

report on the existence of a system to collect all information needed to determine and redetermine eligibility for Medicaid and CHIP. The State Medicaid/CHIP agency will attest to using the PARIS system in determining eligibility in Medicaid or CHIP benefit programs. *Form Number:* CMS–R–74 (OMB control number: 0938–0467); *Frequency:* Occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 55; *Total Annual Responses:* 3,241; *Total Annual Hours:* 1,082. (For policy questions regarding this collection contact: Abby Kahn at 410–786–4321.)

William N. Parham, III,
Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families
 [OMB #: 0970–0145]

Submission for Office of Management and Budget Review; Temporary Assistance for Needy Families (TANF) Program State Plan Guidance

AGENCY: Office of Family Assistance, 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) is requesting a 3-year extension of the Temporary Assistance for Needy Families (TANF) Program State Plan Guidance (TANF Program State Plan; Office of Management and Budget #0970–0145, expiration October 31, 2025). There are no changes requested to this information collection.
DATES: *Comments due December 26, 2025.*

ADDRESSES: The public may view and comment on this information collection request at: https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202511-0970-008. 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: The TANF Program State Plan is a mandatory statement submitted to the Secretary of the U.S. Department of Health and Human Services by the state. 42 U.S.C. 602. It consists of an outline specifying how the state’s TANF program will be administered and operated and contains certain required certifications by the state’s Chief Executive Officer. 42 U.S.C.

602(a). It is used to provide the public with information about the program. 42 U.S.C. 602(c). Authority to require states to submit a State TANF Plan is contained in section 402 of the Social Security Act (42 U.S.C. 602), as amended by the Personal Responsibility and Work Opportunity Reconciliation Act of 1996, Public Law 104–193, 110 Stat. 2105. States are required to submit new plans within a 27-month period. 42 U.S.C. 602(a).

Respondents: The 50 states of the U.S., the District of Columbia, Guam, Puerto Rico, and the U.S. Virgin Islands.

Annual Burden Estimates

The state TANF plan requirements for the 54 states (which includes three territories and the District of Columbia) will create a triennial burden with an average of 18 states responding annually. We estimate the annual burden to be an average of 30 hours per response. We also estimate that the triennial burden of plan amendments for the 54 states, with an average of 18 respondents annually spending approximately 3 hours per response.

Instrument	Total number of respondents per year	Total number of annual responses per respondent	Average burden hours per response	Annual burden hours
Title Amendments	18	1	3	54
State TANF plan	18	1	30	540
Estimated Total Annual Burden Hours				594

Authority: 42 U.S.C. 602.

Mary C. Jones,
ACF/OPRE Certifying Officer.
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration
 [Docket No. FDA–2025–D–5715]

Cross-Center Master Files: Where To Submit; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing the availability of a draft guidance for industry entitled “Cross-Center Master Files: Where to Submit.” The draft guidance provides recommendations to industry, specifically master file holders, regarding where (*i.e.*, to which FDA center) to submit a master file that is referenced in and intended to support more than one regulatory submission for which the lead center for those submissions may vary or where the information in the master file may need to be accessed and reviewed by more than one center to support review of the referencing submission(s). The

recommendations apply to master files submitted to the Center for Biologics Evaluation and Research (CBER), the Center for Drug Evaluation and Research (CDER), the Center for Devices and Radiological Health (CDRH), and certain types of master files submitted to the Center for Veterinary Medicine (CVM).

DATES: Submit either electronic or written comments on the draft guidance by February 24, 2026 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance at any time as follows: