follow the prompts. Once you register, you will receive a confirmation email with the webinar login and teleconference number. The room only has capacity for 115 people, so please register early if you plan to attend in person. Members of the public will not have the opportunity to ask questions or otherwise participate in the meeting.

However, members of the public wishing to comment should follow the steps detailed under the heading Addresses in this publication or contact us via the CECANF Web site at https://

eliminatechildabusefatalities.sites.usa. gov/contact-us/.

Detailed meeting minutes will be posted within 90 days of the meeting on the CECANF Web site in the Events section: https://eliminatechildabuse fatalities.sites.usa.gov/events/

Dated: November 13, 2014.

Karen White,

Executive Assistant.

[FR Doc. 2014–27437 Filed 11–19–14; 8:45 am] BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: HHS-OS-0990-NEW-60D]

Agency Information Collection Activities; Proposed Collection; Public Comment Request

AGENCY: Office of the Assistant Secretary for Health, Office of Adolescent Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, announces plans to submit a new Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting that ICR to OMB, OS seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR. **DATES:** Comments on the ICR must be received on or before January 20, 2015.

ADDRESSES: Submit your comments to Information.CollectionClearance@ hhs.gov or by calling (202) 690–6162. FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff.

Information Collection Clearance stati Information.CollectionClearance@ hhs.gov or (202) 690–6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the document identifier HHS–OS–0990– NEW–60D for reference.

Information Collection Request Title: Pregnancy Assistance Fund Feasibility and Design Study (FADS).

Abstract: The Office of Adolescent Health (OAH), U.S. Department of Health and Human Services (HHS) is requesting approval by OMB on a new collection. The Pregnancy Assistance Fund (PAF) evaluation will provide information about program design, implementation, and impacts through two core components: A rigorous assessment of program impacts and implementation, and a descriptive examination of program design. This proposed information collection request includes instruments related to the indepth implementation study that complements the impact study. The data collected from these instruments will provide a detailed understanding of program implementation.

Need and Proposed Use of the Information: The data will serve two main purposes. First, the information will enable the study team to produce clear, detailed descriptions of each intervention that is evaluated and the counterfactual in each site. This documentation is critical for understanding the meaning of impact estimates. Second, the data will be used to assess fidelity of implementation and the quality of program delivery. This information is essential for determining whether the interventions were implemented well and whether the evaluation provided a good test of each site's intervention.

Likely Respondents: The 140 program administrators and case managers and 200 youth participants in 3 impact study sites.

TOTAL ESTIMATED ANNUALIZED BURDEN-HOURS

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Semi-structured interview	12 17 47' 2 67	2 1 1 12 1	1 1 .5 .5 1.5	24 17 24 12 100.5
Total				177.5

OS specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Darius Taylor,

Information Collection Clearance Officer. [FR Doc. 2014–27473 Filed 11–19–14; 8:45 am] BILLING CODE 4168–11–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-15-0920]

Proposed Data Collections Submitted for Public Comment and Recommendations

The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden, 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. To request more information on the below proposed project or to obtain a copy of the information collection plan and instruments, call 404–639–7570 or send comments to Leroy A. Richardson, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an email to *omb@cdc.gov*.

Comments submitted in response to this notice will be summarized and/or included in the request for Office of Management and Budget (OMB) approval. 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 collected; (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; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information. Written comments should be received within 60 days of this notice.

Proposed Project

Data Collection Through Web Based Surveys for Evaluating Act Against AIDS Social Marketing Campaign Phases Targeting Consumers (Generic ICR, OMB# 0920–0920, Expires 2/28/ 2015)—Extension—National Center for HIV/AIDS, Viral Hepatitis, STD and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In response to the continued HIV epidemic in our country, CDC has launched Act Against AIDS, a 5-year, multifaceted communication campaign to reduce HIV incidence in the United States. CDC plans to release the campaign in phases, with some of the phases running concurrently. Each phase of the campaign will use mass media and direct-to-consumer channels to deliver HIV prevention and testing messages. Some components of the campaign will be designed to provide basic education and increase awareness of HIV/AIDS among the general public, and others will be targeted to specific

subgroups or communities at greatest risk of infection. The current study addresses the need to assess the effectiveness of these social marketing messages aimed at increasing HIV awareness and delivering HIV prevention and testing messages among at-risk populations.

This extension of an ongoing study will evaluate the *Act Against AIDS* (AAA) social marketing campaign aimed at increasing HIV/AIDS awareness, increasing prevention behaviors, and improving HIV testing rates among consumers. A total of 12,000 respondents were originally approved for this generic ICR (0920-0920) and since the original approval date, 1,250 respondents have participated in the surveys under the following mini ICRs: 0920-13AHP; 0920-13YR and 0920-13DD. The information collected from each of the data collections were used to evaluate specific AAA campaign phases. We are requesting additional time to continue to survey other AAA target audiences and campaign phases and measuring exposure to each phase of the campaign and interventions implemented under AAA. Through this extension, we plan to reach the remaining approved 10,750 respondents. In order to obtain the remaining respondents, we anticipate screening approximately 17,915 individuals.

Depending on the target audience for the campaign phase, the study screener will vary. The study screener may address one or more of the following items: race/ethnicity, sexual behavior, and sexual orientation. Each survey will have a core set of items asked in all rounds, as well as a module of questions relating to specific AAA activities and communication initiatives.

Survey respondents will be selected from a combination of sources, including a national opt-in email list sample and respondent lists generated by partnership organizations (e.g., the National Urban League, the National Medical Association). A total of 10,750 participants will self-administer the survey at home on personal computers over a 3-year period.

There is no cost to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Individuals (male and female) aged 18 years and older.	Study Screener	17,915	1	2/60	597