DATES: The meeting will be held on November 30, 2017, 9:00 a.m. to 5:00 p.m., EST and December 1, 2017 9:00 a.m. to 1:00 p.m., EST.

ADDRESSES: CDC Chamblee Campus 4770 Buford Highway, Building 106, Room 1B, Atlanta, GA 30341. The web-conference access for November 30, 2017 is *https://ondieh.adobeconnect.com/bccedcac/.* The web-conference access for December 1, 2017 is *https://ondieh.adobeconnect.com/bccedcac2/.*

FOR FURTHER INFORMATION CONTACT: Jameka Reese Blackmon, MBA,CMP, Designated Federal Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 4770 Buford Highway NE., Mailstop F76, Atlanta, Georgia, 30341–3717, Telephone (770) 488–4740; *grz4@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose: The committee is charged with advising the Secretary, Department of Health and Human Services, and the Director, CDC, regarding the early detection and control of breast and cervical cancer. The committee makes recommendations regarding national program goals and objectives; implementation strategies; and program priorities including surveillance, epidemiologic investigations, education and training, information dissemination, professional interactions and collaborations, and policy.

Matters To Be Considered: The agenda will include discussions on expanded National Breast and Cervical Cancer Early Detection Program (NBCCEDP) strategies to increase breast and cervical cancer screening, review of success with implementing evidence-based interventions in health systems in the Colorectal Cancer Control Program and expanding evaluation to measure impact and sustainability of population based activities. Agenda items are subject to change as priorities dictate.

The Director, Management Analysis and Services Office, 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.

Claudette Grant,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2017–22059 Filed 10–11–17; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Information Comparison With Insurance Data.

OMB No.: 0970–0342.

Description: The Deficit Reduction Act of 2005 amended Section 452 of the Social Security Act (the Act) to authorize the Secretary, through the Federal Parent Locator Service (FPLS), to conduct comparisons of information concerning individuals owing past-due child support with information maintained by insurers (or their agents) concerning insurance claims, settlements, awards, and payments. 42 U.S.C. 652(m)(1).

An insurer may choose to participate in the data comparison using one of the following methods:

- An insurer submits information concerning claims, settlements, awards, and payments to the federal Office of Child Support and Enforcement (OCSE). OCSE compares the information with parents who owe past-due support.
- OCSE sends a file containing information about parents who owe past-due support to the insurer, or their agent to compare with their claims, settlements, awards, and payments. The insurer or their agent sends the matches to OCSE.

On a daily basis, OCSE sends the results of the comparison in the Insurance Match Response Record to child support agencies responsible for collecting past-due support. Child support agencies use the matches to collect past-due support from the insurance proceeds.

The information collection activities pertaining to the information comparison with insurance data are authorized by:

(1) 42 U.S.C. 652(a)(9) which requires the federal Office of Child Support Enforcement (OCSE) to operate the FPLS established by 42 U.S.C. 653(a)(1); and

(2) 42 U.S.C. 652(m) which authorizes OCSE, through the FPLS, to compare information concerning individuals owing past-due support with information maintained by insurers (or their agents) concerning insurance claims, settlements, awards, and payments, and to furnish information resulting from the data matches to the state child support agencies responsible for collecting child support from the individuals.

Respondents: Insurers or their agents, including the U.S. Department of Labor and state agencies administering workers' compensation programs, and the Insurance Services Office (ISO).

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Insurance Match File: Monthly Reporting Electronically	22	12	0.083	22
Insurance Match File: Weekly Reporting Electronically	7	52	0.083	30
Insurance Match File: Daily Reporting Electronically	2	251	0.083	42
Match File: Daily Reporting Manually	80	251	0.1	2,008
Total				2,102

Estimated Total Annual Burden Hours: 2,102 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.

Mary Jones,

Reports Clearance Officer.

[FR Doc. 2017–21960 Filed 10–11–17; 8:45 am]

BILLING CODE 4184–41–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–5568]

Prescription Drug User Fee Act of 2017; Electronic Submissions and Data Standards; Public Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the following public meeting entitled ''Prescription Drug User Fee Act of 2017; Electronic Submissions and Data Standards.'' The topics to be discussed will include the current status of electronic submissions and data standards initiatives to improve the predictability and consistency of the electronic submissions process in support of the human drug review program. FDA is seeking input from a variety of stakeholders—industry, academia, patient advocates, professional societies and other interested parties—as it fulfills its commitment under the Prescription Drug User Fee Act of 2017 (PDUFA) to hold annual public meetings to seek stakeholder input related to enhancing the transparency and accountability of the electronic submission and data standards activities. FDA will use the information from the public meeting to inform the development of the FDA Information Technology (FDA IT) Strategic Plan and electronic submissions gateway target timeframes.

DATES: The public meeting will be held March 21, 2018, from 9 a.m. to 4 p.m. Submit either electronic or written comments regarding this public meeting prior to the meeting through April 18, 2018. See the **SUPPLEMENTARY INFORMATION** section for registration date and information.

ADDRESSES: The public meeting will be held at the FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503, Section A), Silver Spring, MD 20993–0002. Entrance for the public meeting participants (non-FDA employees) is through Building 1, where routine security check procedures will be performed. For parking and security information, please refer to *https://www.fda.gov/AboutFDA/WorkingatFDA/BuildingsandFacilities/WhiteOakCampusInformation/default.htm.*

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 April 18, 2018. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of April 18, 2018. 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–2017–N–5568 for ''Prescription Drug User Fee Act of 2017; Electronic Submissions and Data Standards.'' 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.

- 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 information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Ron Fitzmartin, Center for Drug Evaluation