Department's information collection requirements and provide the requested data in the desired format. The Department is soliciting comments on the proposed information collection request (ICR) that is described below. The Department is especially interested in public comment addressing the following issues: (1) is this collection necessary to the proper functions of the Department; (2) will this information be processed and used in a timely manner; (3) is the estimate of burden accurate; (4) how might the Department enhance the quality, utility, and clarity of the information to be collected; and (5) how might the Department minimize the burden of this collection on the respondents, including through the use of information technology. Please note that written comments received in response to this notice will be considered public records.

Title of Collection: International Resource Information System (IRIS).

OMB Control Number: 1840–0759.

Type of Review: A revision of a currently approved ICR.

Respondents/Affected Public: Private Sector; Individuals and Households.

Total Estimated Number of Annual Responses: 6,126.

Total Estimated Number of Annual Burden Hours: 12,928.

Abstract: Information Resource Information System (IRIS) is an online performance reporting system for grantees of International and Foreign Language Education (IFLE) programs. The site also allows for IFLE program officers to process overseas language requests, travel authorization requests, and grant activation requests. IRIS keeps a record of these requests and also of Foreign Language and Area Studies (FLAS) Fellowship recipients and grantee performance reports.

The International and Foreign Language Education (IFLE) office within the Office of Postsecondary Education, of the U.S. Department of Education (the Department) administers 11 institutional and fellowship grant programs authorized under title VI of the Higher Education Act and section 102(b)(6) of the Mutual Educational and Cultural Exchange Act, also known as Fulbright-Hays (F–H). The purpose of these programs is to strengthen the capability and performance of American education in foreign languages and in area and international studies.

The International Resource Information System (IRIS) is an online database that is used for performance reporting by the grantees funded by these 11 programs. The purpose of IRIS is to provide a centralized and effective way of collecting, reporting, and analyzing annual performance data.

This Information Collection Request is a revision of the previously approved IRIS information collection. The IFLE office underwent a strategic assessment of each program using the specific statute, regulations, and program purposes. Our assessment was guided by the following objectives: reduce the reporting burden for grantees; reduce the burden hours for IFLE staff to review APRs; and, increase the quality and usefulness of the data collected.

The strategic assessment resulted in revisions to the program collections that will significantly reduce reporting burden across programs. We anticipate that the overall annual burden hours will be reduced by 64% from the previous collection clearance.

Dated: October 21, 2024.

Kun Mullan,

PRA Coordinator, Strategic Collections and Clearance, Governance and Strategy Division, Office of Chief Data Officer, Office of Planning, Evaluation and Policy Development.

[FR Doc. 2024–24768 Filed 10–23–24; 8:45 am]

BILLING CODE 4000–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Designation of a Class of Employees for Addition to the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention, Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: HHS gives notice of a decision to designate a class of employees from Metals and Controls Corp. in Attleboro, Massachusetts, as an addition to the Special Exposure Cohort (SEC) under the Energy Employees Occupational Illness Compensation Program Act of 2000.

FOR FURTHER INFORMATION CONTACT: Grady Calhoun, Director, Division of Compensation Analysis and Support, NIOSH, 1090 Tusculum Avenue, MS C–46, Cincinnati, OH 45226–1938, Telephone: (513) 533–6800. Information requests can also be submitted by email to *DCAS@CDC.GOV.*

SUPPLEMENTARY INFORMATION: On September 9, 2024, as provided for under 42 U.S.C. 7384*l*(14)(C), the Secretary of HHS designated the following class of employees as an addition to the SEC:

''All Atomic Weapons Employees who worked at Metals and Controls Corp. in Attleboro, Massachusetts, from January 1, 1968, through September 21, 1995, for a number of work days aggregating at least 250 work days, occurring either solely under this employment or in combination with work days within the parameters established for one or more other classes of employees included in the SEC.''

This designation is effective as of October 9, 2024, unless Congress provides otherwise prior to the effective date. After this effective date, HHS will publish a notice in the **Federal Register** reporting the addition of this class to the SEC or the result of any provision by Congress regarding the decision by HHS to add the class to the SEC.

Authority: 42 U.S.C. 7384q(b). 42 U.S.C. 7384*l*(14)(C).

John J. Howard,

Director, National Institute for Occupational Safety and Health.

[FR Doc. 2024–24643 Filed 10–23–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for Office of Management and Budget Review; Legal and Advocacy Services for Unaccompanied Children (Office of Management and Budget #0970–0565)

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), Administration for Children and Families (ACF), U.S. Department of Health and Human Services is inviting public comment on revisions to an approved information collection. The request consists of several forms that allow the Unaccompanied Children (UC) Program to provide legal and advocacy services to unaccompanied children.

DATES: *Comments due* November 25, 2024. The Office of Management and Budget (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 *infocollection@acf.hhs.gov.* Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: This request is to remove five forms from this collection, add two new forms, move two forms from a different information collection into this collection (with revisions to one of those forms), and revise three existing forms in this collection. ORR also proposes retitling this information collection ''Legal and Advocacy Services for Unaccompanied Children.''

Discontinued Forms

ORR plans to remove the following forms from this information collection:

1. Request for a Flores Bond Hearing (Form LRG–7), Motion to Request a Bond Hearing—Secure or Staff Secure Custody (Form LRG–8A), and Motion to Request a Bond Hearing—Non-Secure Custody (Form LRG–8B): The bond hearing process was replaced with a Risk Determination Hearing process under the UC Program Foundational Rule, 45 CFR 410. These forms were replaced with a new set of form specific to the RDH process, approved under OMB #0970–0633.

2. Motion for Change of Venue (Form L–7): This instrument was created for the UC Path case management system and was intended to be used for filing a motion for change of venue for children transferring to a different ORR care provider program. However, the UC Path system was never implemented, and this form has never been used. In addition, this function is performed by an entity that is party to the proceedings, typically the child's legal representative or Immigration and Customs Enforcement, because the decision to file a change of venue may affect the child's immigration case. A change of venue is filed for cases where a Notice to Appear has been filed. Since neither ORR nor its care provider programs perform this function, the form is not needed.

3. Post Legal Status Plan (Form L–8): The information collected in this form was incorporated into the Legal Services Plan section of the Category 4 Discharge Plan (Form R–9, currently approved under OMB #0970–0552). Therefore, ORR plans to discontinue this form.

New Forms

ORR plans to add the following new forms to this information collection:

1. Case Status Summary for Executive Office for Immigration Review (Form L–9): This form is completed by the Federal Field Specialist (FFS)or care provider and sent to the Executive Office for Immigration Review (EOIR) in advance of a child's immigration hearing. The form provides basic information needed to ensure that EOIR has accurate information on the child's case status. A copy of the form is also shared with the child's legal service provider or attorney of record and child advocate (if applicable).

2. Recommended States List (Form L–11): This form is completed by legal service providers for children transferring to a long-term foster care (LTFC) placement. The form provides a recommended list of preferred placement locations recommended by the legal service provider (LSP) based on the child's potential for immigration relief in each state, type of immigration relief, and status of court hearings or relief petitions. The LSP's recommendation is one of several factors ORR considers when making an LTFC placement determination.

Forms Transferred From a Different Information Collection

1. ORR plans to transfer the ''Notice of Administrative Review (Form P–18)'' into this information collection without revisions. This form is currently approved under OMB #0970–0554.

2. ORR plans to transfer the ''Child Advocate Recommendation and Appointment'' form into this information collection (currently approved under OMB #0970–0553). The currently approved version has been revised to move Section B: Recommendation and Appointment and Section C: ORR Approval into a separate form to better facilitate the referral, recommendation, and appointment process. The separate form containing the information collected in Sections B and C will be completed by fewer than 10 respondents and is, therefore, not subject to Paperwork Reduction Act and is not included in this request.

Additionally, ORR proposes the following revisions to assist its child advocate contractor in supporting referred children and making recommendations. ORR worked directly with the contractor to improve the form and incorporated recommendations the contractor submitted via public comment.

1. Rename the form ''Child Advocate Referral'' (Form L–12A).

2. Replace ''UC'' with ''child'' throughout the form.

3. Add fields for the Title, Email, and Phone Number of the referrer.

4. Make the following revisions related to the child's biographic information:

- Split the field to for the name of the child into three separate fields for First Name, Second, or Middle Name, and Last Name(s)
- Add fields for Also Known As (AKA) and Nicknames or Preferred Names
- Change the Gender field from an open text field to a dropdown field with options for Male, Female, and Nonbinary.
- Add a field for Other Language(s) Spoken

5. Add the following fields related to the child's entry into the United States:

- U.S. Port of Entry Where Child Entered
- Date of Apprehension by DHS

6. Make the following revisions related to the child's placement:

- Reword the Care Provider field to Current Care Provider Facility
- Reword the Admission Date field to Provider Admission Date
- Add the following fields:
  - Is the child in ORR custody?
  - Was the child at another ORR care provider facility?
  - If yes, provide the care provider facility name(s)
  - Add a field for Date of Entry into ORR Custody
  - Child's Length of Care in ORR Custody

7. Add the following fields related to the child's sponsor:

- Does the child have a sponsor?
- If yes, what category has ORR assigned to this sponsor?
- How many potential sponsors has the child had?

8. Add the following fields related to the child's legal representation:

- Does the child have legal representation?
- If yes, provide the following information for the legal representative: Name, Phone, Email

9. Revise the list of reasons for referral to better reflect the most common reasons child advocate referrals are made.

10. Add a text box where referrers can provide additional details regarding the reason for referral.

11. Revise the burden estimate to reflect the number of child advocate referrals made from April 2023 through

March 2024. The annual number of respondents increased from 216 to 300 and the annual number of responses per respondent increased from 5 to 19.

Revisions to Existing Forms

ORR plans to make the following revisions to existing forms in this information collection:

1. Request for Specific Consent to Juvenile Court Jurisdiction (Form L–1):

- Replace ''UC'' with ''child'' throughout the form
- Update language in the form's introductory text and Section D: Next Steps to align with the UC Program Foundational Rule (45 CFR 410).
- Reword the Name of Intended Guardian field to Name of Intended Individual or Entity to be Granted Custody to account for the fact that not all states use the word ''guardian'' to describe an individual or entity granted legal custody of a child. Revision is responsive to public comments.
- Revise the burden estimate to reflect the number of requests for specific consent received in FY 2023. The annual number of respondents decreased from 40 to 31.

2. Specific Consent Request Case Summary (Form L–2):

- Remove the instruction to complete an internal clearance form because that is no longer part of the process.
- Update the email address where the form is submitted.
- Add text fields for the email addresses of the case manager and FFS to better facilitate communication when UC Bureau headquarter staff have follow-up questions.
- Replace ''UC'' with ''child'' throughout the form
- Revise the available dropdown options for the Level of Care field to align with the UC Program Foundational Rule (45 CFR 410).
- Change the Gender field from an open text field to a dropdown field with options for Male, Female, and Nonbinary.
- Remove question 3 (If the child was released from ORR custody into the new custody situation, would there be any risk of escape?) and the mention of ''flight risk'' in Section C to align with the UC Program Foundational Rule. Revision is responsive to public comments.
- Revise the burden estimate to reflect the number of case summaries completed in FY 2023 and account for an increase in the number of care provider facilities. The annual number of respondents increased from 216 to 300 and the annual number of responses per respondent decreased from 0.2 to 0.1.

3. Acknowledgement of Receipt of Legal Resource Guide (Form LRG–4):

- Change form number from LRG–5 to LRG–4
- Retitle form ''Acknowledgement of Receipt of Legal Resource Guide'' (formerly titled ''Legal Service Provider List for UC in ORR Care'')
- Remove the information provided on the first page and the list of legal service providers and their contact information. ORR plans to incorporate this information into a separate document and children will acknowledge receipt of that document in this form.
- Revise the list of documents provided to children to reflect forthcoming revisions and consolidation of legal resource guide documents.
- Remove requirement for children to initial each list item to reduce burden for the child.
- Add instructions to put an ''X'' in the signature line in cases where the child is unable to sign the form and add a text field for the care provider to document the reason the child was unable to sign (*e.g.,* child is two years old). This will assist ORR in monitoring compliance with requirements to complete this form.
- Add a field for care provider program name.
- Revise the burden estimate to account for an increase in the number of care provider facilities and in the number of children placed in ORR care, and report the burden for care providers and unaccompanied children separately to improve accuracy of the estimate. The annual number of respondents increased from 216 to 300 for care providers and 121,669 unaccompanied child respondents were added. The annual number of responses per respondent increased from 556 to 817 for care providers and the responses per respondent for children is two (2).
- ORR plans to translate the form into Spanish and other languages commonly spoken by unaccompanied children.

Revisions to Burden Estimates Only for Existing Forms

1. *Notice of Attorney Representation (Form L–3):*

- Previously, the annual number of respondents was overestimated at 13,000. ORR is changing that estimate to 10,000 (which will still be higher than the number of forms submitted in the previous year) based on the actual number of children who received direct representation through ORR's legal service provider contractor and rounded up to account for an expected increase in direct representation and forms submitted by outside attorneys.

2. *UC Legal Information (Form L–4):*

- Revise the burden estimate to account for an increase in the number of care provider facilities and in the number of children placed in ORR care. The annual number of respondents increased from 216 to 300 and the annual number of responses per respondent increased from 241 to 406.

3. *Legal Service Provider Record (Form L–6):*

- Revise the burden estimate to account for an increase in the number of care provider facilities and in the number of children placed in ORR care. The annual number of respondents increased from 216 to 300 and the annual number of responses per respondent increased from 241 to 406.

Respondents: ORR grantee and contractor staff, unaccompanied children, parents/legal guardians of unaccompanied children, attorneys of record and legal service providers

Annual Burden Estimates:

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual total burden hours
Request for Specific Consent to Juvenile Court Jurisdiction (Form L–1)	31	1.0	0.25	8
Specific Consent Request Case Summary (Form L–2)	300	0.1	0.33	10
Notice of Attorney Representation (Form L–3A)	5,000	1.0	0.25	1,250
Notice of Legal Service Provider Screening (Form L–3B)	5,000	1.0	0.17	850
UC Legal Information (Form L–4)	300	406.0	1.00	121,800
Legal Service Provider Record (Form L–6)	300	406.0	0.08	9,744
Case Status Summary for Executive Office of Immigration Review (Form L–9)	300	5.0	0.17	255

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual total burden hours
Recommended States List (Form L–11)	60	10.0	0.33	198
Child Advocate Referral (Form L–12A)-Respondents	300	19.0	0.25	1,425
Child Advocate Referral (Form L–12A)-Recordkeepers	1	5,601.0	0.33	1,848
Acknowledgment of Receipt of Legal Resource Guide (LRG–4)-Unaccompanied Children	121,669	2.0	0.25	60,835
Acknowledgment of Receipt of Legal Resource Guide (LRG–4)-Care Providers	300	817.0	0.25	61,275
Notice of Administrative Review (Form P–18)	200	1.0	.83	166
Estimated Annual Burden Hours Total				259,664

Authority: 6 U.S.C. 279; 8 U.S.C. 1232; Flores v. Reno Settlement Agreement, No. CV85–4544–RJK (C.D. Cal. 1996).

Mary C. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2024–24588 Filed 10–23–24; 8:45 am]

BILLING CODE 4184–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2024–N–4470]

Agency Information Collection Activities; Proposed Collection; Comment Request; Antimicrobial Animal Drug Sales and Distribution

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) 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 the information collection provisions of our reporting and recordkeeping requirements for antimicrobial animal drug sales and distribution.

DATES: Either electronic or written comments on the collection of information must be submitted by December 23, 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 December 23, 2024. 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 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–2024–N–4470 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Antimicrobial Animal Drug Sales and Distribution.'' 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, 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.*