

(NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In 2010, the National Intimate Partner and Sexual Violence Surveillance System (NISVSS) reported that approximately 6.9 million women and 5.6 million men experienced rape, physical violence and/or stalking by an intimate partner within the last year. The health care costs of Intimate Partner Violence (IPV) exceed \$5.8 billion each year, nearly \$3.9 billion of which is for direct medical and mental health care services.

In order to address this important public health problem, CDC implemented, beginning in 2010, the National Intimate Partner and Sexual Violence Surveillance System that produces national and state level estimates of Intimate Partner Violence (IPV), Sexual Violence (SV) and stalking on an annual basis.

This revision request is multi-faceted. CDC is requesting a continuation of data collection among non-institutionalized

adult men and women aged 18 years or older in the United States assessing lifetime experiences of IPV, SV and stalking with a new and improved data collection tool. The revisions to the survey are aimed at reducing the time and complexity of the instrument, thus reducing the burden on the respondent. The simplified structure of the instrument will also reduce the complexity of the data set, making it more assessable for public use. Additionally, in collaboration with the Department of Defense (DoD), NISVS will collect information regarding the experiences of IPV, SV and stalking among active duty women and men in the military and wives of active duty men. This data collection will take place during the first three months of data collection.

To comply with OMB requirements, CDC is in the process of developing an expert panel to address methodological issues with the NISVS survey. The panel will meet multiple times over the course of the next year. The members of this panel will provide guidance on how

to improve both survey design (methods, sampling frame, recruitment, mode of administration) and content/question wording with the goals of increasing response rates, reducing non-response bias, and maximizing the opportunities across Federal surveys for covering populations of interest. This change request also encompasses the implementation of the panel's recommendations to improve the survey.

In the bi-annual data collection periods, total of 170,000 households will be screened. After determining eligibility and consent, 25,000 will complete the survey. The average burden per screened respondent remains at three minutes (total burden in hours equals 8,500) while the average burden per surveyed respondent is 25 minutes (total burden in hours equals 10,417). The survey will be conducted among English or Spanish speaking male and female adults (18 years and older) living in the United States.

There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Non-Participating Individuals (Screened)	NISVS Survey Instrument	170,000	1	3/60	8,500
Eligible Individuals (Surveyed)	NISVS Survey Instrument	25,000	1	25/60	10,417
Total					18,917

Leroy A. Richardson,

Chief, Information Collection Review Office,
Office of Scientific Integrity, Office of the
Associate Director for Science, Office of the
Director, Centers for Disease Control and
Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Safety and Occupational Health Study Section (SOHSS), National Institute for Occupational Safety and Health (NIOSH or Institute)

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 committee meeting.

Times and Dates:

8:00 a.m.–5:00 p.m., June 16, 2015 (Closed)
8:00 a.m.–5:00 p.m., June 17, 2015 (Closed)

Place: Embassy Suites, 1900 Diagonal Road, Alexandria, Virginia 22314.
Telephone: 703-684-5900, Fax: 703-684-0653.

Purpose: The Safety and Occupational Health Study Section will review, discuss, and evaluate grant application(s) received in response to the Institute's standard grants review and funding cycles pertaining to research issues in occupational safety and health, and allied areas.

It is the intent of NIOSH to support broad-based research endeavors in keeping with the Institute's program goals. This will lead to improved understanding and appreciation for the magnitude of the aggregate health burden associated with occupational injuries and illnesses, as well as to support more focused research projects, which will lead to improvements in the delivery of occupational safety and health services, and the prevention of work-related injury and illness. It is anticipated that research funded will promote these program goals.

Matters for Discussion: The meeting will convene to address matters related to the

conduct of Study Section business and for the study section to consider safety and occupational health-related grant applications.

These portions of the meeting will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, Centers for Disease Control and Prevention, pursuant to Section 10(d) Public Law 92-463.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Price Connor, Ph.D., NIOSH Health Scientist, CDC, 2400 Executive Parkway, Mailstop E-20, Atlanta, Georgia 30345, Telephone: (404) 498-2511, Fax: (404) 498-2571.

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 CDC and the Agency for Toxic Substances and Disease Registry.

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Initial Review

The meeting announced below concerns Integrating Self-Management Education with Cancer Survivorship Care Planning, SIP 15–001, and Using Cancer Registry Data to Promote Proactive Tobacco Cessation among Adult Cancer Survivors, SIP 15–003, initial review.

SUMMARY: This document corrects a notice that was published in the **Federal Register** on April 14, 2015 Volume 80, Number 86, Page 19990. The time and date should have read as follows:

DATES: *Time and Date:* 11:00 a.m.–6:00 p.m., May 12, 2015 (Closed).

FOR FURTHER INFORMATION CONTACT: Brenda Colley Gilbert, Ph.D., M.S.P.H., Director, Extramural Research Program Operations and Services, CDC, 4770

Buford Highway NE., Mailstop F–80, Atlanta, Georgia 30341, Telephone: (770) 488–6295, BJC4@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 the Agency for Toxic Substances and Disease Registry.

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

[FR Doc. 2015–12053 Filed 5–18–15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Assets for Independence Program Performance Progress Report.
OMB No.: New.

Description

The Assets for Independence (AFI) Act (Title IV of the Community Opportunities, Accountability, and Training and Educational Services Act

of 1998, Pub. L. 105–285, [42 U.S.C. 604 note]) requires that organizations operating AFI projects submit annual progress reports.

This request is to create an AFI program specific Performance Progress Report (PPR) to replace the semiannual standard form performance progress report (SF–PPR) and the annual data report. The AFI PPR will collect data on project activities and attributes similar to the reports that it is replacing. The Office of Community Services (OCS) in the Administration for Children and Families (ACF) will use the data collected in the AFI PPR to prepare the annual AFI Report to Congress, to evaluate and monitor the performance of the AFI program overall and of individual projects, and to inform and support technical assistance efforts. The AFI PPR would fulfill AFI Act reporting requirements and program purposes.

The AFI PPR will be submitted quarterly: Three times per year using an abbreviated short form and one time using a long form. Both draft data collection instruments are available for review online at <https://idaresources.acf.hhs.gov/AFIPPR>.

Note: This request does not affect financial reporting requirements for AFI grantees. The SF–425 will still be required semiannually throughout the grant project period with a final report due 90 days after the grant project period ends.

Respondents: Assets for Independence (AFI) program grantees.

ANNUAL BURDEN ESTIMATES				
Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
AFI PPR Short Form	300	3	0.5	450
AFI PPR Long Form	300	1	4	1200

Estimated Total Annual Burden Hours: 1,650.

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.

Karl Koerper,
Reports Clearance Officer.

[FR Doc. 2015–12096 Filed 5–18–15; 8:45 am]

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