(OMB) regulations, DoD, GSA, and NASA invite the public to comment on a revision and renewal concerning termination settlement proposal forms in the FAR. DoD, GSA, and NASA invite comments on: Whether the proposed collection of information is necessary for the proper performance of the functions of Federal Government acquisitions, including whether the information will have practical utility; the accuracy of the estimate of the burden of the proposed information collection; ways to enhance the quality, utility, and clarity of the information to be collected; and ways to minimize the burden of the information collection on respondents, including the use of automated collection techniques or other forms of information technology. OMB has approved this information collection for use through February 29, 2020. DoD, GSA, and NASA propose that OMB extend its approval for use for three additional years beyond the current expiration date.

DATES: DoD, GSA, and NASA will consider all comments received by January 27, 2020.

ADDRESSES: DoD, GSA, and NASA invite interested persons to submit comments on this collection by either of the following methods:

- *Federal eRulemaking Portal:* This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. Go to *http://www.regulations.gov* and follow the instructions on the site.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405. ATTN: Lois Mandell/IC 9000–0012, Termination Settlement Proposal Forms—FAR (SF 1435 through 1440).

Instructions: All items submitted must cite Information Collection 9000–0012, Termination Settlement Proposal Forms—FAR (SF 1435 through 1440). Comments received generally will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two-to-three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

FOR FURTHER INFORMATION CONTACT: Zenaida Delgado, Procurement Analyst, at telephone 202–969–7207, or *zenaida.delgado@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. OMB Control Number, Title, and Any Associated Form(s)

9000–0012, Termination Settlement Proposal Forms—FAR (SF 1435 through 1440).

B. Need and Uses

The termination settlement proposal forms (Standard Forms 1435 through 1440) provide a standardized format for listing essential cost and inventory information needed to support the terminated contractor's negotiation position per the Federal Acquisition Regulation subpart 49.6, Contract Termination Forms and Formats. Submission of the information assures that a contractor will be fairly reimbursed upon settlement of the terminated contract.

C. Annual Burden

Respondents: 4,995.
Total Annual Responses: 14,128.
Total Burden Hours: 33,907.
Obtaining Copies: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0012, Termination Settlement Proposal Forms—FAR (SF 1435 through 1440), in all correspondence.

Dated: November 20, 2019.

Janet Fry,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2019–25580 Filed 11–25–19; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Injury Prevention and Control (NCIPC); Correction

Notice is hereby given of a change in the meeting of the Board of Scientific Counselors, National Center for Injury Prevention and Control (NCIPC); December 4, 2019, 9:00 a.m. to 4:40 p.m., EST; and December 5, 2019, 9:00 a.m. to 11:30 a.m., EST which was published in the **Federal Register** on October 24, 2019, Volume 84, Number 206, page 57021.

The meeting location has been changed from the Hilton Garden Inn, 3342 Peachtree Road NE, Atlanta, Georgia 30326 to the Centers for Disease Control and Prevention, 4770 Buford Highway NE, Chamblee Campus, Building 106, Conference Room 1–B, Atlanta, Georgia 30341. This Federal facility meeting room accommodates 80 people.

For Further Information Contact: Gwendolyn H. Cattledge, Ph.D., M.S.E.H., Deputy Associate Director for Science, NCIPC, CDC, 4770 Buford Highway NE, Mailstop F–63, Atlanta, Georgia 30341; telephone (770) 488–3953; email address: *NCIPCBSC@cdc.gov.*

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2019–25612 Filed 11–25–19; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Child Care and Development Fund (CCDF) Tribal Annual Report—ACF–700 (0970–0430)

AGENCY: Office of Child Care, Administration for Children and Families, HHS.

ACTION: Request for Public Comment.

SUMMARY: The Administration for Children and Families (ACF) is requesting a three-year extension of the form ACF–700: Child Care and Development Fund (CCDF) Tribal Annual Report (OMB #0970–0430, expiration 11/30/2019) with changes.

DATES: *Comments due within 30 days of publication.* 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.

ADDRESSES: 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.

Copies of the proposed collection may be obtained by emailing *infocollection@acf.hhs.gov.* Alternatively, copies can also be obtained 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. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The Child Care and Development Fund (CCDF) Tribal Annual Report (ACF–700) requests Tribal Lead Agencies (TLAs) to provide annual Tribal aggregate information on services provided through the CCDF, which is required by CCDF regulations (45 FR parts 98 and 99). The revised ACF–700 report consists of an introductory section that provides program characteristics and two parts: (1) Administrative Data, and (2) Tribal Narrative. The content and format of the entire form have been revised to address Child Care and Development Block Grant (CCDBG) Act of 2014 changes and to reduce the reporting burden to TLAs.

Information from the ACF–700 will be included in the CCDF Report to Congress, as appropriate, and will be shared with TLAs to inform them of CCDF-funded activities.

Respondents: Tribal Governments.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
ACF–700	138 (Tribes with small allocation)	3	19	7,866	2,622
ACF–700	83 (Tribes with medium/large allocation).	3	26	6,474	2,158

Estimated Total Annual Burden Hours: 4,780.

Authority: 42 U.S.C. 9857.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2019–25607 Filed 11–25–19; 8:45 am]

BILLING CODE 4184–43–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2019–D–5324]

Compliance Policy for Limited Modifications to Certain Marketed Tobacco Products; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the availability of a final guidance for industry entitled ''Compliance Policy for Limited Modifications to Certain Marketed Tobacco Products.'' This guidance describes FDA's compliance policy for premarket review requirements for two types of limited modifications to new tobacco products that were on the market as of August 8, 2016, specifically, modifications to battery-operated tobacco products solely to comply with UL 8139 and modifications to liquid nicotine products solely to comply with the Child Nicotine Poisoning Prevention Act of 2015 (CNPPA) flow restrictor requirements for liquid nicotine containers. This guidance will enable tobacco manufacturers to upgrade their battery-operated tobacco products to UL 8139. It will also enable manufacturers to comply with the CNPPA requirements for flow restrictors for liquid nicotine containers. FDA is issuing this guidance to address battery safety concerns and youth exposure to liquid nicotine toxicity.

DATES: The announcement of the guidance is published in the **Federal Register** on November 26, 2019.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time as follows:

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–2019–D–5324 for ''Compliance Policy for Limited Modifications to Certain Marketed Tobacco Products.'' Received comments 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.

- 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