Statutory Authority: This program is authorized by—

(A) Section 462 of the Homeland Security Act of 2002, which in March 2003, transferred responsibility for the care and custody of Unaccompanied Alien Children from the Commissioner of the former Immigration and Naturalization Service to the Director of ORR of HHS.

(B) The Flores Settlement Agreement, Case No. CV85–4544–RJK (C. D. Cal. 1996), and the William Wilberforce Trafficking Victims Protection Reauthorization Act of 2008 (Pub. L. 110–457), which authorizes post release services under certain conditions to eligible children. All programs must comply with the Flores Settlement Agreement, Case No. CV85–4544–RJK (C.D. Cal. 1996), pertinent regulations and ORR policies and procedures.

Christopher Beach,

Senior Grants Policy Specialist, Division of Grants Policy, Office of Administration.

[FR Doc. 2017–12627 Filed 6–16–17; 8:45 am]

BILLING CODE 4184–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.568]

Reallotment of Fiscal Year 2016 Funds for the Low Income Home Energy Assistance Program (LIHEAP)

AGENCY: Division of Energy Assistance, Office of Community Services (OCS), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS).

ACTION: Notice of determination concerning funds available for reallotment.

SUMMARY: Notice is hereby given of a preliminary determination that funds from the fiscal year (FY) 2016 Low Income Home Energy Assistance Program (LIHEAP) are available for reallotment to states, territories, tribes, and tribal organizations that received FY 2017 direct LIHEAP grants. No subgrantees or other entities may apply for these funds.

DATES: Submit comments on or before July 19, 2017.

ADDRESSES: Comments may be submitted to: J. Janelle George, Acting Director, Office of Community Services, 330 C Street SW., 5th Floor, Mail Room 5425, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Lauren Christopher, Director, Division of Energy Assistance, Office of Community Services, 330 C Street SW., 5th Floor, Mail Room 5425, Washington, DC 20201; telephone (202) 401–4870; email: *lauren.christopher@acf.hhs.gov.*

SUPPLEMENTARY INFORMATION: Section 2607(b)(1) of the Low Income Home Energy Assistance Act (the Act), (42 U.S.C. 8626(b)(1)) requires that, if the Secretary of HHS determines that, as of September 1 of any fiscal year, an amount in excess of 10 percent of the amount awarded to a grantee for that fiscal year (excluding Leveraging and REACH funds) will not be used by the grantee during that fiscal year, then the Secretary must notify the grantee and publish a notice in the **Federal Register** that such funds may be reallotted to LIHEAP grantees during the following fiscal year. If reallotted, the LIHEAP block grant allocation formula will be used to distribute the funds. No funds may be allotted to entities that are not direct LIHEAP grantees during FY 2017.

It has been determined that $3,253,866 in LIHEAP funds may be available for reallotment during FY 2017. This determination is based on FY 2016 Carryover and Reallotment Reports, which showed that 15 grantees reported reallotment funds (State of Arkansas, Association of Village Council Presidents, Cocopah Tribe of Arizona, Eastern Band of Cherokee Indians, State of Georgia, Hoh Indian Tribe, Kalispel Indian Community of the Kalispel Reservation, Oglala Sioux Tribe, Passamaquoddy Tribe at Pleasant Point, Poarch Band of Creeks, Quinault Indian Nation, Sault Ste. Marie Tribe of Chippewa Indians, The Chickasaw Nation, Three Affiliated Tribes of the Ft. Berthold Reservation, and the State of Vermont). Grantees submitted the FY 2016 Carryover and Reallotment Reports to OCS, as required by regulations applicable to LIHEAP at 45 CFR 96.81(b).

The LIHEAP statute allows grantees who have funds unobligated at the end of the federal fiscal year for which they are awarded to request that they be allowed to carry over up to 10 percent of their full-year allotments to the next federal fiscal year. Funds in excess of this amount must be returned to HHS and are subject to reallotment under section 2607(b)(1) of the Act (42 U.S.C. 8626(b)(1)). The amount described in this notice was reported by grantees as unobligated FY 2016 funds in excess of the amount that these grantees could carry over to FY 2017.

In accordance with section 2607(b)(3) of the Act (42 U.S.C. 8626(b)(3)), comments will be accepted for a period of 30 days from the date of publication of this notice.

After considering any comments submitted, all current LIHEAP grantees will be notified of the final reallotment amount redistributed to them for obligation in FY 2017. This decision will be published in an Information Memorandum that gets posted to ACF's Web site.

If funds are reallotted, they will be allocated in accordance with section 2604 of the Act (42 U.S.C. 8623) and must be treated by LIHEAP grantees receiving them as an amount appropriated for FY 2017. As FY 2017 funds, they will be subject to all requirements of the Act, including section 2607(b)(2) (42 U.S.C. 8626(b)(2)), which requires that a grantee obligate at least 90 percent of its total block grant allocation for a fiscal year by the end of the fiscal year for which the funds are appropriated, that is, by September 30, 2017.

ESTIMATED REALLOTMENT AMOUNTS OF FY 2016 LIHEAP FUNDS

Grantee name	Reallotment amount
Arkansas	$726,214
Association of Village Council Presidents	169,410
Cocopah Tribe of Arizona	18
Eastern Band of Cherokee Indians	18,728
Georgia	1,035,739
Hoh Indian Tribe	1,907
Kalispel Indian Community of the Kalispel Reservation	1,558
Oglala Sioux Tribe	23,396
Passamaquoddy Tribe at Pleasant Point	107
Poarch Band of Creeks	70,819

ESTIMATED REALLOTMENT AMOUNTS OF FY 2016 LIHEAP FUNDS—Continued

Grantee name	Reallotment amount
Quinault Indian Nation	4,091
Sault Ste. Marie Tribe of Chippewa Indians	4
The Chickasaw Nation	195,952
Three Affiliated Tribes of the Ft. Berthold Reservation	348,035
Vermont	657,888
Total	3,253,866

Statutory Authority: 42 U.S.C. 8626.

Elizabeth Leo,

Grants Policy Specialist, Division of Grants Policy, Office of Administration.

[FR Doc. 2017–12675 Filed 6–16–17; 8:45 am]

BILLING CODE 4184–80–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0015]

Agency Information Collection Activities; Proposed Collection; Comment Request; Orphan Drugs; Common European Medicines Agency/ Food and Drug Administration Application Form for Orphan Drug Medicinal Product Designation

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on Common European Medicines Agency (EMA)/FDA Application Form for Orphan Drug Medicinal Product Designation (Form FDA 3671).

DATES: Submit either electronic or written comments on the collection of information by August 18, 2017.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before August 18, 2017. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of August 18, 2017. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://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 *https://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):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, 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–2011–N–0015 for ''Orphan Drugs; Common EMA/FDA Application Form for Orphan Medicinal Product Designation (Form FDA 3671)—21 CFR part 316.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff 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 *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. 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: *https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://*