option to retain historic location data if it has obtained affirmative express consent or it ensures that the historic location data is deidentified or rendered non-sensitive. Provision XIII requires Respondent to establish and implement, and thereafter maintain, a comprehensive privacy program that protects the privacy of consumers' personal information.

Provisions XIV–XVII are reporting and compliance provisions, which include recordkeeping requirements and provisions requiring Respondent to provide information or documents necessary for the Commission to monitor compliance.

Provision XVIII states that the Proposed Order will remain in effect for 20 years, with certain exceptions.

The purpose of this analysis is to facilitate public comment on the Proposed Order, and it is not intended to constitute an official interpretation of the complaint or Proposed Order, or to modify the Proposed Order's terms in any way.

By direction of the Commission.

April J. Tabor,

Secretary.

[FR Doc. 2024–01269 Filed 1–22–24; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Injury Prevention and Control; Amended Notice of Meeting

AGENCY: Centers for Disease Control and Prevention, Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: In accordance with the Federal Advisory Committee Act, the Centers for Disease Control and Prevention (CDC) announces an amendment to the following meeting for the Board of Scientific Counselors, National Center for Injury Prevention and Control (BSC, NCIPC).

SUPPLEMENTARY INFORMATION: Notice is hereby given of a change in the meeting of Board of Scientific Counselors, National Center for Injury Prevention and Control; January 11, 2024, first session from 10 a.m. to 12:05 p.m., EST (OPEN), and second session from 1 p.m. to 4:30 p.m., EST (CLOSED), in the original **Federal Register** notice.

The notice of the virtual meeting was published in the **Federal Register** on November 17, 2023, 88 FR 80305.

The meeting notice is being amended. The closed session that was scheduled for January 11, 2024, from 1 p.m. to 4:30 p.m., EST, has been canceled. The notice is being amended to update the **SUMMARY**, **DATES**, **ADDRESSES**, and **SUPPLEMENTARY INFORMATION** and should read as follows:

SUMMARY: In accordance with the Federal Advisory Committee Act, the Centers for Disease Control and Prevention (CDC) announces the following meeting for the Board of Scientific Counselors, National Center for Injury Prevention and Control (BSC, NCIPC). This meeting is open to the public. Time will be available for public comment.

DATES: The meeting will be held on January 11, 2024, from 10 a.m. to 12:05 p.m., EST. The public comment period will be from 11:45 a.m. to 12 p.m., EST.

ADDRESSES: Webinar, Atlanta, Georgia. All participants must register by using the following link to attend the meeting: *https://cdc.zoomgov.com/meeting/register/vJItf-igpjopGsXuGUhsdlIOmRCB2yx509k.*

SUPPLEMENTARY INFORMATION:

Purpose: The Board of Scientific Counselors, National Center for Injury Prevention and Control (BSC, NCIPC) will: (1) conduct, encourage, cooperate with, and assist other appropriate public health authorities, scientific institutions, and scientists in the conduct of research, investigations, experiments, demonstrations, and studies relating to the causes and strategies related to the prevention of injury, overdose, and violence; (2) assist States and other entities in preventing intentional and unintentional injuries, and to promote health and well-being; and (3) make recommendations of grants and cooperative agreements for research and prevention activities related to injury, overdose, and violence. The BSC, NCIPC makes recommendations regarding policies, strategies, objectives, and priorities and reviews progress toward injury, overdose, and violence prevention. The Board also provides advice on the appropriate balance of intramural and extramural research and provides guidance on the needs, structure, progress, and performance of intramural programs. Further, the Board provides guidance on extramural scientific program matters. Additionally, the Board provides second-level scientific and programmatic review of applications for research grants, cooperative agreements, and training grants related to injury, overdose, and violence prevention, and recommends approval of projects that merit further consideration for funding support. The Board also provides feedback and input on strategic plans, resources, and priority publications related to injury, overdose, and violence prevention.

Matters To Be Considered: The meeting will include a discussion on the updated Intimate Partner Violence Research Priorities. Agenda items are subject to change as priorities dictate.

FOR FURTHER INFORMATION CONTACT: Christopher R. Harper, Ph.D., Designated Federal Officer, Board of Scientific Counselors, National Center for Injury Prevention and Control, Centers for Disease Control and Prevention, 4770 Buford Highway NE, Mailstop S–1069, Atlanta, Georgia 30341. Telephone: (404) 718–8330; Email: *ncipcbsc@cdc.gov.*

The Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2024–01165 Filed 1–22–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-24–1071; Docket No. CDC–2024–0002]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a continuing information collection, as required by the Paperwork Reduction

Act of 1995. This notice invites comment on a proposed information collection project titled Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery. This data collection is designed to help CDC collect routine customer feedback on agency service delivery.

DATES: CDC must receive written comments on or before March 25, 2024.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0002 by either of the following methods:

- *Federal eRulemaking Portal: www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *www.regulations.gov.*

Please note: Submit all comments through the Federal eRulemaking portal (*www.regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329; Telephone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
3. Enhance the quality, utility, and clarity of the information to be collected;
4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses; and
5. Assess information collection costs.

Proposed Project

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (OMB Control No. 0920–1071, Exp. 5/31/2024)—Extension—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC/NCEZID is seeking a three-year Extension of OMB Control No. 0920–1071 to continue collecting routine customer feedback on agency service delivery. Executive Order 12862 directs Federal agencies to provide service to the public that matches or exceeds the best service available in the private sector. In order to work continuously to ensure that our programs are effective and meet our customers' needs, the National Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention (CDC) (hereafter the Agency) seeks to obtain OMB approval of a Generic Clearance to collect qualitative feedback on our service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study.

This collection of information is necessary to enable the Agency to garner customer and stakeholder feedback in an efficient, timely manner, in accordance with our commitment to improving service delivery. The information collected from our customers and stakeholders will help ensure that users have an effective, efficient, and satisfying experience with the Agency's programs. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

Since getting approval in May 2021, NCEZID has utilized the 0920–1071 mechanism 15 separate times. In this Extension, CDC requests OMB approval for an estimated 3,850 annual burden hours. There is no cost to respondents other than the time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
General public	Online surveys	1,500	1	30/60	750
	Focus groups	800	1	2	1,600
	In-person surveys	1,000	1	30/60	500
	Usability testing	1,500	1	30/60	750
	Customer comment cards	1,000	1	15/60	250
Total					3,850

Jeffrey M. Zirger,
Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2024–01152 Filed 1–22–24; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Generic Clearance for Financial Reports Used for ACF Mandatory Grant Programs (Office of Management and Budget #: 0970–0510)

AGENCY: 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) proposes to extend approval of the existing overarching generic clearance for Financial Reports used for ACF Mandatory Grant Programs (OMB #0970–0510) as well as all information collections currently approved under the overarching generic. There are no changes to the proposed types of information collection or uses of data as described in the overarching generic, and there are no changes proposed to currently approved information collections for which we are requesting an extension. Burden estimates for the next 3 years have been adjusted based on use to date.

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: ACF programs require detailed financial information from their grantees that allows ACF to monitor various specialized cost categories within each program, to closely manage program activities, and to have sufficient financial information to enable periodic thorough and detailed audits.

The information included on the standard Federal Financial Report Form (SF–425; OMB #4040–0014) provides only minimal, bare-bones, non-program specific financial information insufficient for these purposes. This generic clearance allows ACF programs to efficiently develop and receive approval for financial reports that are tailored to specific funding recipients and the associated needs of the program. This umbrella generic is a mechanism that is available to all ACF mandatory grant programs to use to obtain OMB approval of financial forms. Currently only a small number of ACF's mandatory grant program financial forms are covered under this umbrella; it does *not* cover all ACF mandatory grant program financial forms. Program offices use the information collected under this generic information collection to:

- Monitor program operations and prepare technical assistance and guidance as needed
- Assess the effect of program changes and make informed decision
- Assist in the computation of the grant awards issued to each program's grantees
- Assist in the computation of the Child Support Services program's annual incentive payments
- Determine that child support collections are being properly distributed (Child Support Services Program only)
- Ensure funding recipients are meeting funding requirements established by Congress
- Produce annual financial and statistical reports as may be required by Congress and respond to periodic detailed inquiries from Congress

ACF may require an information collection approved under this generic from funding recipients to obtain or retain benefits.

Prior to a new form being submitted for review under this umbrella generic, ACF will publish a notice in the **Federal Register** announcing the agency's intention to request an OMB review of the form and providing a 14-day period for public comment on that specific request. ACF will review any comments received and address them as appropriate. ACF will provide a copy of any comments received and will provide a description of how comments were considered in the submission form along with the request package for the individual collection. ACF will then follow standard OMB requirements for a generic information collection and submit a generic information collection request for each individual data collection activity under this generic clearance. Each request will include the individual form(s) and instructions, a summary of any comments received, and a short overview of the proposed purpose and use of the data collected. OMB should review requests within 10 days of submission.

Respondents: ACF-funded mandatory grant programs.

Annual Burden Estimates

Find currently approved information collections here: *https://www.reginfo.gov/public/do/PRAICList?ref_nbr=202308-0970-008.* The request to OMB will include an extension request for approved information collections that are planned to continue beyond spring 2024. The current list of ongoing collections follows, but more collections may be approved prior to submission of the extension request to OMB. We will update the list in the subsequent **Federal Register** notice, if needed.

Burden Estimates—Ongoing Requests

Study	Number of respondents	Number of responses per respondent	Average burden hours per response	Burden hours
ACF–196P, TANF Pandemic Emergency Assistance Fund (PEAF) Financial Report for States, Territories and Tribes	137	1	6	822
Child Care and Development Fund (CCDF) ACF–696 Financial Report for States and Territories	56	4	5	1,120
Child Care and Development Fund (CCDF) ACF–696T Financial Report for Tribal Grantees	221	1	7	1,547
Child Support Services Program Financial Reporting Forms (OCSE–34 and OCSE–396)	168	4	14	9,408
Form CB–496: Title IV–E Programs Quarterly Financial Report	67	4	25	6,700