data collected from the updates is also shared with the Division of Grants Management Operations (DGMO) for their assistance in the overall evaluation of each project's progress.

An electronic form is currently being used for progress reporting for the HCOF program. This form provides awardees access to directly input the required status update information in a timely, consistent, and uniform manner. The electronic form minimizes burden to respondents and informs respondents when there are missing data elements prior to submission. We acknowledge a change in the burden estimate due to close out of old projects, and the addition of new projects for FY 2010.

The annual estimate of burden is as follows:

Instrument	Number of respondents	Responses per respondent	Total responses	Hours per response	Total Burden hours
Construction-Related	481	4	1,924	.5	962
Equipment Only	1,238	1	1,238	.5	619
Total	1,719		3,162		1,581

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. Please direct all correspondence to the "attention of the desk officer for HRSA."

Dated: August 3, 2010.

Sahira Rafiullah,

Director, Division of Policy and Information Coordination.

[FR Doc. 2010–19549 Filed 8–6–10; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day-10–0783]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Evaluation of Safe Dates Project—(OMB No. 0920–0783 exp. 6/30/2011)—Revision—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Safe Dates, a dating violence prevention curriculum for 8th and 9th grade students, has been shown to be effective at preventing victimization and perpetration of teen dating violence in one rural North Carolina school district, but appropriateness of the program with urban, high-risk adolescents is unknown. The data collection will require participation from teachers at eight schools who delivered the Safe Dates program and students at one school who received the program. Qualitative data will be collected through student focus groups and teacher interviews. Students will complete a participant profile form to capture basic demographic information. The specific aim of this study is to assess whether the Safe Dates adolescent dating violence prevention program needs modification/adaptation for urban, high-risk adolescents.

Approximately 40 students at one school will participate in focus groups. Two focus groups will consist of 8–10 boys, and two focus groups will include 8–10 girls. Informed written consent from parents for each student's participation and informed written assent from tenth graders for their own participation will be obtained. Twenty teachers will participate in interviews. Students and teachers will be asked about their experiences with the Safe Dates program and ideas they may have about adapting the program for urban schools.

Data collection will occur in July 2010. It is anticipated that study results will be used to determine whether the Safe Dates program should be modified for an urban, high-risk population. There is no cost to respondents other than their time. The total estimated annual burden hours are 849.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Instrument name	Number of respondents	Number of responses per respondent	Average burden per respondent (in hours)	Total response burden (hours)
Student	Effectiveness follow-up survey	1,318	1	35/60	769
	Focus group guide and participant profile form.	40	1	1.5	60
Teacher	Interview guide	20	1	1	20

Dated: August 3, 2010.

Maryam I. Daneshvar,

Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. 2010–19555 Filed 8–6–10; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Center for Substance Abuse Treatment; Notice of Meeting

Pursuant to Public Law 92–463, notice is hereby given that the Substance Abuse and Mental Health Services Administration's (SAMHSA) Center for Substance Abuse Treatment (CSAT) National Advisory Council will meet August 19, 2010, 1–3 p.m. via teleconference.

The meeting will include discussion and evaluation of grant applications reviewed by Initial Review Groups. Therefore, the meeting will be closed to the public as determined by the Administrator, SAMHSA, in accordance with Title 5 U.S.C. 552b(c)(6) and 5 U.S.C. App. 2, Section 10(d).

Substantive program information, a summary of the meeting and a roster of Council members may be obtained as soon as possible after the meeting, either by accessing the SAMHSA Committee Web site at *https://nac.samhsa.gov/CSATcouncil/index.aspx,* or by contacting the CSAT National Advisory Council Designated Federal Official, Ms. Cynthia Graham (*see* contact information below).

Committee Name: SAMHSA's Center for Substance Abuse Treatment National Advisory Council.

Date/Time/Type: August 19, 2010, 1– 3 p.m.: Closed.

Place: SAMHSA Building, 1 Choke Cherry Road, Great Falls Conference Room, Rockville, Maryland 20857.

Contact: Cynthia Graham, M.S., Designated Federal Official, SAMHSA CSAT National Advisory Council, 1 Choke Cherry Road, Room 5–1035, Rockville, Maryland 20857, Telephone: (240) 276–1692, Fax: (240) 276–1690, e-mail: *cynthia.graham@samhsa.hhs.gov.*

Dated: August 3, 2010.

Toian Vaughn,

Committee Management Officer, Substance Abuse and Mental Health, Services Administration.

[FR Doc. 2010–19539 Filed 8–6–10; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Alcohol Abuse and Alcoholism; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App), notice is hereby given of a meeting of the National Advisory Council on Alcohol Abuse and Alcoholism.

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable materials, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Advisory Council on Alcohol Abuse and Alcoholism.

Date: September 22–23, 2010.

Closed: September 22, 2010, 5:30 p.m. to 7:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 5635 Fishers Lane, Bethesda, MD 20892.

Open: September 23, 2010, 9 a.m. to 3 p.m.

Agenda: Presentations and other business of the council.

Place: National Institutes of Health, 5635 Fishers Lane, Bethesda, MD 20892.

Contact Person: Abraham P. Bautista, PhD, Executive Secretary, National Institute on Alcohol Abuse and Alcoholism, National Institutes of Health, 5635 Fishers Lane, Room 2085, Rockville, MD 20892, 301–443–9737. *bautistaa@mail.nih.gov.*

Information is also available on the Institute's/Center's home page: *http:///www.silk.nih.gov/silk/niaaa1/about/roster.htm,* where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos.: 93.271, Alcohol Research Career Development Awards for Scientists and Clinicians; 93.272, Alcohol National Research Service Awards for Research Training; 93.273, Alcohol Research Programs; 93.891, Alcohol Research Center Grants; 93.701, ARRA Related Biomedical Research and Research Support Awards, National Institutes of Health, HHS)

Dated: July 29, 2010.

Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–19558 Filed 8–6–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0001]

Workshop on Optimizing Clinical Trial Design for the Development of Pediatric Cardiovascular Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

The Food and Drug Administration (FDA) and National Institutes of Health (NIH), with support from the American Academy of Pediatrics (AAP), the American College of Cardiology (ACC), and the Society for Cardiovascular Angiography and Interventions (SCAI) are announcing a public workshop entitled "Optimizing Clinical Trial Design for the Development of Pediatric Cardiovascular Devices." The topic to be discussed is pediatric cardiovascular device development. The purpose of the public workshop is to solicit information from clinicians, academia, professional societies, other government agencies, and industry on various efficient and pragmatic clinical trial designs that are conducive to overcoming the challenges in developing devices for the pediatric cardiology market. The information gathered in this and future workshops will help to develop future guidance on optimal designs for pediatric cardiology device trials.

Date and Time: The public workshop will be held on September 30, 2010, from 8 a.m. to 5:30 p.m.

Location: The public workshop will be held at Moscone Center, 747 Howard St., San Francisco, CA 94103.

Contact Person: Francesca Joseph, Office of Orphan Products Development, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 5277, Silver Spring, MD 20903, 301–796–6805, FAX: 301–847–8621, e-mail: *francesca.joseph@fda.hhs.gov*.

Registration: Registration information will be posted on the Internet at *http://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/default.htm*.