

business days in advance at https://www.cdc.gov/cancer/breast/what_cdc_is_doing/conference.htm. Please complete all the required fields before submitting your registration and submit no later than January 31, 2020.

DATES: The meeting will be held on February 6, 2020, 9:00 a.m. to 1:00 p.m., EST.

ADDRESSES: The meeting will be held by teleconference and web conference. The teleconference access is 1-888-606-5944; and the passcode is 8340472. The web conference access is <https://adobeconnect.cdc.gov/rwa641n3jrry/>.

FOR FURTHER INFORMATION CONTACT: Temeika L. Fairley, Ph.D., Designated Federal Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 5770 Buford Highway NE, Mailstop S107-4, Atlanta, Georgia 30341; Telephone (770) 488-4518; Fax: (770) 488-4760; Email: acbcyw@cdc.gov.

SUPPLEMENTARY INFORMATION:

Purpose: The committee provides advice and guidance to the Secretary, HHS; the Assistant Secretary for Health; and the Director, CDC, regarding the formative research, development, implementation and evaluation of evidence-based activities designed to prevent breast cancer (particularly among those at heightened risk) and promote the early detection and support of young women who develop the disease. The advice provided by the Committee will assist in ensuring scientific quality, timeliness, utility, and dissemination of credible appropriate messages and resource materials.

Matters To Be Considered: The agenda will include discussions on current topics related to breast cancer in young women. These will include Mental/Behavioral Health, Sexual Health, Genetics and Genomics, and Provider Engagement. Agenda items are subject to change as priorities dictate.

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. 2019-26919 Filed 12-12-19; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Generic Program-Specific Performance Progress Report (0970-0490)

AGENCY: Office of Planning, Research, and Evaluation; Administration for Children and Families; HHS.

ACTION: Request for public comment.

SUMMARY: This Notice describes the proposal to extend data collection under the Administration for Children and Families (ACF) Generic Program-Specific Performance Progress Report (PPR) (0970-0490). This overarching generic allows ACF program offices to collect performance and progress data from recipients and sub-recipients who receive funding from ACF under a discretionary grant or cooperative agreement. This information is required under 45 CFR 75.342, monitoring and reporting program performance. The generic program-specific PPR was originally approved in January 2017.

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 OPREinfocollection@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: ACF is primarily a grant-making agency that promotes the economic and social well-being of families, children, individuals and communities with partnerships, funding, guidance, training and technical assistance.

Prior to the use of this generic program-specific PPR, a standard ACF PPR (#0970-0406) was used for all ACF discretionary grant and cooperative agreement awards for post award reporting. Historically, on the standard ACF PPR form, ACF required grantees to only respond to a common set of broad questions, which often solicited qualitative or incomplete information. This one-size-fits-all approach did not adequately collect the specific data needed for particular grant programs or allow program offices to assess continuous quality improvement. Different grant programs vary in purpose, target population, and activities. Therefore, a need for program offices to customize performance measurements was identified and the generic program-specific PPR was developed.

ACF program offices have benefited from the ability to create and use a program-specific PPR that is more effective and includes specific data elements that reflects a specific program's indicators, demographics, priorities and objectives. This extension includes extension of previously approved program-specific PPRs under this OMB #. A generic program-specific PPR that can be tailored for program-specific needs allows program offices to collect useful data in a uniform and systematic manner. The reporting format allows program offices to gather uniform program performance data from each grantee, allowing aggregation at the program level to calculate outputs and outcomes, providing a snapshot and allowing for longitudinal analysis.

Data from a tailored program-specific PPR that demonstrates a program's successes and challenges have been useful for accountability purposes, such as required reports to Congress. Moreover, it has been useful for program management and oversight, such as identifying grantees' technical assistance needs and ensuring compliance with Federal and programmatic regulations and policies.

Respondents: ACF Grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hour per response	Total burden hours
New Program Specific PPRs	600	2	4	4,800

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2019–26913 Filed 12–12–19; 8:45 am]

BILLING CODE 4184–79–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Request for Information: Family Caregiving Advisory Council; Correction

AGENCY: Administration for Community Living, HHS.

ACTION: Notice of correction.

SUMMARY: The Administration for Community Living (ACL) published a document in the **Federal Register** on December 9, 2019, requesting information to the Advisory Council to Support Grandparents Raising Grandchildren seeking information to be used in the development of the Initial Report, as required by the Supporting Grandparents Raising Grandchildren Act (SGRG). The ACL wishes to change a line in the titling of the notice in order to avoid confusion for potential commenters.

SUPPLEMENTARY INFORMATION:

Correction: In the **Federal Register** of December 9, 2019, in FR doc. 2019–26437, on page 67270, in the second column, the second line should be changed to “Request for Information: Advisory Council to Support Grandparents Raising Grandchildren.” In addition, the **DATES** section due date is incorrect. It should read as follows: “**DATES:** Comments on the request for information must be submitted by 11:59 p.m. (EST) on February 7, 2019.”

FOR FURTHER INFORMATION CONTACT:
SGRG.Act@acl.hhs.gov

Dated: December 9, 2019.

Lance Robertson,

Administrator and Assistant Secretary for Aging.

[FR Doc. 2019–26880 Filed 12–12–19; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2019–D–4751]

Food and Drug Administration Reauthorization Act Implementation Guidance for Pediatric Studies of Molecularly Targeted Oncology Drugs; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a draft guidance for industry entitled “FDARA Implementation Guidance for Pediatric Studies of Molecularly Targeted Oncology Drugs.” This draft guidance addresses early planning for pediatric evaluation of certain molecularly targeted oncology drugs, including biological products, for which original new drug applications (NDAs) and biologics license applications (BLAs) are expected to be submitted to FDA on or after August 18, 2020, in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (FD&C Act) as amended by the FDA Reauthorization Act of 2017 (FDARA). This guidance addresses the implementation of amendments made by FDARA to the FD&C Act regarding molecularly targeted oncology drugs.

DATES: Submit either electronic or written comments on the draft guidance by February 11, 2020 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance 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–4751 for “FDARA Implementation Guidance for Pediatric Studies of Molecularly Targeted Oncology Drugs.” 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 copies total. One copy will include the information you claim to be confidential with a heading or cover note that states