

support them in the requirements development phase of the acquisition lifecycle.

- The FFRDC must function so effectively as to act as an agent for the sponsor in the design and pursuit of mission goals.
- The FFRDC must provide rapid responsiveness to changing requirements for personnel in all aspects of strategic, technical and program management.
- The FFRDC must recognize government objectives as its own objectives, partnering with the sponsor in pursuit of excellence in public service.
- The FFRDC must allow for non-sponsor, other than CMS, work for operating Divisions within DHHS.

We are publishing this notice in accordance with 48 CFR 5.205(b) of the Federal Acquisition Regulations (FAR), to enable interested members of the public to provide comments on this proposed action. We note that this is the

second of three notices issued under the FAR.

The Request for Proposal will be posted on FedBizOpps in the Summer of 2011. Alternatively, a copy can be received by contacting the person listed in the **FOR FURTHER INFORMATION CONTACT** section above.

Dated: May 4, 2011.

Donald M. Berwick,
Administrator, Centers for Medicare & Medicaid Services.

[FR Doc. 2011-11708 Filed 5-12-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

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

OMB No.: 0970-0177.

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 (OCSE) in monitoring and evaluating State Child Support programs. OCSE is proposing minor changes to the OCSE-157 report instructions for medical support line items that will provide states with the option to define medical support to include private health insurance as well as other health care coverage such as Medicaid, Children's Health Insurance Program (CHIP) and other state coverage plans, and cash medical support. Further legislative or regulatory changes may be necessary to update medical child support policy.

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
Estimated Total Annual Burden Hours:	378

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 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.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2011-11796 Filed 5-12-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2011-N-0015]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Orphan Drugs; Common European Medicines Agency/ Food and Drug Administration Application Form for Orphan Medicinal Product Designation (Form FDA 3671)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by June 13, 2011.