ANNUAL BURDEN ESTIMATES—Continued

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Website Feedback Form	300	1	.083	24.90
Consultant Feedback Form	50	1	.083	4.15
Coordination Feedback Form	100	1	.050	5.00
Focus Group Demographic Survey	25	1	.033	.825
Focus Group Guide	25	1	.750	18.75
Follow-up Feedback Form	300	1	.133	39.90
General Training Feedback Form	150	1	.133	19.95
Interview Guide	25	1	.750	18.75
Pilot Feedback Form	25	1	.150	3.75
Requester Feedback Form	75	1	.117	8.78
Resource Tool Feedback Form	500	1	.033	16.50
SOAR Blended Learning Participant Feedback Form	30	1	.150	4.50
SOAR Conference Feedback Form	500	1	.200	100.00
SOAR Online Participant Feedback Form	1500	1	.100	150.00
SOAR Organizational Feedback Form	20	1	.133	2.66
SOAR Specialized T/TA Feedback Form	200	1	.150	30.00
Webinar Participant Feedback Form	1000	1	.067	67.00
Survivor Impact Consultant Application	20	1	.283	5.66
Expert T/TA Consultant Application	20	1	.267	5.34
Organizational Scholarship Application	10	1	.317	3.17
Professional Development Survivor Scholarship Application	30	1	.333	9.99
Total Annual Burden	5,908			689.15

Estimated Total Annual Burden Hours: 689 hours.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW, Washington, DC 20201. Attention Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2018–13998 Filed 6–28–18; 8:45 am]

BILLING CODE 4184–47–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity

AGENCY: Office of Planning, Research, and Evaluation; ACF; HHS.

ACTION: Request for Public Comment.

TITLE: Temporary Assistance for Needy Families (TANF) Data Innovations (TDI) Project (New Collection).

The Administration for Children and Families (ACF) within the U.S. Department of Health and Human Services (HHS) proposes to collect information as part of the TANF Data Innovations (TDI) project. TDI is an investment to expand the integration, analysis, and use of TANF data to improve program administration, payment integrity, and outcomes for participants.

TDI will start by assessing the needs and readiness of TANF agencies across the country to set up and operate data systems to support program improvement. A key goal of the needs assessment is to help categorize states' readiness to effectively use data and produce evidence. Informed by this assessment and discussions with key stakeholders, TDI will support a broad learning collaborative of state agencies and other entities related to the TANF program, including a range of Technical Assistance (TA) options to help states improve their use of TANF and other program data.

This information collection request will consist of a needs assessment survey to be completed by state TANF agency administrators and staff to gather detailed information about their capacities and needs. These data will help HHS to better understand the challenges and barriers states face in using data and research to inform program decision-making, and they will help the TDI team design future technical assistance activities for TANF agencies to address states' challenges.

Respondents: State TANF Administrators and TANF agency staff. We expect four respondents per state or territory.

ANNUAL BURDEN ESTIMATES

Instrument	Total/annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
Needs Assessment Survey	216	2	0.25	108

Estimated Total Annual Burden Hours: 108.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. Email address: *OPREinfocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION: 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: Sec. 413, Pub. L. 115–31.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2018–14037 Filed 6–28–18; 8:45 am]

BILLING CODE 4184–09–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–2029]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Administrative Practices and Procedures; Formal Evidentiary Public Hearing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by July 30, 2018.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, Fax: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0191. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Amber Sanford, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–8867, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Administrative Practices and Procedures; Formal Evidentiary Public Hearing

OMB Control Number 0910–0191—Extension

The Administrative Procedures Act (5 U.S.C. 553(e)) provides that every Agency shall give an interested person the right to petition for issuance, amendment, or repeal of a rule. Section 10.30 (21 CFR 10.30) sets forth the format and procedures by which an interested person may submit to FDA, in accordance with § 10.20 (21 CFR 10.20), a citizen petition requesting the Commissioner of Food and Drugs (the Commissioner) to issue, amend, or revoke a regulation or order, or to take or refrain from taking any other form of administrative action.

The Commissioner may grant or deny such a petition, in whole or in part, and may grant such other relief or take other action as the petition warrants. Respondents are individuals or households, State or local governments, and not-for-profit institutions or groups.

Section 10.33 (21 CFR 10.33), issued under section 701(a) of the Federal, Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 371(a)), sets forth the format and procedures by which an interested person may request reconsideration of part or all of a decision of the Commissioner on a petition submitted under 21 CFR 10.25 (Initiation of administrative proceedings). A petition for reconsideration must contain a full statement in a well-organized format of the factual and legal grounds upon which the petition relies. The grounds must demonstrate that relevant information and views contained in the administrative record were not previously or not adequately considered by the Commissioner. The respondent must submit a petition no later than 30 days after the decision involved. However, the Commissioner may, for good cause, permit a petition to be filed after 30 days. An interested person who wishes to rely on information or views not included in the administrative record shall submit them with a new petition to modify the decision. FDA uses the information provided in the request to determine whether to grant