

By direction of the Commission.

Donald S. Clark,
Secretary.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.598]

Announcement of the Award of a Single-Source Program Expansion Supplement Grant to Heartland Human Care Services in Chicago, IL

AGENCY: Office of Refugee Resettlement, ACF, HHS.

ACTION: Announcement of the award of a single-source program expansion supplement to Heartland Human Care Services (HHCS) to support expanded services to foreign trafficking victims, potential trafficking victims, and certain family members.

SUMMARY: The Administration for Children and Families (ACF), Office of Refugee Resettlement (ORR) announces the award of a single-source program expansion supplement grant to Heartland Human Care Services in Chicago, Illinois, for a total of \$144,822.

The supplemental funding will ensure that clients' essential needs, such as housing, transportation, communication, food, and medical care, will be met.

DATES: The period of support under these supplements is September 30, 2014 through September 29, 2015.

FOR FURTHER INFORMATION CONTACT: Maggie Wynne, Director, Division of Anti-Trafficking in Persons, Office of Refugee Resettlement, 901 D Street SW., Washington, DC 20024, Telephone (202) 401-4664. Email: maggie.wynne@acf.hhs.gov.

SUPPLEMENTARY INFORMATION: The National Human Trafficking Victim Assistance Program (NHTVAP) provides funding for comprehensive case management services to victims of trafficking and certain family members on a per capita basis. The NHTVAP

grantees help clients gain access to housing, employability services, mental health screening and therapy, medical care, and some legal services. During FY 2015, a grantee, Heartland Human Care Services (HHCS), served more clients than it had planned for in its budget for the year. Without the additional funding, HHCS would have to make significant cuts in services to current clients and limit the enrollment of new clients. With the supplemental funding, HHCS will be able to ensure that all of the clients' essential needs will be met.

Statutory Authority: Trafficking Victims Protection Act of 2000 (TVPA), as amended, Section 107(b)(1)(B), 22 U.S.C. 7105(b)(1)(B), authorizes funding for benefits and services to foreign victims of severe forms of trafficking in persons in the United States, potential victims of trafficking seeking HHS Certification, and certain family members.

Christopher Beach,
Senior Grants Policy Specialist, Office of Administration.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects: Application for Grants to States.

Title: State Access and Visitation Grant Application.

OMB No.: 0970-NEW.

Description: The Personal Responsibility and Work Opportunity Reconciliation Act of 1996 (PRWORA) created the "Grants to States for Access and Visitation" program (AV grant program). Funding for the program began in FY 1997 with a capped, annual entitlement of \$10 million. The statutory goal of the program is to provide funds to states that will enable them to provide services for the purpose of increasing noncustodial parent (NCP) access to and visitation with their children. State governors decide which

state entity will be responsible for implementing the AV grant program in addition to determining who will be served, what services will be provided, and whether the services will be statewide or in local jurisdictions. The statute specifies certain activities which may be funded including: Voluntary and mandatory mediation, counseling, education, the development of parenting plans, supervised visitation, and the development of guidelines for visitation and alternative custody arrangements. Even though OCSE manages this program, the funding for the AV grant is separate from funding for federal and state administration of the Child Support program.

Section 469B(e)(3) of the Social Security Act (Pub. L. 104-193) requires that each state receiving an AV grant award shall monitor, evaluate and report on such programs in accordance with regulations. Additionally, the Catalog of Federal Domestic Assistance, states that there is an application requirement for Grants to States for Access and Visitation Programs (93.597). The application process will assist OCSE in complying with this requirement and will reflect a greater emphasis on program efficiency, coordination of services, and increased attention to family safety.

The application will require states to submit a program plan, indicating how they anticipate spending their funds within the program statute and regulations. The applications will cover three fiscal years and any changes made to the plan during the three year period will require a notification of change to OCSE.

OCSE will review the applications to ensure that planned services meet the requirements laid out in Section 469B(e)(3) of the Social Security Act (Pub. L. 104-193). This review will include monitoring of program compliance and the safe delivery of services. In addition to monitoring, the report will also assist in OCSE's ability to provide technical assistance to states that would like assistance.

Respondents: Recipients of the Access & Visitation Grant (54 states and territories).

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Fillable word document	54	1	10	540

Estimated Total Annual Burden Hours: 540.

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.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2015-N-4602]

Streamlining Regulations for Good Manufacturing Practices for Hearing Aids; Public Workshop; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop; request for comments.

SUMMARY: The Food and Drug Administration (FDA) is announcing the following public workshop entitled "Streamlining Regulations for Good Manufacturing Practices (GMPs) for Hearing Aids." The topic to be

discussed is the appropriate level of good manufacturing practices (GMPs) regulation to ensure the safety and effectiveness of air-conduction hearing aid devices.

DATES: The public workshop will be held on April 21, 2016, from 8:30 a.m. to 5 p.m. Submit either electronic or written comments on the public workshop by May 19, 2016.

ADDRESSES: The public workshop will be held at FDA's White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993-0002. Entrance for the public meeting participants (non-FDA employees) is through Building 1 where routine security check procedures will be performed. For parking and security information, please refer to <http://www.fda.gov/AboutFDA/WorkingatFDA/BuildingsandFacilities/WhiteOakCampusInformation/ucm241740.htm>.

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-2015-N-4602 for "Streamlining Regulations for Good Manufacturing Practices (GMPs) for Hearing Aids." Received comments will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <http://www.regulations.gov> or at the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

- *Confidential Submissions*—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <http://www.regulations.gov>. Submit both copies to the Division of Dockets Management. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as "confidential." Any information marked as "confidential" will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <http://www.fda.gov/regulatoryinformation/dockets/default.htm>.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to <http://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the "Search" box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.