

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 fourteen (14) 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 Secretary, ACMH, Tower Building, 1101 Wootton Parkway, Suite 600, Rockville, Maryland 20852, prior to close of business October 13, 2009.

Dated: August 31, 2009.

Garth Graham,

Deputy Assistant Secretary for Minority Health, Office of Minority Health, Office of Public Health and Science, Office of the Secretary, U.S. Department of Health and Human Services.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-09-0607]

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) 395-6974. Written comments should be received within 30 days of this notice.

Proposed Project

The National Violent Death Reporting System (NVDRS)—[OMB# 0920-0607, exp.01/31/2010]—Revision—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Violence is an important public health problem. In the United States, homicide and suicide are the second and third leading causes of death, respectively, in the 1-34-year-old age group. Unfortunately, public health agencies do not know much more about the problem than the numbers and the sex, race, and age of the victims, all information obtainable from the standard death certificate. Death certificates, however, carry no information about key facts necessary for prevention such as the relationship of the victim and suspect and the circumstances of the deaths, thereby making it impossible to discern anything but the gross contours of the problem. Furthermore, death certificates are typically available 20 months after the completion of a single calendar year. Official publications of national violent death rates, e.g. those in Morbidity and Mortality Weekly Report, rarely use data that is less than two years old. Public health interventions aimed at a moving target last seen two years ago may well miss the mark.

Local and Federal criminal justice agencies such as the Federal Bureau of Investigation (FBI) provide slightly more information about homicides, but they do not routinely collect standardized data about suicides, which are in fact much more common than homicides. The FBI's Supplemental Homicide Report system (SHRs) does collect basic information about the victim-suspect relationship and circumstances, like death certificates, it does not link violent deaths that are part of one incident such as homicide-suicides. It also is a voluntary system in which

some 10-20 percent of police departments nationwide do not participate. The FBI's National Incident Based Reporting System (NIBRS) addresses some of these deficiencies, but it covers less of the country than SHRs, still includes only homicides, and collects only police information. Also, the Bureau of Justice Statistics Reports do not use data that is less than two years old.

CDC therefore proposes to continue a state-based surveillance system for violent deaths that will provide more detailed and timely information. It taps into the case records held by medical examiners/coroners, police, and crime labs. Data is collected centrally by each state in the system, stripped of identifiers, and then sent to the CDC. Information is collected from these records about the characteristics of the victims and suspects, the circumstances of the deaths, and the weapons involved. States use standardized data elements and software designed by CDC. Ultimately, this information will guide states in designing programs that reduce multiple forms of violence.

Neither victim families nor suspects are contacted to collect this information. It all comes from existing records and is collected by state health department staff or their subcontractors. Health departments incur an average of 2.0 hours per death in identifying the deaths from death certificates, contacting the police and medical examiners to get copies of or to view the relevant records, abstracting all the records, various data processing tasks, various administrative tasks, data utilization, training, communications, etc. Public agencies working with NVDRS states incur an average of 0.5 hours per death to retrieve and then refile records.

This revision is a request to allow 10 new state health departments to be added to the currently funded 17 if funding becomes available. This may bring the total to 27 by the year 2012. There are no costs to respondents other than their time. The total estimated annual burden hours are 67,500.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
State Health Departments	27	1,000	2.0
Public Agencies	27	1,000	30/60

Dated: September 9, 2009.

Maryam I. Daneshvar,
Acting Reports Clearance Officer, Centers for Disease Control and Prevention.
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2008-N-0487]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food Safety Survey

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by October 15, 2009.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202-395-6974, or e-mailed to *oira_submission@omb.eop.gov*. All comments should be identified with the OMB control number 0910-0345. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Information Management (HFA-710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-796-3794, *JonnaLynn.Capezzuto@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Food Safety Survey—(OMB Control Number 0910-0345—Reinstatement)

Under section 903(b)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393(b)(2)), FDA is authorized to conduct research relating to foods and to conduct educational and public information programs relating to the safety of the nation's food supply. The Food Safety Survey is a nationally representative survey of consumers' knowledge, attitudes, and beliefs about food safety. Previous versions of the survey were collected in 1988, 1993, 1998, 2001, and 2006. Data from the previous surveys are being used to evaluate two Healthy People 2010 objectives: (1) Increase the proportion of consumers who follow key food safety practices (Objective 10-5), and (2) reduce severe allergic reactions to food among adults (Objective 10-4b). Additionally, data are used to measure trends in consumer food safety habits including hand and cutting board washing, cooking practices, and use of food thermometers. Finally, data are used to evaluate educational messages and to inform policymakers about consumer attitudes about novel technologies such as food irradiation and biotechnology.

Since 2006, there have been several high profile recalls of FDA-regulated food due to contamination. Information

about food recalls does not always reach the intended audience (Refs. 1, 2, and 3). The Food Safety Survey planned for 2009 will look specifically at reasons why consumers do not always heed food recall alerts. A new food recall module will be added that contains new questions to learn about how recent food recalls have affected consumer confidence in the food supply and what effect, if any, they have on consumers' home food safety behaviors. This information will help FDA develop strategies to more effectively communicate food recall information to the public.

The methods for the 2009 version of the Food Safety Survey will be the same as for the previous Food Safety Surveys. A nationally representative sample of 4,000 adults in households with telephones will be selected at random and interviewed by telephone. This survey will include an oversample of Hispanics with a minimum of 500 Hispanics sampled. Additionally, 200 initial nonrespondents will be asked to participate in a short version of the survey to conduct a nonresponse analysis. Participation will be voluntary. Cognitive interviews and a pretest will be conducted prior to fielding the survey.

In the **Federal Register** of September 17, 2008 (73 FR 53878), FDA published a 60-day notice requesting public comment on the proposed collection of information. The agency received one comment that was not responsive to the comment request on the information collection provisions.

FDA estimates the burden of this collection of information as follows:

The total estimated burden imposed by this collection of information is 1,541 hours (table 1 of this document).

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN¹

Activity	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Cognitive Interview	20	1	20	1	20
Pretest	27	1	27	0.5	14
Screener	10,000	1	10,000	.0167	167
Survey	4,000	1	4,000	.33	1,320
Nonresponse	200	1	200	.10	20
Total					1,541

¹ There are no capital costs or operating and maintenance costs associated with this collection of information.

Prior to finalizing the survey, FDA will conduct 20 cognitive interviews each requiring an average of 1 hour per

respondent for a total of 20 hours. Before the survey is fielded, a small pretest of 27 individuals, each lasting

half an hour (0.5 hour), will be conducted. The survey screener is estimated to take 1 minute or less per