

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 Discussed: The agenda will include discussions on evidence-based recommendations and the public health aspects of breast cancer in young women including biology, genomics, prevention, early diagnosis, treatment, and survivorship; appropriate venues to educate women at increased risk for developing breast cancer at younger ages; and approaches to increase awareness of clinicians/practitioners regarding topics such as breast health, symptoms, diagnosis, and treatment of breast cancer in young women.

Agenda items are subject to change as priorities dictate.

In order to assure that sufficient space and materials are available for meeting attendees, CDC is requesting that potential attendees register to attend this meeting at the following Web site: http://www.cdc.gov/cancer/breast/what_cdc_is_doing/conference.htm.

Contact Person for More Information: Temeika L. Fairley, PhD, 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.

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.

Dated: December 28, 2010.

Elaine Baker,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: National Survey of Child and Adolescent Well-Being—Second Cohort (NSCAW II).

OMB No.: 0970-0202.

Billing Accounting Code (BAC): 418422 (G994426).

Description: The Department of Health and Human Services (HHS)

intends to collect follow-up data on a sample of children and families for the National Survey of Child and Adolescent Wellbeing (NSCAW). The NSCAW was authorized under Section 427 of the Personal Responsibility and Work Opportunities Reconciliation Act of 1996. The NSCAW is the only source of nationally representative, firsthand information about the functioning and well-being, service needs, and service utilization of children and families who come to the attention of the child welfare system. Information is collected about children's cognitive, social, emotional, behavioral, and adaptive functioning, as well as family and community factors that are likely to influence their functioning. Family service needs and service utilization also are addressed in the data collection.

Selection of the current NSCAW sample and baseline data collection began in 2007 with a final sample size of 5,873 children. The proposed data collection will allow for follow-up of this sample 36 months post-baseline, will follow the same format as that used in the baseline round and the 18-month follow-up, and will employ, with only modest revisions, the same instruments that were used in the previous rounds. Data from NSCAW are made available to the research community through licensing arrangements from the National Data Archive on Child Abuse and Neglect at Cornell University.

Respondents: Children and their associated permanent or foster caregivers, caseworkers, and teachers.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Child Interview	1,424	1	1.33	1,894
Caregiver Interview	1,424	1	1.9	2,704
Caseworker Interview	285	3	1	855
Teacher Questionnaire	855	1	.50	428

Estimated Total Annual Burden Hours: 5, 882

In compliance with the requirements of Section 3506(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:** OPRE Reports Clearance Officer. **E-mail address:** OPREinfocollection@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.

Dated: December 29, 2010.

Steven M. Hanmer,

Reports Clearance Officer.

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