

### Subcontracting Reporting System (eSRS).

Contractors must use the SSR in lieu of the SF 295, with the exception of those contracts noted in FAR 4.606(c)(5) which requires that actions, pursuant to other authority, will not be entered in Federal Procurement Data System (e.g., reporting of the information would compromise national security). Those contract actions noted in FAR 4.606(c)(5) will continue to use the Standard Form 295.

## II. Annual Reporting Burden

Based on information from eSRS and an estimate of the use of eSRS, an upward adjustment is being made to the number of respondents, but a downward adjustment is being made to the average burden hours for reporting and recordkeeping per response. As a result, a downward adjustment is being made to the estimated annual reporting burden since the notice regarding an extension to this clearance published in the **Federal Register** at 75 FR 9603, on March 3, 2010.

*Respondents:* 129,009.

*Responses per Respondent:* 1.

*Total Responses:* 129,009.

*Average Burden Hours per Response:* 9.0.

*Total Burden Hours:* 1,161,081.

*Obtaining Copies of Proposals:*

Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (MVCB), 1275 First Street NE., Washington, DC 20417, telephone (202) 501-4755. Please cite OMB Control Number 9000-0007, Summary Subcontract Report, in all correspondence.

Dated: November 8, 2012.

**William Clark,**

*Acting Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.*

[FR Doc. 2012-28067 Filed 11-16-12; 8:45 am]

**BILLING CODE 6820-EP-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Meeting of the Advisory Committee on Minority Health

**AGENCY:** Office of Minority Health, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

**ACTION:** Notice of meeting.

**SUMMARY:** As stipulated by the Federal Advisory Committee Act, the

Department of Health and Human Services (DHHS) is hereby giving notice that the Advisory Committee on Minority Health (ACMH) will hold a meeting. This meeting will be open to the public. Preregistration is required for both public attendance and comment. Any individual who wishes to attend the meeting and/or participate in the public comment session should email [acmh@osophs.dhhs.gov](mailto:acmh@osophs.dhhs.gov).

**DATES:** The meeting will be held on Thursday, November 29, 2012 from 9:00 a.m. to 5:00 p.m. and Friday, November 30, 2012 from 9:00 a.m. to 1:00 p.m.

**ADDRESSES:** The meeting will be held at the Doubletree Hotel, 1515 Rhode Island Avenue, Washington, DC 20005.

**FOR FURTHER INFORMATION CONTACT:** Ms. Monica A. Baltimore, Tower Building, 1101 Wootton Parkway, Suite 600, Rockville, Maryland 20852. Phone: 240-453-2882, Fax: 240-453-2883.

**SUPPLEMENTARY INFORMATION:** In accordance with Public Law 105-392, the ACMH was established to provide advice to the Deputy Assistant Secretary for Minority Health in improving the health of each racial and ethnic minority group and on the development of goals and specific program activities of the Office of Minority Health.

Topics to be discussed during these meetings will include strategies to improve the health of racial and ethnic minority populations through the development of health policies and programs that will help eliminate health disparities, as well as other related issues.

Public attendance at this meeting is 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 designated contact person at least seven (7) business days prior to the meeting. Members of the public will have an opportunity to provide comments at the meeting. Public comments will be limited to three minutes per speaker. Individuals who would like to submit written statements should mail or fax their comments to the Office of Minority Health at least seven (7) business days prior to the meeting. Any members of the public who wish to have printed material distributed to ACMH committee members should submit their materials to the Executive Director, ACMH, Tower Building, 1101 Wootton Parkway, Suite 600, Rockville, Maryland 20852, prior to close of business November 26, 2012.

Dated: November 9, 2012.

**Monica A. Baltimore,**

*Executive Director, Advisory Committee on Minority Health.*

[FR Doc. 2012-27988 Filed 11-16-12; 8:45 am]

**BILLING CODE 4150-29-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[30Day-13-0008]

### 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 (404) 639-7570 or send an email to [omb@cdc.gov](mailto:omb@cdc.gov). Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395-5806. Written comments should be received within 30 days of this notice.

### Proposed Project

Emergency Epidemic Investigations (0920-0008)—Revision—Scientific Education and Professional Development Program Office (SEPDPPO), Office of Surveillance, Epidemiology, and Laboratory Services (OSELS), Centers for Disease Control and Prevention (CDC).

### Background and Brief Description

One of the objectives of CDC's epidemic services is to provide for the prevention and control of epidemics, and protect the population from public health crises such as human-made or natural biological disasters and chemical emergencies. CDC meets this objective, in part, by training investigators, maintaining laboratory capabilities for identifying potential problems, collecting and analyzing data, and recommending appropriate actions to protect the public's health. When state, local, or foreign health authorities request help in controlling an epidemic or solving other health problems, CDC dispatches skilled epidemiologists from the Epidemic Intelligence Service (EIS) to investigate and resolve the problem. Resolving public health problems rapidly ensures cost-effective health care and enhances health promotion and disease prevention.

The purpose of the Emergency Epidemic Investigation data collection

project is to collect data on the conditions surrounding and preceding the onset of a problem. The data must be collected in a timely fashion so that information can be used to develop prevention and control techniques, to interrupt disease transmission and to help identify the cause of an outbreak. Since the events necessitating the collections of information are of an emergency nature, most data collection is done by direct interview or written questionnaire and are one-time efforts related to a specific outbreak or circumstance. If during the emergency investigation, the need for further study is recognized, a project is designed and separate OMB clearance is required. Interviews are conducted to be as unobtrusive as possible and only the minimal information necessary is collected. The Emergency Epidemic Investigations data collection project is the principal source of data on outbreaks of infectious and noninfectious diseases, injuries,

nutrition, environmental health, and occupational problems.

Each investigation contributes to the general knowledge about a particular type of problem or emergency, so that data collections are designed taking into account knowledge gained during similar situations in the past. Some questionnaires have been standardized, such as investigations of outbreaks aboard aircraft or cruise vessels.

The Emergency Epidemic Investigations data collection project provides a range of data on the characteristics of outbreaks and those affected by outbreaks. Data collected include demographic characteristics of the affected population, exposure to the causative agent(s), transmission patterns, and severity of the outbreak. These data, together with trend data, may be used to monitor the effects of change in the health care system, plan health services, improve the availability of medical services, and assess the health status of the population.

Users of the Emergency Epidemic Investigations data include, but are not limited to, Epidemic Intelligence Service (EIS) officers of the CDC, who investigate the patterns of disease or injury, the level of risky behaviors, causative agents, the transmission of the condition, and the impact of interventions. EIS is a two-year program of training and service in applied epidemiology through CDC, primarily for persons holding doctoral degrees.

Predicting the number of epidemic investigations that might occur in any given year is difficult. The previous three years' experience shows an annualized burden of 3,750 hours and respondent total of 15,000. After completion of the Epi-Aid investigation, using the Epi-Aid Satisfaction Survey for Requesting Officials, data are collected from 100 state and local health officials for an annualized total of 25 burden hours. For this clearance, CDC is requesting 3,775 total burden hours.

**ESTIMATED ANNUALIZED BURDEN HOURS**

Respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Requestors of Epi-Aids .....	Epi-Aid Satisfaction Survey for Requesting Official.	100	1	15/60
General Public .....	Emergency Epidemic Investigations .....	15,000	1	15/60

Dated: November 13, 2012.

**Ron A. Otten,**

Director, Office of Scientific Integrity (OSI), Office of the Associate Director for Science (OADS), Office of the Director, Centers for Disease Control and Prevention.

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**BILLING CODE 4163-18-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[60Day-13-13BU]

**Proposed Data Collections Submitted for Public Comment and Recommendations**

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of

the data collection plans and instruments, call 404-639-7570 or send comments to Ron Otten, 1600 Clifton Road, MS D-74, Atlanta, GA 30333 or send an email to [omb@cdc.gov](mailto:omb@cdc.gov).

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; 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. Written comments should be received within 60 days of this notice.

**Proposed Project**

Determining Causes of Sudden, Unexpected Infant Death: A National Survey of U.S. Medical Examiners and Coroners—New—National Center for Chronic Disease Prevention and Health

Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

Each year in the United States, approximately 4,200 infants die suddenly without any cause that is immediately obvious. Half of these sudden unexpected infant deaths (SUID) are attributed to Sudden Infant Death Syndrome (SIDS), which is the leading cause of death in infants between one and twelve months of age. Reducing deaths caused by SIDS and other SUID such as accidental suffocation are important public health priorities.

Between 1990 and 2001, the rate of SIDS in the U.S. decreased from 1.3 per 1,000 live births to 0.56 deaths per 1,000 live births. The 50% decline in SIDS is attributed to the success of the "Back to Sleep" campaign, launched in 1994, during which prone sleeping for infants decreased from about 75% in 1992 to 12% in 2002. SIDS has continued to decline slightly and in 2009 was estimated to be 0.525 deaths per 1,000, however, post-neonatal mortality due to other causes increased, particularly in 1999-2001. Further examination of the cause-specific age at death and month of