bioethics.gov. Persons needing special assistance, such as sign language interpretation or other special accommodations, should contact NBAC staff at the address or telephone number listed below as soon as possible.

FOR FURTHER INFORMATION CONTACT: Ms. Patricia Norris, National Bioethics Advisory Commission, 6100 Executive Boulevard, Suite 5B01, Rockville, Maryland 20892–7508, telephone 301–402–4242, fax number 301–480–6900.

Dated: July 1, 1999.

Eric M. Meslin,

Executive Director, National Bioethics Advisory Commission.

[FR Doc. 99–17243 Filed 7–7–99; 8:45 am]

BILLING CODE 4160–17–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY–15–99]

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) 639–7090. Send written comments to CDC, Desk Officer; Human Resources and Housing Branch, New Executive Office Building, Room 10235; Washington, DC 20503. Written comments should be received within 30 days of this notice.

Proposed Project

1. *Young People in Alternative Education Settings: Preventing HIV and Other STDs*—New—The National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Division of Adolescent and School Health—The purpose of this request is to obtain OMB clearance to conduct a randomized trial of a curriculum to reduce behaviors related to HIV/STD transmission among 14 to 18 year old students in 30 court and community schools in Northern California. Participants will respond to surveys of attitudes, knowledge, and behavior related to HIV/STD transmission and prevention at baseline and at 6, 12, and 18 month post-tests. Reduction of behaviors among adolescents related to HIV and STD transmission, and reduction of the prevalence of STDs is the focus of at least seven objectives in *Healthy People 2000: Midcourse Review and 1995 Revisions.* There have been few studies assessing the effectiveness of curricula to reduce HIV/STD related risk behaviors in this high-risk adolescent population. Data gathered from this study will provide information about how HIV/STD risk behavior may be effectively reduced among alternative school students. The total annual burden hours are 7,680.

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hrs.)
Alternative school students	1,920	4	1.0

2. *Evaluation of Customer Satisfaction of the Agency for Toxic Substances and Disease Registry (ATSDR) Internet Home Page and Links*—New—ATSDR proposes to conduct consumer satisfaction research around its Internet site. Information on the site focuses on prevention of exposure and adverse human health effects and diminished quality of life associated with exposure to hazardous substances from waste sites, unplanned releases, and other sources of pollution present in the environment. The site is designed to serve the general public, persons at risk for exposure to hazardous substances, and health professionals. This research will ensure that these audiences find the information easy to access, clear, informative, and useful. Specifically, the research will examine whether the information is presented in an appropriate technological format and whether it meets the needs, wants, and preferences of visitors or ''customers'' to the Internet site.

The initial 60 day **Federal Register** Notice was solely for the evaluation of the National Center for HIV, STD, and TB Prevention (NCHSTP) web-site, but after an internal meeting the instrument has been modified for use on the ATSDR web-site. The total annual burden hours are 2,000.

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
Visitors to ATSDR Internet Site	12,000	1	0.1667

3. *Use of Laboratory Information Systems (LIS) to Transmit Infectious Diseases Test Results (HL7 Messages) to Public Health Agencies*—New—Public Health Program Office (PHPPO), Division of Laboratory Systems. CDC proposes to gather data through the use of a mail/telephone survey of all United States vendors of LIS used for recording and processing microbiology data. The use of a mail/telephone-assisted survey instrument will be an efficient, cost-effective approach for performing the data collection. No computerized data collection systems have been developed for this survey because the number of respondents is small. Instead, trained telephone interviewers knowledgeable about LIS and about the specific messages that CDC is interested in transmitting will gather data. The interviewers will have the flexibility to answer technical questions, probe for further information and provide explanations of coding vocabularies, security needs and other issues that may not be readily understood by the LIS vendors.

The data will provide the government, LIS vendors, laboratory practitioners,

committees that make recommendations regarding messaging and other stakeholders with information about the projected costs to vendors and laboratories and about the time frames required for and the barriers to implementation.

CDC will use the survey to gauge the technological readiness and the cost factors affecting secure electronic transmission of infectious disease data to government agencies. These transmissions will act as part of an early warning system leading to more timely response to infectious disease outbreaks. This survey responds to President Clinton's request for the increased use of modern technology to identify and prevent outbreaks of food-borne illness. The total annual burden hours are 121.

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hrs.)
Contact Information Form	56	1	0.1667
Mail Survey	56	2	0.50
Telephone Follow-up	56	2	0.50

Nancy Cheal,

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

[FR Doc. 99–17273 Filed 7–7–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Revised Vessel Sanitation Operations Manual; Public Meeting

The National Center for Environmental Health (NCEH) of the Centers for Disease Control and Prevention (CDC) announces the following meeting.

Name: Discussion of the second draft of the revised Vessel Sanitation Operations Manual—Public meeting between CDC and the cruise ship industry, private sanitation consultants, and other interested parties.

Time and Date: 9 a.m.–4:30 p.m., October 5, 1999; 9 a.m.–4:30 p.m., October 6, 1999; 9 a.m.–12 noon, October 7, 1999.

Place: Auditorium, Port Everglades Administration Building, 1850 Eller Drive, Fort Lauderdale, Florida 33316.

Status: Open to the public, limited by the space available. The meeting room accommodates approximately 100 people.

Purpose: CDC announced its intention to revise the *Vessel Sanitation Operations Manual* in the **Federal Register** of July 9, 1998 (Volume 63, Number 131). Input and comments from the public were requested of and received from the cruise ship industry, private sanitation consultants, and other interested parties, and were discussed in detail at a public meeting held in Fort Lauderdale on April 14–16, 1999. On the basis of comments received, VSP staff have written a second draft of the revised manual and will discuss the revisions at this public meeting.

Matters To Be Discussed: Agenda items will include a thorough discussion of each section of the second draft of the revised operations manual. A copy of the second draft will be available for review by August 6, 1999. To obtain a copy, contact the VSP in Atlanta at the address or phone number below, or go to the VSP Home Page on the Internet at *http://www.cdc.gov/nceh/programs/vsp*.

For a period of 15 days following the meeting, through October 22, 1999, the official record of the meeting will remain open so that additional materials or comments may be submitted to be made part of the record of the meeting. VSP staff will then finalize the revised operations manual and publish the final in the **Federal Register**.

Advanced registration for this important meeting is encouraged. If you plan to attend, please provide your name, title, company name, mailing address, telephone number, facsimile number, and E-mail address to Dorothy Johnson, Management Assistant, facsimile 770/488–4127 or E-mail: *dgj0@cdc.gov*.

Contact Person for More Information: David Forney, Acting Chief, VSP, telephone 770/488–7333 or E-mail: *dlf1@cdc.gov;* or Daniel Harper, Senior Environmental Health Officer, VSP, telephone 770/488–3524, E-mail: *dmh2@cdc.gov;* or write to us at Vessel Sanitation Program, CDC, 4770 Buford Highway, NE, M/S F–16, Atlanta, Georgia 30341–3724.

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 the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: June 30, 1999.

Carolyn J. Russell,

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

[FR Doc. 99–17274 Filed 7–7–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–1110]

Agency Information Collection Activities; Announcement of OMB Approval; Current Good Manufacturing Practice Regulations for Finished Pharmaceuticals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''CGMP Regulations for Finished Pharmaceuticals'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA).

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of April 19, 1999 (64 FR 19180), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0139. The approval expires on June 30, 2002. A copy of the supporting statement for this information collection is available on the Internet at ''http://www.fda.gov/ohrms/dockets''.