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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[Office of Management and Budget #: 0970–0599]

Submission for Office of Management and Budget Review; Office of Refugee Resettlement Services for Survivors of Torture Program Data Points and Performance Progress Report

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 Administration for Children and Families' (ACF) Office of Refugee Resettlement (ORR) intends to continue collecting demographic, programmatic, and outcome data on Services for Survivors of Torture (SOT) grant recipients and the clients they serve. ORR collects information from the grantee cohort under the Survivors of Torture Program Data Points (PDP) and Program Performance Progress Report (PPR) (Office of Management and Budget (OMB) #0970–0599; Expiration date: February 28, 2026) to learn more about the populations served; the types and effectiveness of services provided; methods, challenges, and facilitators of implementing services; and grant recipients' progress towards programmatic goals. Revisions are proposed as described in the discussion section that follows.

DATES: *Comments due* March 26, 2026.

ADDRESSES: The public may view and comment on this information collection request at: *https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202602-0970-009.* 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: ORR proposes to continue to use the PDP Form and PPR, with revisions, to collect data on the Services for SOT grant recipients and their clients.

The recipients will continue to report their PDP through the ORR Refugee Arrivals Data System (RADS), an information technology platform used for enhanced data collection and record keeping.

Grant recipients will provide aggregated data on new and continuing clients annually, including demographic information, characteristics related to experiences of torture, services received, length of service, and wellbeing across six outcome domains.

Grant recipients will also provide information about community attendance at trainings and pro-bono services donated to the program. In the PPR, grant recipients will provide program narrative and program metric information on grant-funded activities and progress towards grant goals semi-annually.

Information collected will be used in aggregate by ORR to provide reports to stakeholders, including a required Report to Congress, and responses to funding requests.

ORR has made changes to the data collection, which include removing a total of twelve subcategories for two program indicators and reducing the frequency of reporting percentage-based outcomes in the program metrics. ORR has also added one subcategory in one program indicator. Overall, these changes have reduced the estimated reporting burden by 30 percent.

Respondents: Services for SOT grant programs (this may include non-profit social service, health, and higher education organizations, states, municipalities, and for-profit organizations).

Annual Burden Estimates: Estimated annual burden has been updated to reflect a reduction in estimated time per response from an average of 6 hours per response to an average of 4 hours per response.

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
PDP Form	35	1	4.2	147
PPRs—Parts A and B	35	2	4.2	294
Total Annual Burden				441

Authority: Section 5(a) of the ''Torture Victims Relief Act of 1998,'' Public Law 105–320 (22 U.S.C. 2152 note) Assistance for Treatment of Torture Victims.

Mary C. Jones,

ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2026–N–0686]

Agency Information Collection Activities; Proposed Collection; Comment Request; Current Good Manufacturing Practice in Manufacturing, Packaging, Labeling, or Holding Operations for Dietary Supplements

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) 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 FDA's regulations regarding current good