

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 the CDC Reports Clearance Officer on (404) 498-1210.

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. Send comments to Anne O'Connor, CDC Assistant Reports

Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project: Youth Risk Behavior Survey (YRBS) Methodological Study—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention, (CDC). CDC intends to conduct a methodological study in the Spring of 2004 to assess the effects of setting and mode of survey administration on the reporting of health-risk behaviors among adolescents, and thereby, to provide methodological guidance for future surveys, especially surveys of adolescents. In 2000, the Office of the Assistant Secretary for Planning and Evaluation (ASPE) commissioned five expert papers written on the topic "Examining Substance Abuse Data Collection Methodologies." The papers focused on the YRBS, the National Survey on Drug Use and Health

(NSDUH, formerly the National Household Survey on Drug Abuse, or NHSDA), and Monitoring the Future (MTF). A consensus among the authors was that disparate results across the studies are most likely a product of methodological differences across the surveys. This YRBS Methodological Study is designed to measure the extent to which the prevalence of health-risk behaviors among students varies by whether the survey is administered in schools vs. students' homes (setting), and by whether the survey is administered using paper-and-pencil questionnaire booklets vs. computer-assisted self-interviewing (mode). Approximately 5,376 high school students will be given questionnaires in one of the four setting/mode combinations. Elucidation of the impact of these factors on prevalence will assist in reducing response effects and improving the quality of the YRBS data. There are no costs to respondents.

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)	Total burden (in hrs.)
High school students	5,376	1	45/60	4,032
School administrators	104	1	45/60	78
Total				4,110

Dated: June 9, 2003.

Thomas A. Bartenfeld,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-03-77]

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 the CDC Reports Clearance Officer on (404) 498-1210.

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. Send comments to Anne O'Connor, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project: Hemophilia Treatment Center Laboratory Survey—New—National Center for Infectious Diseases (NCID), Centers for Disease Control and Prevention (CDC). Up to two million women in the United States may have an inherited bleeding disorder and not know it. Many women learn to

live with the problems their bleeding causes, such as heavy periods, and not realize that they could have a bleeding disorder. Other women may have more serious bleeding problems such as hemorrhages after childbirth or surgery, and some have hysterectomies to end their heavy periods. With proper diagnosis, women with bleeding disorders could avoid these complications and surgeries. Management of bleeding in these women can decrease heavy periods and can improve quality of life.

The most common bleeding disorder is called Von Willebrand disease (VWD). VWD is caused by a deficiency or defect in the body's ability to make a protein, von Willebrand factor, which helps blood clot. The symptoms of VWD can range in severity; however, 90 percent of people who have this disease have the mild form. VWD occurs in men and women equally, but women are more likely to notice the symptoms of VWD due to heavy or abnormal bleeding during their menstrual periods and after childbirth. There are many gynecological and physical causes for heavy periods, such as endometriosis, thyroid problems and cancer; however,

the cause is not identified in half the cases. A CDC-Emory University survey found that gynecologists rarely considered bleeding disorders as a cause of heavy menstrual bleeding. However, recent research from Europe and CDC has shown that 15–20 percent of women with heavy periods have inherited bleeding disorders. Women with VWD interviewed by CDC reported an average of 16 years between the onset of bleeding symptoms and diagnosis of a bleeding disorder. CDC and the National Hemophilia Foundation have been working to encourage gynecologists to consider bleeding disorders in women who have menorrhagia. As a result, the American College of Obstetricians and Gynecologists has recently recommended screening for VWD in these women.

An important part of increasing the awareness among physicians and their

patients with heavy periods who may have an underlying bleeding disorder is referral for appropriate diagnosis. Federally funded Hemophilia Treatment Centers (HTCs) are thought to be the best source for appropriate laboratory diagnosis, however, the following concerns have been raised: (1) Anecdotal reports from HTC providers describe reduced capacity of in-house laboratory support and access to specialty coagulation laboratory tests that are essential for appropriate diagnosis of bleeding disorders; (2) A CDC, Public Health Practice Program Office (PHPPO), study demonstrated reduced capacity to perform specific coagulation tests through their survey of hospital laboratories, but it is impossible to know if HTCs have higher capacity than the hospitals studied; (3) HTCs report that changes in third party

payer policies, especially health maintenance organizations, are dictating the source of laboratory testing requiring shipment of laboratory specimens to sites away from the hospital that reduce the quality of the sample and effect the reliability of the results. It is important to assess the HTCs and determine their capabilities and barriers to delivering comprehensive care to patients with bleeding disorders.

The proposed study will involve the 135 federally funded HTCs. The study participants are composed of medical directors, adult hematologists, pediatric hematologists, and coagulation laboratory technicians. A survey will be distributed to the above personnel to ascertain their perceptions of lab capabilities and procedures. There will be no cost to respondents.

Respondents	Number of respondents	Number responses per respondent	Average burden per response (in hrs.)	Total burden (in hrs.)
HTC medical directors & coagulation technicians	325	1	20/60	108
Total				108

Dated: June 9, 2003.

Thomas A. Bartenfeld,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 03-14912 Filed 6-12-03; 8:45 am]

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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: Cooperative Agreement for Development of the National Violent Death Reporting System, Program Announcement #03038

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 meeting:

Name: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Cooperative Agreement for Development of the National Violent Death Reporting System, Program Announcement #03038.

Times and Dates: 8:30 a.m.–8:50 a.m., June 30, 2003. (Open); 8:50 a.m.–4 p.m., June 30, 2003. (Closed)

Place: The Westin Atlanta North at Perimeter, 7 Concourse Parkway, Atlanta, GA 30328, Telephone 770.395.3900.

Status: 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, CDC, pursuant to Public Law 92-463.

Matters To Be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to Program Announcement #03038.

For Further Information Contact: James Belloni, Deputy Director, Office of Program Management and Operations, National Center For Injury Prevention and Control, CDC, 4770 Buford Highway NE., MS-K-62, Atlanta, GA 30341, Telephone 770.488.4538.

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.

Dated: June 5, 2003.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 03-14927 Filed 6-12-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Science and Program Review Subcommittee (SPRS) and the Advisory Committee for Injury Prevention and Control (ACIPC): Meetings

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 Subcommittee and Committee meetings.

Name: Science and Program Review Subcommittee to ACIPC.

Time and Dates: 6:30 p.m.–9 p.m., June 22, 2003; 9 a.m.–12 p.m., June 23, 2003.

Place: The Westin Atlanta Airport, 4736 Best Road, Atlanta, Georgia 30337, telephone (404) 762-7676.

Status: Open: 6:30 p.m.–7 p.m., June 22, 2003.

Closed: 7 p.m.–9 p.m., June 22, through 12 p.m., June 23, 2003.

Purpose: The Subcommittee provides advice on the needs, structure, progress, and performance of the National Center for Injury Prevention and Control (NCIPC) programs. They also provide second-level scientific and programmatic review for applications for research grants, cooperative agreements, and training grants related to injury control and violence prevention, and recommends