the HHS Executive Development Program, including developmental activities, rotational assignments, and the Candidate Development Program; and (13) advises on development of executive succession planning activities.

Title 42 and Immigration Activity (CAJQG3). (1) Provides leadership, technical assistance, guidance, and consultation in the administration of policies and procedures for appointment of individuals through the distinguished consultants, experts, consultants, and fellows under Title 42 appointment authorities; (2) provides technical guidance and visa-assistance for employment based, CDC-sponsored visas; (3) administers and manages the Exchange Visitor Program; (4) works closely with the US Office of Exchange and Cultural Affairs, US Citizenship and Immigration Services, US Department of Homeland Security, US Department of State, Office of the Secretary/DHHS, and US Department of Labor) to facilitate immigration procedures; (5) reviews, processes and files H–1B, 0–1, and Green Card (I–140) Petitions with the U.S. Citizenship and Immigration Services; (6) provides advisory services and guidance on employment based green card petitions in the Alien of Extraordinary Ability category; (7) issues DS–2019s (Certificate of Eligibility for J–1 Exchange Visitor Status) through the Student and Exchange Visitor Information System to non US citizens seeking CDC J–1 visa sponsorship; (8) coordinates and provides consultations and guidance on Interested Government Agency Waivers; (9) provides Immigration Training Workshops to CDC/ATSDR Administrative Staff; (10) determines the appointment mechanism, legal status, and work authorizations for 5,000+ non US citizens through the Visitors and Management System; and (11) administers and manages the Guest Researcher and Oak Ridge Institute for Science and Education Program.

Dated: April 22, 2014.

Sherri A. Berger,

MSPH, Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2014–10178 Filed 5–5–14; 8:45 am]

BILLING CODE 4163–18–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Request for Assistance for Child Victims of Human Trafficking.

OMB No.: 0970–0362.

Description: The William Wilberforce Trafficking Victims Protection Reauthorization Act (TVPRA) of 2008, Public Law 110–457, directs the U.S. Secretary of Health and Human Services (HHS), upon receipt of credible information that an alien child may have been subjected to a severe form of trafficking in persons and is seeking Federal assistance available to victims of trafficking, to promptly determine if the child is eligible for interim assistance. The law further directs the Secretary of HHS to determine if a child receiving interim assistance is eligible for assistance as a victim of a severe form of trafficking in persons after consultation with the Attorney General, the Secretary of Homeland Security, and nongovernmental organizations with expertise on victims of severe form of trafficking.

In developing procedures for collecting the necessary information from potential child victims of trafficking, their case managers, attorneys, or other representatives to allow HHS to grant interim eligibility, HHS devised a form. HHS has determined that the use of a standard form to collect information is the best way to ensure requestors are notified of their option to request assistance for child victims of trafficking and to make prompt and consistent determinations about the child's eligibility for assistance.

Specifically, the form asks the requestor for his/her identifying information, for information on the child, information describing the type of trafficking and circumstances surrounding the situation, and the strengths and needs of the child. The form also asks the requestor to verify the information contained in the form because the information could be the basis for a determination of an alien child's eligibility for federally funded benefits. Finally, the form takes into consideration the need to compile information regarding a child's circumstances and experiences in a non-directive, child-friendly way, and assists the requestor in assessing whether the child may have been subjected to trafficking in persons.

The information provided through the completion of a Request for Assistance for Child Victims of Human Trafficking form will enable HHS to make prompt determinations regarding the eligibility of an alien child for interim assistance, inform HHS' determination regarding the child's eligibility for assistance as a victim of a severe form of trafficking in persons, facilitate the required consultation process, and enable HHS to assess potential child protection issues. HHS proposes to make several small, technical changes to the form, including the elimination of an unnecessary paragraph and updated references to the Trafficking Victims Protection Act of 2000, as amended, to reflect changes to that law.

Respondents: Representatives of governmental and nongovernmental entities providing social, legal, or protective services to alien persons under the age of 18 (children) in the United States who are neither U.S. citizens nor Lawful Permanent Residents and who may have been subjected to severe forms of trafficking in persons.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Request for Assistance for Child Victims of Human Trafficking	40	1	1	40

Estimated Total Annual Burden Hours: 40.

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. 2014–10307 Filed 5–5–14; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0530]

Center for Devices and Radiological Health Guidance Development and Prioritization; Public Workshop; Requests for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop; request for comments.

The Food and Drug Administration (FDA) is announcing the following public workshop entitled ''Center for Devices and Radiological Health Guidance Development and Prioritization Public Workshop.'' The topics to be discussed include the FDA's Center for Devices and Radiological Health's (CDRH) guidance development process, guidance development best practices for FDA, CDRH, and CDRH stakeholders, and CDRH guidance priorities and priority development.

Date and Time: The public workshop will be held on June 5, 2014, from 9 a.m. to 3 p.m.

Location: The public workshop will be held at the FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (rm. 1503A), Silver Spring, MD 20993–0002. Entrance for public workshop 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.*

Contact Person: Cathy Norcio, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, rm. 5448, Silver Spring, MD 20993–0002, 301–796–5446, email: *Catherine.norcio@fda.hhs.gov.*

Registration: Registration is free and available on a first-come, first-served basis. Persons interested in attending this public workshop must register online by 5 p.m., EDT, May 29, 2014. Early registration is recommended because facilities are limited and, therefore, FDA may limit the number of participants from each organization. If time and space permits, onsite registration on the day of the workshop will be provided beginning at 8 a.m.

If you need special accommodations due to a disability, please contact Susan Monahan (301–796–5661 or email: *susan.monahan@fda.hhs.gov*) no later than May 22, 2014.

To register for the public workshop, please visit FDA's Medical Devices News & Events—Workshops & Conferences calendar at *http://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/default.htm.* (Select this public workshop from the posted events list.) Please provide complete contact information for each attendee, including name, title, affiliation, email, and telephone number. Those without Internet access should contact Susan Monahan to register (see *Registration* contact for special accommodations). Registrants will receive confirmation after they have been accepted. You will be notified if you are on a waiting list.

Streaming Webcast of the Public Workshop: This public workshop will also be Webcast. Persons interested in viewing the Webcast must register online by 5 p.m., EDT, May 29, 2014. Early registration is recommended because Webcast connections are limited. Organizations are requested to register all participants, but to view using one connection per location. Webcast participants will be sent technical system requirements after registration and will be sent connection access information no later than June 2, 2014. If you have never attended a Connect Pro event before, test your connection at *https://collaboration.fda.gov/common/help/en/support/meeting_test.htm.* To get a quick overview of the Connect Pro program, visit *http://www.adobe.com/go/connectpro_overview.* (FDA has verified the Web site addresses in this document, but FDA is not responsible for any subsequent changes to the Web sites after this document publishes in the **Federal Register**.)

Comments: FDA is holding this public workshop to obtain feedback on CDRH's guidance development and guidance prioritization processes. In order to permit the widest possible opportunity to obtain public comment, FDA is soliciting either electronic or written comments on all aspects of the public workshop topics. The deadline for submitting comments related to this public workshop is July 7, 2014.

Regardless of attendance at the public workshop, interested persons may submit either electronic comments regarding this document to *http://www.regulations.gov* or written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. It is only necessary to send one set of comments. Please identify comments with the docket number found in brackets in the heading of this document. Received comments may be viewed in person in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at *http://www.regulations.gov.*

Transcripts: Please be advised that as soon as a transcript is available, it will be accessible at *http://www.regulations.gov.* It may be viewed at the Division of Dockets Management (see *Comments*). A transcript will also be available in either hardcopy or on CD–ROM, after submission of a Freedom of Information request. Written requests are to be sent to the Division of Freedom of Information (ELEM–1029), Food and Drug Administration, 12420 Parklawn Dr., Element Bldg., Rockville, MD 20857. A link to the transcripts will also be available approximately 45 days after the public workshop on the Internet at *http://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/default.htm.* (Select this public workshop from the posted events list.)

SUPPLEMENTARY INFORMATION:

I. Background

Guidance documents are documents issued by FDA and prepared for FDA staff and/or FDA stakeholders. They describe the Agency's interpretation of, or policy on, a regulatory issue (see § 10.115(b) (21 CFR 10.115(b))). Unlike statutes and regulations, guidances themselves do not create legally binding requirements (see § 10.115(d)). Nevertheless, guidance documents are important because they assist both staff and industry in understanding FDA's current thinking on certain topics. FDA's Good Guidance Practices regulation (§ 10.115) governs the development and issuance of guidance, and it gives interested parties a number