• Disseminate insights related to human trafficking cases and trends to inform anti-trafficking strategies and policies; and

• Provide information to Congress, other federal agencies, stakeholders, the public, and other countries on the aggregate outputs and outcomes of the NHTH operations.

In accordance with the Paperwork Reduction Act of 1995 (Pub. L. 104–13) and OMB regulations at 5 CFR part 1320, ACF published a notice in the **Federal Register** to announce the agency's intention to request OMB review of this information collection activity and provide a sixty-day period for public comment (86 FR 38489). During the notice and comment period, one comment was received from the NHTH grantee. The comment did not pertain to the burden estimate for respondents (signalers to the NHTH), rather the burden on the recordkeeper (the NHTH grantee).

To be responsive to this comment and reduce the burden on the recordkeeper, OTIP modified the collection to remove several of the data elements that were initially proposed. Where OTIP has requested any new data (*e.g.,* data the grantee is not already providing to OTIP as a condition of award), particularly, for existing data to be further disaggregated, it is in the interest of allowing OTIP to:

• Monitor performance and operational issues;

• Generate more timely insights into trends related to victim demographics and service needs, and the impact of particular intra- and inter-agency efforts, messaging campaigns, trainings, and other anti-trafficking efforts on NHTH signals, and;

• Respond to congressional inquiries and other ad hoc inquiries without submitting burdensome individual requests to the NHTH.

Respondents: Potential victims, representatives of governmental entities, law enforcement, first responders, members of the community, representatives of nongovernmental entities providing social, legal, or protective services to individuals in the United States who may have been subjected to severe forms of trafficking in persons utilize the NHTH as signalers.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents (signalers)	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
National Human Trafficking Hotline (NHTH) Performance Indicators	585,300	1	0.43333333	253,630	84,543

Estimated Total Annual Burden Hours: 84,543.

Authority: 22 U.S.C. 7105.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–27646 Filed 12–21–21; 8:45 am]

BILLING CODE 4184–47–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Unaccompanied Children (UC) Program Budget Workbook Template (New Collection)

AGENCY: Office of Refugee Resettlement, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families' (ACF) Office of Refugee Resettlement (ORR) is requesting clearance for the proposed new collection titled ''UC Program Budget Workbook'' to streamline budget details and justifications of applicants to funding opportunities.

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.

SUPPLEMENTARY INFORMATION:

Description: The UC Program Budget Workbook will streamline the budget detail and justification documentation for applicants to any upcoming Notice of Funding Opportunity (NOFO). This new information collection will provide guidance to the applicant as well as a fillable form to insert calculations and budget line items. With the assistance of this template, the review of applications will be expedited since documentation will be clearer and more unified. Additionally, this will facilitate the completion of applications that may not otherwise be completed due to lack of budget documentation guidance in past NOFOs.

Respondents: New and existing applicants to NOFOs for residential services for UC.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
UC Program Budget Form	120	3	90	32,400	10,800

Estimated Total Annual Burden Hours: 10,800.

Authority: 8 U.S.C. 1522 of the Immigration and Nationality Act (the Act) (Title IV, Sec. 412 of the Act) and 45 CFR 400.28(b).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–27767 Filed 12–21–21; 8:45 am]

BILLING CODE 4184–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–D–0756]

Validation and Verification of Analytical Testing Methods Used for Tobacco Products; Draft Guidance for Industry; Availability; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for comments.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing the availability of a draft guidance for industry entitled ''Validation and Verification of Analytical Testing Methods used for Tobacco Products'' and requesting comments, including scientific and other information, concerning the recommendations set forth in the draft guidance. The draft guidance, when finalized, would provide information and recommendations related to the validation and verification of analytical test methods, including analytical testing of tobacco product constituents, ingredients, and additives, as well as stability testing of tobacco products. This draft guidance would help industry produce more consistent and reliable analytical data used to support regulatory submissions for finished tobacco products.

DATES: Submit either electronic or written comments on the draft guidance by February 22, 2022 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance at any time as follows:

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–2021–D–0756 for ''Validation and Verification of Analytical Testing Methods used for Tobacco Products.'' Received comments 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, 240–402–7500.

- 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.govinfo.gov/content/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://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of the draft guidance to the Center for Tobacco Products, Food and Drug Administration, 10903 New Hampshire Ave., Document Control Center, Bldg. 71, Rm. G335, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your request or include a fax number to which the guidance document may be sent. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

FOR FURTHER INFORMATION CONTACT: Nathan Mease or Matthew Brenner, Center for Tobacco Products, Food and Drug Administration, Document Control Center, 10903 New Hampshire Ave., Bldg. 71, Rm. G335, Silver Spring, MD 20993–0002, 1–877–287–1373, email: *CTPRegulations@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

We are announcing the availability of a draft document entitled ''Validation and Verification of Analytical Testing Methods used for Tobacco Products; Draft Guidance for Industry.'' This draft guidance, when finalized, provides information and recommendations on how tobacco product manufacturers can produce validation and verification data for the analytical procedures and