

Hours per Recordkeeper: 100.

Total Annual Recordkeeping hours: 508,000.

C. Public Comment

Public comments are particularly invited on: Whether this collection of information is necessary; whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies of Proposals:

Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202-501-4755. Please cite OMB Control No. 9000-0094, Debarment and Suspension and Other Responsibility Matters, in all correspondence.

Dated: February 17, 2016.

Lorin S. Curit,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2016-03579 Filed 2-19-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Disaster Information Collection Form.

OMB No.: 0970-NEW.

Description: This is a request by the Administration for Children and Families (ACF) for a generic clearance for the Disaster Information Collection Form. An approval for a generic clearance is being requested because each of the thirteen program offices within ACF has a slightly different need for information about program impact information collection during a disaster.

ACF oversees more than 60 programs that affect the normal day to day operations of families, children, individuals and communities in the United States. Many of these programs

encourage grantees or state administrators to develop emergency preparedness plans, but do not have statutory authority to require these plans be in place. ACF facilitates the inclusion of emergency preparedness planning and training efforts for ACF programs.

Presidential Policy Directive-8 (PPD-8) provides federal guidance and planning procedures under established phases—protection, preparedness, response, recovery, and mitigation. The Disaster Information Collection Forms addressed in this clearance process provide assessment of ACF programs in disaster response, and recovery.

ACF/Office of Human Services Emergency Preparedness and Response (OHSEPR) has a requirement under PPD-8, the National Response Framework, and the National Disaster Recovery Framework to report disaster impacts to ACF-supported human services programs to the HHS Secretary's Operation Center (SOC) and interagency partners. ACF/OHSEPR works in partnership with the Assistant Secretary for Preparedness and Response (ASPR), and the Federal Emergency Management Agency (FEMA) to report assessments of disaster impacted ACF programs and the status of continuity of services and recovery.

Respondents: State administrators, and/or ACF grantees.

ANNUAL BURDEN ESTIMATES

[The burden cap for the Disaster Information Collection Form is estimated based on a single disaster per year. The estimate is for approximately 10 state administrators, or grantees to go through all of the applicable questions with the Regional and Central Office staff. Some ACF programs have more questions and may have more respondents.]

Table with 5 columns: Instrument, Number of respondents, Number of responses per respondent, Burden hours per response, Total burden hours. Row 1: Disaster Information Collection Form, 10, 15, 0.08 hours (5 minutes), 1.25 hours (75 minutes).

An estimate of the number of disasters that would warrant data collection is difficult to calculate due to the unpredictable nature of disasters. For example, in 2012, there were 95 disasters nationwide but OHSEPR did not collect data on all of them because they had minimal effects on ACF programs.

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. All requests should be identified by the title of the information collection. Email 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 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. 2016-03455 Filed 2-19-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Guidance for Tribal TANF.

OMB No.: 0970-0157.

Description: 42 U.S.C. 612 (Section 412 of the Social Security Act) requires each Indian Tribe that elects to administer and operate a TANF program to submit a TANF Tribal Plan. The TANF Tribal Plan is a mandatory

statement submitted to the Secretary by the Indian Tribe, which consists of an outline of how the Indian Tribes TANF program will be administered and operated. It is used by the Secretary to determine whether the plan is

approvable and to determine that the Indian Tribe is eligible to receive a TANF assistance grant. It is also made available to the public.

Respondents: Indian Tribes applying to operate a TANF program.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Request for State Data Needed to Determine the Amount of a Tribal Family Assistance Grant	24	1	68	1632

Estimated Total Annual Burden Hours: 1632.

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, 330 C Street SW., Washington DC 20201. 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. 2016-03453 Filed 2-19-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2014-E-0934]

Determination of Regulatory Review Period for Purposes of Patent Extension; SUPERA PERIPHERAL STENT SYSTEM

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined the regulatory review period for SUPERA PERIPHERAL STENT SYSTEM and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of an application to the Director of the U.S. Patent and Trademark Office (USPTO), Department of Commerce, for the extension of a patent which claims that medical device.

DATES: Anyone with knowledge that any of the dates as published (see the **SUPPLEMENTARY INFORMATION** section) are incorrect may submit either electronic or written comments and ask for a redetermination by April 22, 2016. Furthermore, any interested person may petition FDA for a determination regarding whether the applicant for extension acted with due diligence during the regulatory review period by August 22, 2016. See "Petitions" in the **SUPPLEMENTARY INFORMATION** section for more information.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments.

Comments submitted electronically, including attachments, to <http://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <http://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA-2014-E-0934 for "Determination of Regulatory Review Period for Purposes of Patent Extension; SUPERA PERIPHERAL STENT SYSTEM". Received comments will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at