level implementing agencies will collect qualitative data to understand contextual factors and the impetus behind the design and implementation of the initiatives. Finally, cost workbooks completed by center-based CCEE setting administrators will collect cost data to assess the costs associated with implementing the initiative. This information collection will support ACF and the CCEE field in understanding whether workforce support strategies that increase compensation affect the retention and well-being of the CCEE workforce. This information will help to inform federal, state, and local initiatives to build and retain a qualified CCEE workforce.

Respondents: CCEE center-based directors, administrators, teachers; CCEE home-based owners and caregivers; CCEE key informants.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Avg. burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
1. Follow-up center director survey	75	2	0.75	113	75
2. Follow-up lead and assistant teacher survey	1,000	2	0.75	1,500	1,000
3. Follow-up home-based owner and caregiver survey	95	2	0.75	143	95
4. One-on-one center director interview	15	1	1	15	10
5. One-on-one lead and assistant teacher interview	25	1	1	25	17
6. One-on-one home-based owner and caregiver interview	25	1	1.5	38	25
7. One-on-one key informant interview	5	1	1	5	3
8. Center-based setting costs workbook	16	1	5	80	53

Estimated Total Annual Burden Hours: 1,278.

Authority: Head Start Act 640 [42 U.S.C. 9835] and 649 [42 U.S.C. 9844]; appropriated by the Consolidated Appropriations Act of 2022. Head Start Act as amended by the Improving Head Start for School Readiness Act of 2007 (Pub. L. 110 134).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2023–15555 Filed 7–21–23; 8:45 am]

BILLING CODE 4184–22–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; ORR–3 and ORR–4 Report Forms for the Unaccompanied Refugee Minors Program (OMB #0970–0034)

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

ACTION: Request for public comments.

SUMMARY: The Office of Refugee Resettlement (ORR) is requesting a 3-year extension of the ORR–3 and ORR–4 Report Forms for the Unaccompanied Refugee Minors Program (OMB#: 0970–0034, expiration 02/29/2024). There are no changes requested to the report forms, but ORR proposes minor revisions to the form instructions to improve clarity, including additional guidance for providers on how to assess youth functioning.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: You can obtain copies of the proposed collection of information and submit comments by emailing *infocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The ORR–3 Report is submitted within 30 days of the minor's initial placement in the state, within 60 days of a reportable change in the minor's case (*e.g.,* change in legal responsibility, change in foster home placement, change in immigration data), and within 60 days of termination from the program. The ORR–4 Report is submitted every 12 months beginning on the first anniversary of the initial placement date, to record outcomes of the minor's progress.

Respondents: Unaccompanied Refugee Minors (URM) State Agencies, URM Provider Agencies, and youth participants.

Annual Burden Estimates

URM STATE AGENCIES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
ORR–3 URM Placement Report	15	432	0.25	1,620	540
ORR–4 URM Outcomes Report	15	282	0.50	2,115	705

Estimated Total Annual Burden Hours (URM State Agencies): 1,245.

URM PROVIDER AGENCIES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
ORR–3 URM Placement Report	24	270	0.50	3,240	1,080
ORR–4 URM Outcomes Report	24	162	1.0	3,888	1,296

Estimated Total Annual Burden Hours (URM Provider Agencies): 2,376.

YOUTH PARTICIPANTS

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
ORR–4 URM Outcomes Report	1032	3	0.50	1,548	516

Estimated Total Annual Burden Hours (Youth Participants): 516.

Total Estimated Annual Burden Hours: 4,137.

Comments: The Department specifically requests comments on (a) whether the proposed 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 estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority: 8 U.S.C. 1522(d).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2023–15556 Filed 7–21–23; 8:45 am]

BILLING CODE 4184–89–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–N–2873]

Public Meeting and Listening Session for Developing the Food and Drug Administration's Center for Tobacco Products' Strategic Plan; Request for Comments

AGENCY: Food and Drug Administration, Department of Health and Human Services (HHS).

ACTION: Notification of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the following virtual listening session entitled ''Public Meeting and Listening Session for Developing FDA's Center for Tobacco Products' Strategic Plan.'' The purpose of the listening session is to obtain feedback on the proposed strategic goals that are being used to develop FDA's Center for Tobacco Products' (CTP) comprehensive Strategic Plan. FDA will provide information on the proposed goals and provide the public an opportunity to provide open public comment.

DATES: The virtual listening session will be held on August 22, 2023, beginning at 10 a.m. Eastern Time. Additional details, such as the time of the listening session and registration information, is available at: *https://www.fda.gov/tobacco-products/ctp-newsroom/listening-session-developing-fdas-center-tobacco-products-strategic-plan-08222023.* All requests to make open public comment must be received by August 14, 2023, at 11:59 p.m. Eastern Time. Either electronic or written comments on this listening session must be submitted to the docket by August 29, 2023. See the **SUPPLEMENTARY INFORMATION** section for registration date and information.

ADDRESSES: The listening session will be held virtually and more information will be posted here: *https://www.fda.gov/tobacco-products/ctp-newsroom/listening-session-developing-fdas-center-tobacco-products-strategic-plan-08222023.*

You may submit written 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 August 29, 2023. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received 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