relationships that provide young people, who have an incarcerated parent with caring adult volunteers.

Additional information about this program and its purpose can be located on the following Web site: *http://www.acf.hhs.gov/programs/fysb.*

Contact for Further Information: Gloria Watkins, Family Youth and Services Bureau, 1250 Maryland Ave., SW., Washington, DC 20047. Telephone: (202) 205–9546. E-mail: *Gloria.Watkins@acf.hhs.gov*.

Dated: May 12, 2009.

Maiso L. Bryant,

Acting Commissioner, Administration on Children, Youth and Families.

[FR Doc. E9–11816 Filed 5–20–09; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0210]

Temporary Deferment of Activities Relating to Medical Device Submissions

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the Center for Devices and Radiological Health (CDRH) will be moving from various Rockville, Maryland locations to Building 66 at 10903 New Hampshire Avenue in Silver Spring, Maryland from approximately mid May 2009 until the beginning of August 2009. Offices will progressively move over weekends during this period. Specifically, moves will occur on Friday, Saturday, and Sunday except on holiday weekends. During the period required for relocation of files, equipment, and agency personnel, the Center for Devices and Radiological Health will not officially receive premarket submissions on the Friday of a move weekend and the Monday after a move weekend.

FOR FURTHER INFORMATION CONTACT: Marjorie Shulman, Center for Devices and Radiological Health (HFZ–404), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 240–276–4186 or *Marjorie.shulman@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

CDRH is responsible for activities under sections 510, 513, 515, and 520 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360, 360c, 360e, and 360j). These activities include, but are not limited to:

1. Advising the Director, CDRH, and other FDA officials on all medical device submissions, such as premarket notification submissions under section 510(k) of the act, device classifications under section 513 of the act, premarket approval applications (PMA's) and product development protocols (PDP's) under section 515 of the act, and clinical investigations under section 520 of the act;

2. Determining substantial equivalence for premarket notification submissions;

3. Planning, conducting, and coordinating CDRH actions regarding PMA's, PDP's, and investigational device exemption approvals, denials, or withdrawals of approval;

4. Monitoring sponsors' compliance with regulatory requirements; and

5. Conducting a continuing review, surveillance, and medical evaluation of the labeling, clinical experience, and required reports submitted by sponsors holding approved applications.

In an effort to consolidate CDRH offices, FDA is moving various CDRH offices from their present Rockville, Maryland locations to Building 66 at 10903 New Hampshire Avenue in Silver Spring, Maryland. Offices will progressively move, during weekends, during this period. Specifically, moves will occur on Friday, Saturday, and Sunday except on holiday weekends. During the period required for relocation of files, equipment, and agency personnel, the agency, specifically the Center for Devices and Radiological Health, will not officially receive submissions on the Friday of a move weekend and the Monday after a move weekend. Although mail will be delivered to a CDRH address during the move, CDRH will not be able to receive it on Fridays and Mondays, and will have limited capacity on Tuesday. Accordingly, mail delivered on Friday or Monday will be logged in on a staggered basis to preserve equity in the order of receipt and manageability of the accumulated workload. Specifically, mail delivered on Friday or Monday will be received on Tuesday and mail delivered on Tuesday will be received by Wednesday. Mail delivered on Wednesdays and Thursdays will remain unaffected.

The new mailing address for submissions and updated telephone contact information may be found by accessing *www.fda.gov/cdrh/whiteoakmove*.

II. Comments

Persons who may be affected by this temporary deferment should contact FDA with any questions they may have regarding CDRH's move to the White Oak, Maryland. These persons should call CDRH's Division of Small Manufacturers, International, and Consumer Assistance at 800–638–2041 (in Maryland, 240–276–3150).

Dated: May 13, 2009.

Daniel G. Schultz,

Director, Center for Devices and Radiological Health.

[FR Doc. E9–11840 Filed 5–20–09; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Accreditation and Approval of Intertek USA, Inc., as a Commercial Gauger and Laboratory

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security.

ACTION: Notice of accreditation and approval of Intertek USA, Inc., as a commercial gauger and laboratory.

SUMMARY: Notice is hereby given that, pursuant to 19 CFR 151.12 and 19 CFR 151.13, Intertek USA, Inc., 101 20th Street South, Texas City, TX 77590, has been approved to gauge and accredited to test petroleum and petroleum products, organic chemicals and vegetable oils for customs purposes, in accordance with the provisions of 19 CFR 151.12 and 19 CFR 151.13. Anyone wishing to employ this entity to conduct laboratory analyses and gauger services should request and receive written assurances from the entity that it is accredited or approved by the U.S. Customs and Border Protection to conduct the specific test or gauger service requested. Alternatively, inquires regarding the specific test or gauger service this entity is accredited or approved to perform may be directed to the U.S. Customs and Border Protection by calling (202) 344–1060. The inquiry may also be sent to *cbp.labhq@dhs.gov.* Please reference the Web site listed below for a complete listing of CBP approved gaugers and accredited laboratories.*http://cbp.gov/xp/cgov/import/operations_support/labs_scientific_svcs/commercial_gaugers/*.

DATES: The accreditation and approval of Intertek USA, Inc., as commercial gauger and laboratory became effective

on February 18, 2009. The next triennial inspection date will be scheduled for February 2012.

FOR FURTHER INFORMATION CONTACT: Anthony Malana, Laboratories and Scientific Services, U.S. Customs and Border Protection, 1300 Pennsylvania Avenue, NW., Suite 1500N, Washington, DC 20229, 202–344–1060.

Dated: May 15, 2009.

Ira S. Reese,

Executive Director, Laboratories and Scientific Services.

[FR Doc. E9–11932 Filed 5–20–09; 8:45 am]

BILLING CODE 9111–14–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Information Collection Activities: Deferral of Duty on Large Yachts Imported for Sale

AGENCY: U.S. Customs and Border Protection (CBP), Department of Homeland Security.

ACTION: 60-Day notice and request for comments; Extension of an existing information collection: 1651–0080.

SUMMARY: As part of its continuing effort to reduce paperwork and respondent burden, CBP invites the general public and other Federal agencies to comment on an information collection requirement concerning the Deferral of Duty on Large Yachts Imported for Sale. This request for comment is being made pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104–13; 44 U.S.C. 3505(c)(2)).

DATES: Written comments should be received on or before July 20, 2009, to be assured of consideration.

ADDRESSES: Direct all written comments to U.S. Customs and Border Protection, Attn: Tracey Denning, Office of Regulations and Rulings, 799 9th Street, NW., 7th Floor, Washington, DC 20229–1177.

FOR FURTHER INFORMATION CONTACT: Requests for additional information should be directed to Tracey Denning, U.S. Customs and Border Protection, Office of Regulations and Rulings, 799 9th Street, NW., 7th Floor, Washington, DC 20229–1177, at 202–325–0265.

SUPPLEMENTARY INFORMATION: CBP invites the general public and other Federal agencies to comment on proposed and/or continuing information collections pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104–13; 44 U.S.C. 3505(c)(2)). The comments should address: (a) Whether the 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 estimates of the burden of the collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden including the use of automated collection techniques or the use of other forms of information technology; and (e) estimates of capital or start-up costs and costs of operations, maintenance, and purchase of services to provide information. The comments that are submitted will be summarized and included in the request for Office of Management and Budget (OMB) approval. All comments will become a matter of public record. In this document the CBP is soliciting comments concerning the following information collection:

Title: Deferral of Duty on Large Yachts Imported for Sale.

OMB Number: 1651–0080.

Form Number: None.

Abstract: Section 2406(a) of the Miscellaneous Trade and Technical Corrections Act of 1999 provides that an otherwise dutiable ''large yacht'' may be imported without the payment of duty if the yacht is imported with the intention to offer for sale at a boat show in the U.S.

Current Actions: There are no changes to the information collection. This submission is being made to extend the expiration date.

Type of Review: Extension (without change).

Affected Public: Business and non-profit institutions.

Estimated Number of Respondents: 100.

Estimated Time per Respondent: 1 hour.

Estimated Total Annual Burden Hours: 100.

Dated: May 14, 2009.

Tracey Denning,

Agency Clearance Officer, Customs and Border Protection.

[FR Doc. E9–11771 Filed 5–20–09; 8:45 am]

BILLING CODE 9111–14–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Citizenship and Immigration Services

Agency Information Collection Activities: Form I–643, Extension of an Existing Information Collection; Comment Request

ACTION: 60-Day Notice of Information Collection Under Review: Form I–643, Health and Human Services Statistical Data for Refugee/Asylee Adjusting Status; OMB Control No. 1615–0070.

The Department of Homeland Security, U.S. Citizenship and Immigration Services has submitted the following information collection request for review and clearance in accordance with the Paperwork Reduction Act of 1995. The information collection is published to obtain comments from the public and affected agencies. Comments are encouraged and will be accepted for sixty days until July 20, 2009.

During this 60-day period, USCIS will be evaluating whether to revise the Form I–643. Should USCIS decide to revise the Form I–634, we will advise the public when we publish the 30-day notice in the **Federal Register** in accordance with the Paperwork Reduction Act. The public will then have 30 days to comment on any revisions to the Form I–643.

Written comments and suggestions regarding the item contained in this notice, and especially regarding the estimated public burden and associated response time, should be directed to the Department of Homeland Security (DHS), USCIS, Chief, Regulatory Products Division, Clearance Office, 111 Massachusetts Avenue, NW., Washington, DC 20529–2210. Comments may also be submitted to DHS via facsimile to 202–272–8352, or via e-mail at *rfs.regs@dhs.gov.* When submitting comments by e-mail, please add the OMB Control Number 1615–0070 in the subject box.

Written comments and suggestions from the public and affected agencies concerning the collection of information should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;