URM STATE/REPLACEMENT DESIGNEE AGENCIES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
ORR–3 Report	16	117	.2	374
ORR–4 Report	16	96	.25	384
Estimated Total Annual Burden Hours				758

URM PROVIDER AGENCIES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
ORR–3 Report	24	78	.4	749
ORR–4 Report	24	64	.6	922
Estimated Total Annual Burden Hours				1,670

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: 8 U.S.C. 1822(d).

Mary C. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2026–15843 Filed 8–4–26; 8:45 am]

BILLING CODE 4184–89–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[Office of Management and Budget #: 0970–0356]

Proposed Information Collection Activity; Formative Data Collections for ACF Research and Evaluation

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 data collection under the existing overarching generic clearance for Formative Data Collections for ACF Research and Evaluation (Office of Management and Budget (OMB)#: 0970–0356). There are no changes proposed to the intended purpose or use of the potential data collections, but burden has been adjusted to reflect agency priorities.

DATES: Comments due October 5, 2026.

ADDRESSES: In compliance with the requirements of the Paperwork Reduction Act (PRA) of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above. 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 promote the economic and social well-being of families, children, individuals, and communities. ACF conducts research on the programs, and with the populations they serve, through rigorous research and evaluation projects. These include evaluations of existing programs, evaluations of innovative approaches to helping low-income children and families, research syntheses, and descriptive and exploratory studies. ACF's research offers further understanding of current programs and service populations, explores options for program improvement, and assesses alternative policy and program designs. ACF regularly undertakes a variety of new research projects related to welfare, employment and self-sufficiency, Head Start, child care, healthy marriage and responsible fatherhood, family and youth services, home visiting, child welfare, trafficking, community services, and other areas of interest to ACF.

Under this generic clearance, ACF engages in a variety of formative data collections with researchers, practitioners, technical assistance providers, service providers, and potential participants throughout the field to fulfill the following goals: (1) inform the development of ACF research, (2) maintain a research agenda that is rigorous and relevant, (3) ensure that research products are as current as possible, and (4) inform the provision of technical assistance and supports around research and evaluation. ACF envisions using a variety of techniques including semi-structured discussions, focus groups, surveys, and telephone or in-person interviews, in order to reach these goals.

Information collected under this overarching generic is meant to inform ACF research activities and may be incorporated into documents or presentations that are made public. The following are some examples of ways in which we may share information resulting from these data collections: research design documents or reports; research or technical assistance plans; background materials for technical workgroups; concept maps, process maps, or conceptual frameworks; contextualization of research findings from a follow-up data collection that has full PRA approval; informational reports to technical assistance providers; or project specific reports, or other documents relevant to the field, such as federal leadership and staff, grantees, local implementing agencies.

Following standard OMB requirements, ACF has and will continue to submit to OMB information about individual information collection activities proposed under the generic

clearance. ACF will provide OMB with a copy of the individual instruments or questionnaires, as well as other materials describing the project. ACF requests OMB's review within 10 days of submission of individual requests under this generic.

Find currently approved information collections here: *https://www.reginfo.gov/public/do/PRAICList?ref_nbr=202504-0970-028.*

Respondents: Respondents could include key groups involved in ACF projects and programs, state or local government officials, service providers, participants in ACF programs or similar comparison groups, experts in fields pertaining to ACF research and programs, or others involved in conducting ACF research or evaluation projects.

Annual Burden Estimates

Burden Estimates—Ongoing Requests

The request to OMB will include an extension request for approved information collections that are planned to continue beyond the submission of this request to OMB (estimated November 2026). At this time, the following are expected to be included for an extension. This will be updated, if needed, for the 30-day comment period to reflect new GenICs or changes in plans for the following information collections.

Study	Responses	Burden hours
Center for Indigenous Research Collaborations and Learning in Home Visiting (CIRCLE–HV)	285	162
National and State Survey of Child and Adolescent Well-Being (NSSCAW): Informing Site Selection and Recruitment Processes	78	97
Title IV–E Prevention Services Clearinghouse	71	88
Totals	434	347

Total Burden Estimates—New Requests

Burden for potential new requests has been adjusted to reflect agency priorities, resulting in a 30 percent decrease compared to the currently approved burden for new requests.

	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours
Semi-Structured Discussions and Focus Groups	2,800	1	1.5	4,200
Interviews	1,050	1	1	1,050
Questionnaires/Surveys	800	1	.5	400
Totals	4,650	.....	.....	5,650

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, section 1110. [42 U.S.C. 1310].

Mary C. Jones,

ACF/OPRE Certifying Officer.

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BILLING CODE 4184–79–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2026–N–7605]

Establishment of a Quantitative Medicine Innovation Network: Scope, Feasibility, and Opportunities; Request for Information

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for information.

SUMMARY: The Center for Drug Evaluation and Research (CDER) within the Food and Drug Administration (FDA or Agency), through its Quantitative Medicine Center of Excellence (QM CoE), is announcing a request for information to inform the development of a Quantitative Medicine Innovation Network (QMIN). The vision for the QMIN is to create a cross-sector collaborative platform that accelerates the translation of quantitative medicine approaches to transform drug development, regulatory science, and clinical decision-making for patient benefit. The purpose of this request is to obtain feedback from industry, academia, healthcare providers, patient groups, and other interested parties on identifying critical community needs and establishing mechanisms for continuous engagement, supporting collaborative demonstration projects or research on innovative approaches aimed at addressing high priority areas through the application of QM approaches, and exploring mechanisms to leverage community expertise and capacity to promote QM innovation and adoption. Information submitted in response to this request will be considered in shaping the structure, priorities, and activities of the QMIN.

DATES: Either electronic or written comments on the notice must be submitted by November 3, 2026.

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 November 3, 2026. Comments received