for entry into the OSCAR system. The data is analyzed by the CMS regional offices and by the CMS central office components for program evaluation and monitoring purposes. The information is also available to the public upon request. *Form Number:* CMS–643 (OMB control number: 0938–0379); *Frequency:* Yearly; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 6,153; *Total Annual Responses:* 2,051; *Total Annual Hours:* 49,224. (For policy questions regarding this collection contact Thomas Pryor at 410–786–1132.)

Dated: September 20, 2022.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2022–20667 Filed 9–22–22; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Tribal Child Support Enforcement Direct Funding Request: 45 CFR 309 (OMB #0970–0218)

AGENCY: Office of Child Support Enforcement, Administration for Children and Families, Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Office of Child Support Enforcement (OCSE), Administration for Children and Families (ACF) is requesting a 3-year extension of the Tribal Child Support Enforcement Direct Funding Requests (Office of Management and Budget (OMB) #0970–0218, expiration March 31, 2023) with revisions. We are proposing to provide an optional Table of Contents and Cover Sheet for plan pages.

DATES: *Comments due within 60 days of publication.* 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.

ADDRESSES: You can obtain copies of the proposed collection of information and submit comments by emailing *ocse.tribal@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The final rule within 45 CFR part 309, published in the **Federal Register** on March 30, 2004, contains a regulatory reporting requirement that, in order to receive funding for a Tribal IV–D program, a tribe or tribal organization must submit a plan describing how the tribe or tribal organization meets or plans to meet the objectives of section 455(f) of the Social Security Act, including establishing paternity; establishing, modifying, and enforcing support orders; and locating noncustodial parents. The plan is required for all tribes requesting funding; however, once a tribe has met the requirements to operate a comprehensive program, a new plan is not required annually unless a tribe makes changes to its title IV–D program. If a tribe or tribal organization intends to make any substantial or material changes, a Tribal IV–D plan amendment must be submitted for approval. Tribes and tribal organizations must have an approved plan and submit any required plan amendments to receive funding to operate a Tribal IV–D program.With this request to extend approval of this information collection, OCSE is proposing a change to the paperwork by providing optional plan pages. The optional plan pages organize the Tribal IV–D plan, identify required attachments, and streamline plan amendment submissions. Tribes and tribal organizations who choose to participate will attest to complying with the regulatory requirements in 45 CFR, Parts 309 and 310 and submit plan amendments for changes to the required attachments identified in the Table of Contents. The optional plan pages organize the Tribal IV–D plan, identify required attachments, and streamline plan amendment submissions.

Respondents: Tribes and tribal Organizations.

Annual Burden Estimates

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
45 CFR 309–New Plan	2	1	480	960
45 CFR 309–Amended Plan	60	1	105	6300

Estimated Total Annual Burden Hours: 7,260.

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: Title IV–D of the Social Security Act; 45 CFR 309.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–20613 Filed 9–22–22; 8:45 am]

BILLING CODE 4184–41–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.652]

Announcement of the Intent To Award a Single-Source Grant to the Center for Adoption Support and Education in Burtonsville, MD

AGENCY: Children's Bureau (CB), Administration for Children, Youth and Families (ACYF), Administration for Children and Families (ACF), Department of Health and Human Services (HHS).

ACTION: Notice of Issuance of a Single-source cooperative agreement to the

Center for Adoption Support and Education (C.A.S.E) for the continued expansion and use of the National Adoption Competency Mental Health Training Initiative across the nation.

SUMMARY: The ACF, ACYF, Children's Bureau, Division of Capacity Building announces the intent to award a single-source cooperative agreement in the amount of up to $10,000,000 to the Center for Adoption Support and Education in Burtonsville, MD. The purpose of this award is to continue to scale the web-based National Training Initiative for Adoption Competent Mental Health Training program to remaining states in the nation, refine and update the curriculum as needed and perform an updated evaluation regarding current use of the curriculum.

DATES: The proposed period of performance is September 30, 2022 to September 29, 2027.

FOR FURTHER INFORMATION CONTACT: June Dorn, National Adoption Specialist, Division of Capacity Building, 333 C St. SW, Suite 3521B, Washington, DC 20201. Telephone: (202) 205–9450; Email: *June.Dorn@acf.hhs.gov.*

SUPPLEMENTARY INFORMATION: Award funds will support the continued expansion and use of the National Training Initiative on Mental Health Competency across the nation. This curriculum was developed through an initial 5-year grant to C.A.S.E. and pilot tested prior to being supplemented and extended for 3 additional years for the further dissemination and integration of this important web-based training in the state child welfare training systems so there would be consistent use of the knowledge imparted by it across all child welfare systems.

This funding will allow for the continued scale up of the web-based trainings for the child welfare workforce and mental health practitioners to remaining states in the nation; the refinement and update of the curricula as needed; and additional and updated evaluation of the curricula.

Statutory Authority: Title II, section 203 of the Child Abuse Prevention and Treatment and Adoption Reform Act of 1978 (42 U.S.C. 5113(b)), as amended by CAPTA Reauthorization Act of 2010.

Elizabeth A. Leo,

Senior Grants Policy Specialist, Office of Grants Policy, Office of Administration.

[FR Doc. 2022–20587 Filed 9–22–22; 8:45 am]

BILLING CODE 4184–44–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–N–1886]

Agency Information Collection Activities; Proposed Collection; Comment Request; Endorser Status and Actual Use in Direct-to-Consumer Television Ads

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 and to allow 60 days for public comment in response to the notice. This notice solicits comments on a proposed study entitled ''Endorser Status and Actual Use in Direct-to-Consumer Television Ads.''

DATES: Submit either electronic or written comments on the collection of information by November 22, 2022.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before November 22, 2022. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of November 22, 2022. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2022–N–1886 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Endorser Status and Actual Use in Direct-to-Consumer Television Ads.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this