A. Purpose

The General Services Administration Acquisition Regulation (GSAR) requires construction and building services contractors to submit a release of claims before final payment is made to ensure contractors are paid in accordance with their contract requirements and for work performed. GSA Form 1142, Release of Claims is used to achieve uniformity and consistency in the release of claims process.

B. Annual Reporting Burden

Respondents: 1,427.

Responses per Respondent: 1.

Annual Responses: 1,427.

Hours per Response: 0.50.

Total Burden Hours: 714.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the GSAR, and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; and ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the Regulatory Secretariat Division (MVCB), at *GSARegSec@gsa.gov.*

Please cite OMB Control No. 3090–0080; Release of Claims for Construction and Building Service Contracts, in all correspondence.

Jeffrey A. Koses,

Senior Procurement Executive, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2024–10636 Filed 5–14–24; 8:45 am]

BILLING CODE 6820–61–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for Office of Management and Budget Review; Head Start REACH: Strengthening Outreach, Recruitment, and Engagement Approaches With Families—Mixed Methods Study (New Collection)

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, U.S. Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Administration for Children and Families (ACF) within the U.S. Department of Health and Human Services (HHS) is proposing to collect data on different approaches that Head Start programs use to recruit, select, and enroll families, and the ways in which such practices reflect programs' community contexts. We are not attempting to recruit a nationally representative sample. Instead, the study will aim to obtain a variety of eligibility, recruitment, selection, enrollment, and attendance (ERSEA) practices and experiences to explore how these practices and experiences intersect with different adversities, demographic characteristics, and community contexts.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about 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.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. You can also obtain copies of the proposed collection of information by emailing *OPREinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: Building on information collected previously through case studies (OMB #0970–0580), the Head Start REACH: Strengthening Outreach, Recruitment, and Engagement Approaches with Families Project is proposing to conduct a mixed-methods study to expand understanding of (1) how Head Start programs implement recruitment, selection, and enrollment practices; and (2) the ways in which practices reflect programs' community contexts. The mixed-methods study would achieve several goals including (1) providing in-depth contextual information about recruitment, selection, and enrollment practices and experiences; (2) identifying promising recruitment, selection, and enrollment practices and experiences; and (3) informing training and technical assistance regarding recruitment, selection, and enrollment challenges and needs. We will aim to collect information from 60 Head Start and Early Head Start programs in 15 geographic areas in states, from Head Start regions I–X, located in census tracts where the rate of deep poverty is high.

We will collect information about the characteristics of families in Head Start programs and their communities; programs' enrollment numbers and goals; programs' use and perceived effectiveness of and challenges with recruitment, selection, and enrollment practices; promising recruitment, selection, and enrollment practices for potential future replication; families' reasons for choosing Head Start and experiences with and perceptions of recruitment, selection, and enrollment practices; and how community partner staff support recruitment, selection, and enrollment of families into Head Start. The findings are intended to help Head Start programs understand how to support the needs of families facing adversities. We will disseminate findings in a report, research brief, and presentations or briefings.

Respondents: Head Start program directors (one per program), ERSEA lead staff (one per program), Head Start parents/caregivers (up to 10 per program), and staff from community organizations with which Head Start programs partner for ERSEA activities (up to 3 per program).

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total/annual burden (in hours)
Program director survey (*Instrument 1)*	60	1	0.17	10.2
ERSEA lead staff survey (*Instrument 2)*	60	1	0.75	45
Onsite coordination [a]	60	1	1.5	90
Head Start parent/caregiver survey *(Instrument 3)*	600	1	0.5	300
Community partner survey *(Instrument 4)*	180	1	0.25	45
ERSEA lead staff focus group guide *(Instrument 5)*	24	1	1.5	36
Estimated Total Annual Burden Hours				526.2

[a] There is no instrument associated with this activity. We will ask each program director to nominate a staff person who will help coordinate data collection activities. This line accounts for the time of the onsite coordinator.

Authority: Head Start Act Section 640 [42 U.S.C. 9835].

Mary C. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2024–10578 Filed 5–14–24; 8:45 am]

BILLING CODE 4184–22–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2020–N–0026]

Issuance of Priority Review Voucher; Rare Pediatric Disease Product; OJEMDA (tovorafenib)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the issuance of a priority review voucher to the sponsor of a rare pediatric disease product application. The Federal Food, Drug, and Cosmetic Act (FD&C Act) authorizes FDA to award priority review vouchers to sponsors of approved rare pediatric disease product applications that meet certain criteria. FDA is required to publish notice of the award of the priority review voucher. FDA has determined that OJEMDA (tovorafenib), approved on April 23, 2024, manufactured by Day One Biopharmaceuticals, Inc., meets the criteria for a priority review voucher.

FOR FURTHER INFORMATION CONTACT: Cathryn Lee, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002, 301–796–1394.

SUPPLEMENTARY INFORMATION: FDA is announcing the issuance of a priority review voucher to the sponsor of an approved rare pediatric disease product application. Under section 529 of the FD&C Act (21 U.S.C. 360ff), FDA will award priority review vouchers to sponsors of approved rare pediatric disease product applications that meet certain criteria. FDA has determined that OJEMDA (tovorafenib), manufactured by Day One Biopharmaceuticals, Inc., meets the criteria for a priority review voucher. OJEMDA (tovorafenib) is indicated for the treatment of patients 6 months of age and older with relapsed or refractory pediatric low-grade glioma harboring a BRAF fusion or rearrangement, or BRAF V600 mutation.

For further information about the Rare Pediatric Disease Priority Review Voucher Program and for a link to the full text of section 529 of the FD&C Act, go to *http://www.fda.gov/ForIndustry/DevelopingProductsforRareDiseasesConditions/RarePediatricDiseasePriorityVoucherProgram/default.htm.* For further information about OJEMDA (tovorafenib), go to the ''*Drugs@FDA*'' website at *http://www.accessdata.fda.gov/scripts/cder/daf/.*

Dated: May 9, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024–10583 Filed 5–14–24; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2024–N–2032]

Agency Information Collection Activities; Proposed Collection; Comment Request; Food and Cosmetic Export Certificate Application Process

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (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 information collection associated with export certificate applications for FDA-regulated human food and cosmetic products.

DATES: Either electronic or written comments on the collection of information must be submitted by July 15, 2024.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of July 15, 2024. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered