the Secretary on the types of problems and inquiries encountered by consumers'' (Sec. 2793 (d)). Analysis of this data reporting will help identify patterns of practice in the insurance marketplaces and uncover suspected patterns of noncompliance. HHS must share program data reports with the Departments of Labor and Treasury, and State regulators. Program data also can offer CCIIO one indication of the effectiveness of State enforcement, affording opportunities to provide technical assistance and support to State insurance regulators and, in extreme cases, inform the need to trigger federal enforcement. *Form Number:* CMS–10333 (OMB Control Number: 0938–1097); *Frequency:* Annually, Quarterly; *Affected Public:* Private Sector: State, Local, or Tribal Governments; *Number of Respondents:* 51; *Total Annual Responses:* 459; *Total Annual Hours:* 9,588. (For policy questions regarding this collection contact Lateefa Dawkins at 301–492–4262.)

Dated: October 28, 2015.

William N. Parham, III,

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

[FR Doc. 2015–27859 Filed 10–30–15; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Evaluation of the Child Welfare Capacity Building Collaborative.

OMB No.: New Collection.

Description: The Evaluation of the Child Welfare Capacity Building Collaborative is sponsored by the Children's Bureau, Administration for Children and Families of the U.S. Department of Health and Human Services. The Capacity Building Collaborative includes three centers (Center for States, Center for Tribes, Center for Courts) funded by the Children's Bureau to provide national child welfare expertise and evidence-informed training and technical assistance services to State, Tribal and Territorial public child welfare agencies and Court Improvement Programs (CIPs). The Centers offer a wide array of services including, but not limited to: Web-based content and resources, product development and dissemination, self-directed and group-based training, virtual learning and peer networking events, and tailored consultation and coaching. During the project period the Centers' services will be evaluated by both Center-specific evaluations and a Cross-Center Evaluation. The Center-specific evaluations are designed to collect data on Center-specific processes and outcomes. The Cross-Center Evaluation is designed to respond to a set of cross-cutting evaluation questions posed by the Children's Bureau. The Cross-Center Evaluation will examine: The extent to which key partners across and within the Centers are collaborating; whether the capacity building service interventions offered by the Centers are evaluable; the degree to which Centers follow common protocols; whether service interventions are delivered or performed as designed; how satisfied recipients are with the services received; how effective the service interventions were; which service approaches were most effective and under what conditions; and the costs of services.

The Cross-Center Evaluation is utilizing a longitudinal mixed methods approach to evaluate the Centers' services as they develop and mature over the course of the study period. Multiple data collection strategies will be used to efficiently capture quantitative and qualitative data to enable analyses that address each evaluation question. Proposed Cross-Center Evaluation data sources for this effort include (1) satisfaction surveys to assess recipients' satisfaction with services, such as the Learning Experiences Satisfaction Survey; (2) a leadership interview, administered to all State child welfare directors, Tribal child welfare directors, and CIP coordinators that are receiving services from the Centers; and (3) a collaboration survey, an annual Web-based survey administered to the directors and staff of the three Centers. Center-specific data sources for this effort include (1) assessment tools such as the Tribal Organizational Assessment Caseworker Interview; and (2) service-specific feedback forms, such as the Center for States Intensive Projects instrument and the Center for Courts CQI Workshops instrument.

Respondents: Respondents of data collection instruments will include (1) child welfare and judicial professionals that use the Centers' Web pages, products, and online courses, that participate in virtual or in-person trainings or peer events, and that receive brief or intensive tailored services from the Centers; (2) State child welfare directors, Tribal child welfare directors, and CIP coordinators that are receiving services from the Centers; and (3) the directors and staff of the three Capacity Building Centers. The proposed data collection will span four years.

ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Webpage & Product Satisfaction Survey	1,560	1	.08	125
Learning Experiences Satisfaction Survey	500	1	.33	165
Learning Experience Module Survey	900	1	.08	72
Peer Event Satisfaction Survey	5,502	1	.08	441
Assessment & Capacity Building Plan Satisfaction Survey	450	1	.066	30
Center for Tribes Contact Form	50	1	.05	3
Center for Tribes Demographic Survey	20	1	1.75	35
Tribal Organizational Assessment Caseworker Interview	20	1	1.25	25
Tribal Organizational Assessment Community Provider Interview	16	1	1.25	20
Tribal Organizational Assessment Community Member/Elder Interview	12	1	1.0	12
Tribal Organizational Assessment Family Interview	14	1	1.0	14
Center for States Information and Referral	12	1	.05	1
Center for States Intensive Projects	330	2	.33	218
Center for States Constituency Groups	400	2	.33	264
Center for States Consultant Feedback Form	156	1	.13	21
Center for States Brief Services	125	1	.33	42

ANNUAL BURDEN ESTIMATES—Continued

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
CIP Annual Meeting Survey	200	1	.13	26
Center for Courts CQI Workshops	48	1	.17	8
Leadership Interview—States	13	2	1	26
Leadership Interview—CIPs	13	2	1	26
Leadership Interview—Tribes	8	2	1.25	20
Leadership Interview Part II—Tribes	8	2	.67	11
Annual Collaboration Survey	230	1	.36	83

Estimated Total Annual Burden Hours: 1,688.

In compliance with the requirements of Section 506(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. 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, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

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.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2015–27833 Filed 10–30–15; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–D–2270]

The Drug Supply Chain Security Act Implementation: Product Tracing Requirements for Dispensers—Compliance Policy; Updated Guidance for Industry, Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability; revised guidance document.

SUMMARY: The Food and Drug Administration (FDA or we) is issuing a revised guidance document that extends the compliance policy described in the guidance for industry entitled ''DSCSA Implementation: Product Tracing Requirements for Dispensers—Compliance Policy.'' This revised guidance announces FDA's intention with regard to enforcement of certain product tracing requirements of the Federal Food, Drug, and Cosmetic Act (FD&C Act) added by the Drug Supply Chain Security Act (DSCSA). FDA does not intend to take action against dispensers who, prior to March 1, 2016, accept ownership of product without receiving transaction information, transaction history, and transaction statements (product tracing information), prior to or at the time of a transaction, or do not capture and maintain the product tracing information, as required by the FD&C Act.

DATES: Effective November 2, 2015. For information about enforcement dates, please see the **SUPPLEMENTARY INFORMATION** section.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *http://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 *http://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):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, 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–2015–D–2270 for ''The Drug Supply Chain Security Act Implementation: Product Tracing Requirements for Dispensers—Compliance Policy; Revised Guidance for Industry.'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *http://www.regulations.gov* or at the