DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: ANA Project Impact Assessment Survey.

OMB No.: 0970–0379.

Description: The information collected by the Project Impact Assessment Survey is needed for two main reasons: (1) To collect crucial information required to report on the Administration for Native Americans' (ANA) established Government Performance and Results Act (GPRA) measures, and (2) to properly abide by ANA's congressionally-mandated statute (42 United States Code 2991 et seq.) found within the Native American Programs Act of 1974, as amended, which states that ANA will evaluate projects assisted through ANA grant dollars ''including evaluations that describe and measure the impact of such projects, their effectiveness in achieving stated goals, their impact on related programs, and their structure and mechanisms for delivery of services.'' The information collected with this survey will fulfill ANA's statutory requirement and will also serve as an important planning and performance tool for ANA.

The ANA Project Impact Assessment Survey, previously approved under information collection number OMB CN: 0970–0379, expires on 7/31/2013. This notice is issued to support ANA's continued use of the survey, with minor changes to eliminate duplication of reporting and improve the clarity and content of questions. These minor changes will not result in additional reporting burden on ANA grantees.

Respondents: Tribal Governments, Native American nonprofit organizations, and Tribal Colleges and Universities.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ANA Project Impact Assessment Survey	85	1	6	510

Estimated Total Annual Burden Hours: 510.

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, Fax: 202–395–7285, Email: *OIRA_SUBMISSION@OMB.EOP.GOV.* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2013–10252 Filed 4–30–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.676]

Announcement of the Award of Three Urgent Single-Source Grants To Support Shelter Care for Unaccompanied Alien Children

AGENCY: Office of Refugee Resettlement, Administration for Children and Families, Department of Health and Human Services.

ACTION: The Office of Refugee Resettlement (ORR) announces the award of three urgent single-source grants from the Unaccompanied Alien Children's Program to KidsPeace in Bethlehem, PA, St. Peter and Joseph Children's Home in San Antonio, TX, and Seton Home in San Antonio, TX.

SUMMARY: The Administration for Children and Families (ACF), Office of Refugee Resettlement (ORR) announces the award of three urgent single awards to the following organizations.

Organization	Location	Amount
KidsPeace	Bethlehem, PA	$3,105,096
St. Peter and Joseph Children's Home	San Antonio, TX	1,630,000
Seton Home	San Antonio, TX	650,000

These urgent awards will support the expansion of bed capacity and supportive services to meet the number of unaccompanied alien children referrals from the Department of Homeland Security (DHS). The funding program is mandated by section 462 of the Homeland Security Act to ensure appropriate placement of all referrals from the DHS. The program is tied to DHS apprehension strategies and to the sporadic numbers of border crossers.

DATES: The period of support provided by these awards is April 15, 2013 through September 30, 2013.

FOR FURTHER INFORMATION CONTACT: Jallyn Sualog, Acting Director, Division of Children's Services, Office of Refugee

Resettlement, 901 D Street SW., Washington, DC 20447, Telephone (202) 401–4997. Email: *jallyn.sualog@acf.hhs.gov*.

SUPPLEMENTARY INFORMATION: Since the beginning of FY 2013, the Unaccompanied Alien Children (UAC) program has seen a dramatic increase in the number of DHS referrals. The influx of border crossers referred by DHS has grown beyond anticipated rates resulting in the need for a significant increase in the number of shelter beds and supportive services for the children.

The UAC program has specific requirements for the provision of services to unaccompanied alien children. The named organizations were chosen for the noncompetitive awards because they already have the infrastructure, licensing, and appropriate levels of trained staff to meet service requirements and the urgent need for expanded services in order to respond to the increased numbers of unaccompanied children. The immediate provision of services will alleviate the buildup of children held in border patrol stations while awaiting placement in shelter care.

Statutory Authority: Section 462 of the Homeland Security Act, (6 U.S.C. 279) and sections 235(c) and 235(d) of the William Wilberforce Trafficking Victims Protection Reauthorization Act of 2008, (8 U.S.C. 1232(c) and 1232(d)).

Eskinder Negash,

Director, Office of Refugee Resettlement.

[FR Doc. 2013–10311 Filed 4–30–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0427]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Medical Devices; Inspection by Accredited Persons Program

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Medical Devices; Inspection by Accredited Persons Program'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Daniel Gittleson, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–5156, *daniel.gittleson@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: On January 31, 2013, the Agency submitted a proposed collection of information entitled ''Medical Devices; Inspection by Accredited Persons Program'' to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0510. The approval expires on April 30, 2016. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: April 26, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–10248 Filed 4–30–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0976]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance: Emergency Use Authorization of Medical Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Emergency Use Authorization of Medical Products'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–7726, *ila.mizrachi@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: On January 8, 2013, the Agency submitted a proposed collection of information entitled ''Emergency Use Authorization of Medical Products'' to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0595. The approval expires on March 31, 2016. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: April 26, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–10247 Filed 4–30–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0867]

Ashley Brandon Foyle: Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is issuing an order under the Federal Food, Drug, and Cosmetic Act (the FD&C Act) debarring Ashley Brandon Foyle for 5 years from providing services in any capacity to a person that has an approved or pending drug product application. FDA bases this order on a finding that Mr. Foyle was convicted of introducing and delivering for introduction into interstate commerce a misbranded drug, which relates to the development or approval, including the process for development or approval, of drug products and to the regulation of drug products under the FD&C Act. In addition, FDA determined that the type of conduct that served as the basis for Mr. Foyle's conviction undermines the process for the regulation of drugs. Mr. Foyle was given notice of the proposed debarment and an opportunity to request a hearing within the prescribed timeframe by regulation but failed to respond. Mr. Foyle's failure to respond constitutes a waiver of his right to a hearing concerning this action.

DATES: This order is effective May 1, 2013.

ADDRESSES: Submit applications for termination of debarment to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Kenny Shade, Office of Regulatory