DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee on Breast Cancer in Young Women (ACBCYW)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC), announces the following meeting of the aforementioned committee:

Time and Date: 1:00 p.m.–5:00 p.m. EST, December 13, 2016.

Place: This meeting will be held via Teleconference and web access. Teleconference and web access login information is as follows:


Conference number: PWXW1545545. Audience passcode: 3895011.

Participants can join the event directly at: https://www.mymeetings.com/nc/join/.

Conference number: PWXW1545545. Audience passcode: 3895011.

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 for Discussion: The agenda will include discussions on the current and emerging topics related to breast cancer in young women. These will include public health communication, breast cancer in young women digital and social media campaigns, and CDC updates. Committee workgroups will report findings to the committee.

Agenda items are subject to change as priorities dictate.

Online Registration Required: All ACBCYW Meeting participants must register for the meeting online at least 3 business days in advance at http://www.cdc.gov/cancer/breast/what_cdc_is_doing/meetings.htm. Please complete all the required fields before submitting your registration and submit no later than December 8, 2016.

Contact Person for More Information: Temeika L. Fairley, Ph.D., Designated Federal Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 5770 Buford Hwy, NE., Mailstop K52, Atlanta, Georgia 30341, Telephone (770) 488–4518, Fax (770) 488–4760. Email: acbcyw@cdc.gov.

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 Agency for Toxic Substances and Disease Registry.

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

[D74, Atlanta, Georgia 30329. Phone: 404–639–7570.
Email: omb@cdc.gov.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–17–17BZ]; Docket No. CDC–2016–0104]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project entitled “Project Pride.” This project is funded by CDC at 12 health departments in the United States. The health departments will report standardized program monitoring and evaluation (M&E) data to CDC. CDC is requesting approval to collect standardized HIV prevention program evaluation data from funded health departments.

DATES: Written comments must be received on or before January 3, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2017–0104 by any of the following methods:

• Federal eRulemaking Portal: Regulations.gov. Follow the instructions for submitting comments.

• Mail: Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to Regulations.gov, including any personal information provided. For access to the docket to read background documents or comments received, go to Regulations.gov.

Please note: All public comment should be submitted through the Federal eRulemaking portal (Regulations.gov) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact the Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: omb@cdc.gov.

SUPPLEMENTARY INFORMATION:

Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the Federal Register concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

Comments are invited 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) ways to enhance the quality, utility, and clarity of the information to be