implementation of the Common Formats with members of the public, including software developers and other interested parties. Agenda topics will include discussion of ICD–11's incorporation of patient safety. Active participation and discussion by meeting participants is encouraged.

AHRQ requests that interested persons send an email to *SDMeetings@infinityconferences.com* for registration information. Before the meeting, an agenda and logistical information will be provided to registrants.

Dated: June 22, 2023.

Marquita Cullom,

Associate Director.

[FR Doc. 2023–13716 Filed 6–27–23; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Child Care and Development Fund Plan Preprint for States/Territories for FFY 2025–2027 (ACF–118) and Extension of Child Care and Development Fund Plan Preprint for States/Territories for FFY 2022–2024 (OMB #0970–0114)

AGENCY: Office of Child Care; Administration for Children and Families; U.S. Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Administration for Children and Families (ACF) is requesting an extension without changes of the form ACF–118: Child Care and Development Fund Plan Preprint for States/Territories for FFY 2022–2024 (OMB #0970–0114, expiration 02/29/2024), and an additional 3-year extension of the form ACF–118: Child Care and Development Fund Plan Preprint for States/Territories for FFY 2025–2027. There are changes requested to the form ACF–118: Child Care and Development Fund Plan Preprint for States/Territories for FFY 2025–2027 to improve formatting, collect additional information about program implementation, and streamline questions.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of section 3506(c)(2)(A) 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 Child Care and Development Fund (CCDF) Plan (the Plan) for States and Territories is required from each CCDF Lead agency in accordance with Section 658E of the Child Care and Development Block Grant Act of 1990 (CCDBG Act), as amended, CCDBG Act of 2014 (Pub. L. 113–186), and 42 U.S.C. 9858. The Plan, submitted on the ACF–118, is required triennially and remains in effect for 3 years. The Plan provides ACF and the public with a description of and assurance about the states' and territories' child care programs. These Plans are the applications for CCDF funds.

At this time, the ACF Office of Child Care (OCC) is proposing an extension of the approval of the currently approved CCDF Plan Preprint for FFY 2022–2024 to allow states and territories to continue to submit amendments through September 30, 2024, as required. There are no changes proposed to the FFY 2022–2024 Plan Preprint. In addition, OCC is requesting comments on the proposed CCDF Plan Preprint for FFY 2025–2027. Updates were made to clarify questions, enhance the ability to align data with OCC monitoring data, reflect equity and other OCC priorities, ensure alignment with federal requirements, and facilitate grantee submission in the Child Care Automated Reporting System (CARS) data system.

Respondents: State and Territory Lead Agencies.

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
Child Care and Development Fund for States and Territories (ACF–118)	56	0.33	200	3,696	1,232

Estimated Total Annual Burden Hours: 1,232; however, since Plans are required triennially, and remain in effect for 3 years, the actual *Total* Burden Hours is 3,696.

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: Pub. L. 113–186 and 42 U.S.C. 9858.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2023–13676 Filed 6–27–23; 8:45 am]

BILLING CODE 4184–87–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Judicial, Court, and Attorney Measures of Performance: Feedback and Implementation (New Collection)

AGENCY: Children's Bureau, Administration for Children and Families, United States Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Children's Bureau, Administration for Children and

Families (ACF), United States Department of Health and Human Services, is proposing to collect data for a new descriptive study, Judicial, Court, and Attorney Measures of Performance (JCAMP): Feedback and Implementation. This expands on earlier work around technical assistance, as approved under Office of Management and Budget #: 0970–0593.

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: This study will expand on a collection from field testing sites that informed the development of a suite of measures and tools, which became the JCAMP (OMB #0970–0970–0593 [1]). The data collection proposed here will further those efforts now that that suite of documents has been released. Specifically, this effort will (1) collect information from JCAMP implementation teams to understand their experiences with JCAMP implementation support, and (2) collect information from parents and children with child welfare cases, foster/kinship caregivers, judges, case workers, parent attorneys, children's attorneys, and child welfare agency attorneys to gather information for JCAMP measures selected for use by jurisdictions (jurisdictions will collect only the data elements relevant to them). This will be accomplished using eleven instruments:

JCAMP Feedback Survey: Members of JCAMP implementation teams will answer questions about their experiences with JCAMP written materials, technical assistance, and the eJCAMP online platform.

Parent Experience Survey: A brief survey that collects data post-hearing about parent experiences in court including, strategies used by judges to engage families, satisfaction with their legal representation, and collects demographic information.

Parent Court Experience Question Bank: This question bank includes options for items to include on a survey of parents with child welfare cases. Sites will select items that align with their chosen JCAMP measures. It is expected that surveys created from this bank will include up to 30 questions.

Parent Focus Group Guide: This focus group guide includes questions for parents with child welfare cases about their experiences with the child welfare court process.

Youth Post-Hearing Short Survey: This brief survey asks youth about their experiences immediately following hearings and collects demographic information (for example to allow assessment of equity aspects of judicial and legal practice and differences among age groups).

Youth Experience Survey: This survey collects information from youth with child welfare cases about their experiences with the child welfare court process and collects demographic information (for example to allow assessment of equity aspects of judicial and legal practice and differences among age groups).

Youth Court Experience Question Bank: This question bank includes options for items to include on a survey of youth with child welfare cases. Sites will select items that align with their chosen JCAMP measures. It is expected that surveys created from this bank will include up to 30 questions.

Youth Focus Group Guide: This focus group guide includes questions for youth with child welfare cases about their experiences with the child welfare court process.

Caregiver Survey: This survey collects information from adults caring for children with child welfare cases about their experiences with the child welfare court process and demographic information.

Stakeholder Survey: This survey collects data regarding judges' and attorneys' experiences in court including, persons present at hearings, judicial engagement strategies used with parents, children, and caregivers, the practices of parent, child, and agency attorneys during hearings, typical timelines to permanency, and case processing activities.

Stakeholder Focus Group Guide: This focus group guide asks judges, parent attorneys, children's attorneys, and child welfare agency attorneys their perceptions of the child welfare court system, including how families are engaged, how families receive due process, the quality of legal representation, safety decision-making, and permanency decision-making.

Other than the JCAMP Feedback Survey, all other instruments will be used for jurisdiction program and practice improvements.

Respondents: Respondents consist of Court Improvement Program administrators and staff, parents, youth, adult caregivers, judges, case workers, parent attorneys, children's attorneys, and agency attorneys.

ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
JCAMP Feedback Survey	100	1	0.25	25
Parent Experience Survey	250	1	0.17	42.5
Parent Court Experience Question Bank	250	1	0.17	42.5
Parent Focus Group Guide	80	1	1	80
Youth Post-Hearing Survey Short	250	1	0.08	20
Youth Experience Survey	250	1	0.17	42.5
Youth Court Experience Question Bank	250	1	0.17	42.5
Youth Focus Group Guide	80	1	1	80
Caregiver Survey	250	1	0.08	20
Stakeholder Survey	1,500	1	0.17	255
Stakeholder Focus Group Guide	400	1	1	400

[1] *https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202203-0970-010.*

Estimated Total Annual Burden Hours: 1,050.

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: Section 5106, Public Law 111–320, the Child Abuse Prevention and Treatment Act Reauthorization Act of 2010, and titles IV–B and IV–E of the Social Security Act.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2023–13677 Filed 6–27–23; 8:45 am]

BILLING CODE 4184–29–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–N–0895]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Imports and Electronic Import Entries

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments (including recommendations) on the collection of information by July 28, 2023.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function. The OMB control number for this information collection is 0910–0046. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Imports and Electronic Import Entries

OMB Control Number 0910–0046—Revision

This information collection supports Agency regulations found in 21 CFR part 1, subparts D (21 CFR 1.70 through 1.81) and E (21 CFR 1.83 through 1.101), governing FDA import activities and related Agency guidance. Specifically, the regulations prescribe the required data elements that respondents must submit when importing, or offering for import, an FDA-regulated article into the United States. Review of the data elements allows FDA to continue to meet its responsibilities pertaining to current submission requirements established by the U.S. Customs and Border Protection (CBP) related to the submission of entry information in using its Automated Commercial Environment (ACE) system, or any CBP-authorized electronic data interchange system. The regulations were recently revised through rulemaking to include data elements associated with import entries for veterinary devices (RIN 0910–AH66).

Respondents (ACE filers) submit important and useful information about FDA-regulated products being imported or offered for import into the United States so that we may effectively and efficiently review products and determine their admissibility. In addition, and as set forth in the regulations, certain product types are subject to additional data elements (for example, 21 CFR 1.77 prescribes additional data elements for radiation-emitting products), as well as those data elements applicable to all products.

The information collection also includes our weekly entry filing program (WEF). More detailed information on Foreign Trade Zones (FTZ)/WEF, is available at *https://www.fda.gov/industry/import-basics/foreign-trade-zonesweekly-entry-filing.* The WEF program allows entry filers to file a single entry estimating the amount of merchandise anticipated to be removed from an FTZ and offered for U.S. consumption during a 7-day period. To participate, we recommend respondents who wish to file a weekly entry of FDA-regulated products with CBP to first request a preliminary assessment from FDA. As part of the assessment, we also recommend submitting specific data elements, as discussed in the assessment. The information helps us appropriately route submissions within the Agency. Information on whether a product is stored or manufactured in the zone is necessary for FDA to determine the applicable admissibility requirements. The FTZ and port information is necessary to ensure that basic requirements in 19 CFR part 146 are met. The importer of record (IOR) and manufacturer FDA establishment identification number information is requested by FDA to expedite the admissibility review. Requests to participate in the WEF process are submitted to the FDA Import Division Office covering the intended port of entry.

The information collection also includes our Import Trade Auxiliary Communication System (ITACS). ITACS is used by the import trade community and was implemented to improve communication with FDA. By utilizing ITACS, respondents to the information collection have the ability to establish an account and electronically check the status of FDA-regulated entries and lines, submit entry documentation, submit the location of goods availability for those lines targeted for examination by FDA, and check the estimated laboratory analysis completion dates for lines that have been sampled. For further information regarding ITACS, please visit our website at *https://www.fda.gov/industry/import-systems/itacs.*

The information collection also includes burden associated with the use of Form FDA 766 entitled ''Application for Authorization to Relabel or Recondition Non-compliant Articles'' as the collection instrument for 21 CFR 1.95. Form FDA 766 facilitates collection of information associated with certain general enforcement provisions for importing FDA-regulated articles into the United States. The form is available at *https://www.fda.gov/industry/actions-enforcement/reconditioning.*

Relatedly, we are revising the information collection to include burden associated with the use of proposed electronic Form FDA 5054