076 Filed 8–13–15; 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: State Court Improvement Program.

OMB No.: 0970–0307.

Description: The Court Improvement Program (CIP) is a mandatory formula grant funded under section 438 of the Social Security Act, and most recently reauthorized under the Child and Family Services Improvement and Innovation Act of 2012 (Pub. L. 112–34). The purpose of the CIP is to facilitate the handling of child welfare cases in the courts. All 50 states, Puerto Rico, and the District of Columbia receive grants under the program. The program requires two submissions annually from grantees that constitute information collections under the Paperwork Reduction Act.

The purpose of this notice is to request an extension of the Office of Management and Budget Control Number 0907–0307 permitting continued use of the information collections requires by ACF–CB–PI–12–02. The burden estimates are provided below. The Administration on Children, Youth, and Families anticipates issuing a new Program Instruction following reauthorization of the program in federal fiscal year 2017.

Respondents: State Courts.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Application	52	1	92	4,784
Annual Reports	52	1	86	4,472

Estimated Total Annual Burden Hours: 9,256.

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 Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email 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.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2015–20073 Filed 8–13–15; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Proposed Projects

Title: Office of Child Support Enforcement Child Support Portal Registration.

OMB No.: 0970–0370.

Description: The federal Office of Child Support Enforcement (OCSE), Division of Federal Systems maintains the Child Support Portal, which contains a variety of child support applications to help enforce state child support cases. To securely access child support applications, authorized users must register to use the Child Support Portal. Information collected from the registration form is used to authenticate and authorize the users.

The OCSE Child Support Portal Registration information collection activities are authorized by 42 U.S.C. 653(m)(2), which requires the Secretary to establish and implement safeguards to restrict access to confidential information in the Federal Parent Locator Service to authorized persons and to restrict use of such information to authorized purposes.

Respondents: Employers, Financial Institutions, Insurers, Tribal, and State Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Registration Screens	299	1	0.15	44.85

Estimated Total Annual Burden Hours: 45.

Additional Information

Copies of the proposed collection may be obtained 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.

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, Email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2015–20121 Filed 8–13–15; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–N–0001]

National Mammography Quality Assurance Advisory Committee, Renewal

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; renewal of advisory committee.

SUMMARY: The Food and Drug Administration (FDA) is announcing the renewal of the National Mammography Quality Assurance Advisory Committee by the Commissioner of Food and Drugs (the Commissioner). The Commissioner has determined that it is in the public interest to renew the National Mammography Quality Assurance Advisory Committee for an additional 2 years beyond the charter expiration date. The new charter will be in effect until July 6, 2017.

DATES: Authority for the National Mammography Quality Assurance Advisory Committee will expire on July 6, 2017 unless the Commissioner formally determines that renewal is in the public interest.

FOR FURTHER INFORMATION CONTACT: Sara J. Anderson, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg 66, Rm. 1643, Silver Spring, MD, 20993, *Sara.Anderson@fda.hhs.gov,* 301 796–7047.

SUPPLEMENTARY INFORMATION: Under 41 CFR 102–3.65 and approval by the Department of Health and Human Services under 45 CFR part 11 and by the General Services Administration, FDA is announcing the renewal of the National Mammography Quality Assurance Advisory Committee. The committee is a statutory Federal advisory committee established to provide advice to the Commissioner.

The Secretary and, by delegation, the Assistant Secretary for the Office of Public Health and Science, and the Commissioner of Food and Drugs are charged with the administration of the Federal Food, Drug and Cosmetic Act and various provisions of the Public Health Service Act. The Mammography Quality Standards Act of 1992 amends the Public Health Service Act to establish national uniform quality and safety standards for mammography facilities. The National Mammography Quality Assurance Advisory Committee advises the Secretary and, by delegation, the Commissioner of Food and Drugs in discharging their responsibilities with respect to establishing a mammography facilities certification program.

The Committee shall advise the Food and Drug Administration on:

A. Developing appropriate quality standards and regulations for mammography facilities;

B. Developing appropriate standards and regulations for bodies accrediting mammography facilities under this program;

C. Developing regulations with respect to sanctions;

D. Developing procedures for monitoring compliance with standards;

E. Establishing a mechanism to investigate consumer complaints;

F. Reporting new developments concerning breast imaging which should be considered in the oversight of mammography facilities;

G. Determining whether there exists a shortage of mammography facilities in rural and health professional shortage areas and determining the effects of personnel on access to the services of such facilities in such areas;

H. Determining whether there will exist a sufficient number of medical physicists after October 1, 1999; and

I. Determining the costs and benefits of compliance with these requirements.

The Committee shall consist of a core of 15 members, including the Chair. Members and the Chair are selected by the Commissioner or designee from among physicians, practitioners, and other health professionals, whose clinical practice, research specialization, or professional expertise includes a significant focus on mammography. Members will be invited to serve for overlapping terms of up to four years. Almost all non-Federal members of this committee serve as Special Government Employees. The core of voting members shall include at least 4 individuals from among national breast cancer or consumer health organizations with expertise in mammography, and at least 2 practicing physicians who provide mammography services. In addition to the voting members, the Committee shall include 2 nonvoting industry representatives who have expertise in mammography equipment. The Committee may include one technically qualified member, selected by the Commissioner or designee, who is identified with consumer interests.

Further information regarding the most recent charter and other information can be found at *http://www.fda.gov/AdvisoryCommittees/CommitteesMeetingMaterials/Radiation-EmittingProducts/NationalMammographyQualityAssuranceAdvisoryCommittee/ucm124611.htm* or by contacting the Designated Federal Officer (see **FOR FURTHER INFORMATION CONTACT**). In light of the fact that no change has been made to the committee name or description of duties, no