momentum depends on the continuity of the same team managing the registry.

Summary of the award:

Recipient: Michigan State University.

Purpose of the Award: The purpose of this award is to support activities related to the Flint Registry, a comprehensive public health registry of residents who were exposed to lead-contaminated water from the Flint water system during April 25, 2014, to October 15, 2015.

Amount of Award: $3.6 million in Federal Fiscal Year (FYY) 2022 funds, with a total estimated $18 million over the 5-year period of performance, subject to availability of funds.

Authority: This program is authorized under Section 2203(b) of Public Law 114–322, the Water Infrastructure Improvements for the Nation (WIIN) Act of 2016; 42 U.S.C. Section 300j–27(b).

Period of Performance: August 1, 2022, through July 27, 2027.

Dated: January 25, 2022.

Terrance Perry,

Chief Grants Management Officer, Centers for Disease Control and Prevention.

[FR Doc. 2022–01800 Filed 1–27–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2022–0014]

Notice of Intent To Prepare a Supplemental Environmental Impact Statement and Request for Comments

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

ACTION: Notice and request for comment.

SUMMARY: Pursuant to the requirements of the National Environmental Policy Act of 1969 (NEPA) as implemented by the Council on Environmental Quality (CEQ) regulations (Title 40 CFR Section 1507.3) and the Department of Health and Human Services (HHS) General Administration Manual Part 30 ''Environmental Procedures,'' dated February 25, 2000, the Centers for Disease Control and Prevention (CDC) within HHS announces its intent to prepare a Supplemental Environmental Impact Statement (SEIS) to address changes proposed since completing the 2014 *Final Environmental Impact Statement (EIS) for Centers for Disease Control and Prevention Roybal Campus 2025 Master Plan* (2014 Final EIS) and issuing the Record of Decision dated November 7, 2014. The 2014 Final EIS analyzed the potential impacts associated with implementing a new long-range Master Plan to guide the future physical development of the Roybal Campus for the planning horizon of 2015 to 2025.

DATES: Written scoping comments will be accepted through February 28, 2022.

ADDRESSES: You may submit comments, identified by Docket Number CDC–2022–0014, by either of the following methods:

- *Federal eRulemaking Portal: https://www.regulations.gov/.* Follow the instructions for submitting comments.
- *U.S. Mail:* Thayra Riley, NEPA Coordinator Office of Safety, Security, and Asset Management (OSSAM), Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H20–4, Atlanta, Georgia 30329.

Instructions: All submissions received must include the Agency name and Docket Number (CDC–2022–0014). CDC will post, without change, all relevant comments to *https://www.regulations.gov,* including any personal information provided. Do not submit comments by email. CDC does not accept comments by email. For access to the docket to read background documents or comments received, go to *https://www.regulations.gov.*

FOR FURTHER INFORMATION CONTACT: Thayra Riley, NEPA Coordinator, Office of Safety, Security, and Asset Management (OSSAM), Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H20–4, Atlanta, Georgia 30329, email: *cdc-roybalga-seis@cdc.gov,* or telephone: 770–488–8170.

SUPPLEMENTARY INFORMATION: CDC intends to prepare an SEIS to analyze the potential impacts of proposed components that were not analyzed in the 2014 Final EIS. Proposed components that will be analyzed in the SEIS include the addition of a Hospital, Medical, and Infectious Waste Incinerator in a new laboratory and two emergency standby power diesel generators. The construction of a new laboratory was included in the 2014 Final EIS and will not be re-evaluated in the SEIS.

In accordance with NEPA as implemented by CEQ regulations (Title 40 CFR Section 1507.3) and HHS environmental procedures, CDC will prepare an SEIS to analyze the effects of proposed components that were not analyzed in the 2014 Final EIS. The potential impacts of construction and operation of these components on the natural and built environment will be evaluated.

Under the NEPA, federal agencies are required to evaluate the environmental effects of their proposed actions and a range of feasible alternatives to the proposed actions prior to making a final decision about what actions to take. The SEIS will incorporate the 2014 Final EIS by reference and will build upon that document to focus on specific resource areas that would have potential effects different from those analyzed in the 2014 Final EIS. Areas of anticipated concern include, but are not limited to, the following: Air quality, climate change and sustainability, environmental justice, and hazardous materials and hazardous waste.

Scoping Process

The scoping process is a requirement of NEPA and serves to identify the full range of environmental issues and inform the interested or affected parties of the proposed action. During the scoping process, CDC will actively seek input from the public, interested persons, organizations, and federal, state, and regional agencies to identify environmental concerns to be addressed in the SEIS. The purpose of this Notice is to inform interested or affected parties of CDC's plan to prepare the SEIS for component changes to a new laboratory that were not analyzed in the 2014 Final EIS, to provide information on the nature of the proposed actions, and to initiate the scoping process. NEPA does not require a public scoping meeting for an SEIS, and CDC will not conduct a public scoping meeting. CDC will conduct a virtual public meeting during the public comment period for the draft SEIS (a separate notice will be published for that comment period).

Dated: January 25, 2022.

Angela K. Oliver,

Executive Secretary, Centers for Disease Control and Prevention.

[FR Doc. 2022–01790 Filed 1–27–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB #0970–0531]

Proposed Information Collection Activity; Formative Data Collections for ACF Program Support

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF) plans to submit a request to the Office of Management and Budget (OMB) to extend approval of the existing overarching generic clearance for Formative Data Collections for ACF Program Support (OMB #0970–0531; expiration date 7/31/2022). ACF proposes minor updates to the description of potential generic information collections under the overarching generic and to the estimated number of respondents.

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 *OPREinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The goals of the generic information collections under this approval are to obtain information about program and grantee processes or needs and to inform the following types of activities, among others:

- Delivery of targeted assistance and/or workflows related to program and grantee processes. This could include the development and refinement of recordkeeping and communication systems.
- Planning for provision of programmatic or evaluation-related training or technical assistance (T/TA).
- Obtaining input on the development of program performance measures from grantees or others with experience or vested interest.
- Obtaining feedback about processes and/or practices to inform ACF program development or support, or ACF research.
- Use of rapid-cycle testing activities to strengthen programs in preparation for summative evaluations.

ACF uses a variety of techniques such as semi-structured discussions, focus groups, surveys, templates, open-ended requests, and telephone or in-person interviews in order to reach these goals.

Information collected under this overarching generic is meant to inform ACF activities and may be incorporated into documents or presentations that are made public such as through conference presentations, websites, or social media. The following are some examples of ways in which we may share information resulting from these data collections: Technical assistance plans, presentations, infographics, project specific reports, or other documents relevant to those involved with or interested in ACF programs such as federal leadership and staff, grantees, local implementing agencies, and/or T/TA providers.

Following standard OMB requirements, the Office of Planning, Research, and Evaluation will submit a change request for each individual data collection activity under this generic clearance. Each request will include the individual instrument(s), a justification specific to the individual information collection, and any supplementary documents. OMB should review requests within 10 days of submission.

The proposed types and the purpose of generic information collections submitted under this umbrella generic remain the same. Minor revisions are based on experiences over the past 3 years. These include:

- Updated burden estimates
- Broadened the description to make clearer the intention to broadly include respondents with knowledge, experience, or interest in ACF programs to allow ACF to learn about needs and processes related to ACF programs from those not necessarily funded by ACF

Respondents: Example respondents include current or prospective service providers, training or T/TA providers, grantees, contractors, current and potential participants in ACF programs or similar comparison groups, experts in fields pertaining to ACF programs, key groups involved in ACF projects and programs, individuals engaged in program re-design or demonstration development for evaluation, state or local government officials, or others involved in or prospectively involved in ACF programs.

BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total burden (in hours)
Semi-Structured Discussions and Focus Groups	10,000	1	2	20,000
Interviews	4,500	1	1	4,500
Questionnaires/Surveys	8,000	1.5	.5	6,000
Templates and Open-ended Requests	1,000	1	10	10,000

Estimated Total Annual Burden Hours: 40,500.

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: Social Security Act, Sec 1110 [42 U.S.C. 1310].

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–01777 Filed 1–27–22; 8:45 am]

BILLING CODE 4184–79–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Privacy Act of 1974; System of Records

AGENCY: Administration for Children and Families (ACF), Department of Health and Human Services (HHS).

ACTION: Notice of a modified system of records.