Respondents	Number of respondents	Number of responses/respondent	Average burden per response (in hours)
Laboratories	200	1	45/60

Dated: June 8, 2000.

Nancy Cheal,

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

[FR Doc. 00–15114 Filed 6–14–00; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY–46–00]

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 Projects

Possible Estuary-Associated Syndrome (PEAS) Surveillance—Extension—National Center for Environmental Health (NCEH)—In 1997, scientists found a newly identified microorganism, the dinoflagellate Pfiesteria piscicida, in water samples taken from a bay tributary. The presence of large numbers of this organism (a bloom) was purportedly associated with observations of thousands of dead fish as well as with reports of a wide range of adverse human health effects. Reports of this purported association created excessive public concern about exposure to estuarine waters and a general distrust in seafood that prompted a flood of inquiries to public health and environmental quality agencies.

Since 1997, the Centers for Disease Control and Prevention (CDC) has been working with the States of Delaware, Florida, Maryland, North Carolina, South Carolina, and Virginia in a series of meetings, workshops, and conference calls to design, implement, evaluate, and revise surveillance activities to provide a quantitative estimate of the public health burden associated with responding to Pfiesteria-related events, including blooms, fish kills, and people with health complaints. Cooperative agreement funds were awarded to these states to develop a multi-state surveillance system to examine the effects of Pfiesteria blooms upon humans and to expand the scientific knowledge of the human health effects of Pfiesteria. Specifically, the states will quantify the burden of PEAS on their health agencies by enumerating the number of contacts involving public and professional requests for information as well as symptoms involved in self-reporting. In collaboration with the state health departments, NCEH has developed a standardized data collection instrument that the states may use to collect and store the surveillance data. NCEH has requested that the states report specific data elements back at regular intervals so that NCEH can compile the data and issue periodic aggregate reports. CDC/NCEH is requesting a 3-year clearance. The total annual burden hours are 99.

Type of burden	Number of respondents	Number of responses	Average burden/response (in hours)
Information only calls	800	1	5/60
Symptomatic Reports—telephone interview	80	1	25/60

Dated: June 8, 2000.

Nancy Cheal,

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

[FR Doc. 00–15115 Filed 6–14–00; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Announcement Number 00127]

Developing a Model System for the Collection, Analysis, and Dissemination of Data on Genetic Tests, Notice of Availability of Funds; Amendment

A notice announcing the availability of Fiscal Year 2000 funds to fund a cooperative agreement program for Developing a Model System for the Collection, Analysis, and Dissemination of Data on Genetic Tests, which was published in the **Federal Register** on June 8, 2000, (Vol. 65, No. 111, Pages 36446–36448). The notice is amended as follows:

On page 36447, First Column, under Section D. Program Requirements, Item 1(c), change to read: Using experts, identify six to ten DNA-based tests to guide in the finding, collection, and analysis of relevant data (published and unpublished).

On page 36447, Third Column, under Section G. Evaluation Criteria, Item 4(c), change to read: Identify available data for the DNA-based tests, identifying data gaps, and developing common data formats,

Dated: June 9, 2000.

Henry S. Cassell III,

Acting Director, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 00–15116 Filed 6–14–00; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Guide to Community Preventive Services Task Force; Notice of Meeting

Office of the Director, Centers for Disease Control and Prevention (CDC), announces the following meeting:

Name: Guide to Community Preventive Services (GCPS) Task Force Meeting.

Times and Dates: 9 a.m.–5 p.m., June 21, 2000. 8:30 a.m.–3:15 p.m., June 22, 2000.

Place: The Radisson Hotel Atlanta Airport, 5010 Old National Highway, Atlanta, Georgia 30349, telephone (404) 761–4000.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 40 people.

Purpose: The mission of the Task Force is to develop and publish a Guide to Community Preventive Services, which is based on the best available scientific evidence and current expertise regarding essential public health services and what works in the delivery of those services.

Matters to be Discussed: Agenda items include: Final recommendation approvals on the Tobacco Chapter, recommendation approvals for the Oral Health, Physical Activity, Sociocultural Environment, and Sexual Behavior Chapters, an update on dissemination/evaluation plans, updates for the following chapters: Nutrition, Alcohol Use and Misuse, Prevention of Mental Disorders, Cancer, Motor Vehicle Occupant Injury (seat belts and Alcohol Impaired Driving), Violent and Abusive Behavior, and Diabetes, summaries of economic evaluations from the Tobacco Chapter, a general update on economic evaluations from other chapters, and briefings on cross-cutting activities including methods development and development of a cross-cutting chapter.

Agenda items are subject to change as priorities dictate.

Contact Person for Additional Information: Bradford Myers, Deputy Director, Community Guide Section, Division of Prevention Research and Analytic Methods, Epidemiology Program Office, CDC, 4770 Buford Highway, M/S K–73, Atlanta, Georgia 30341, telephone 770/488–8189.

Persons interested in reserving a space for this meeting should call 770/488–8189 by close of business on June 19, 2000.

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 12, 2000.

John Burckhardt,

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

[FR Doc. 00–15245 Filed 6–14–00; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Blood Donor Recruitment Practices; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of meeting.

The Food and Drug Administration (FDA) is announcing the following public workshop: Blood Donor Recruitment Practices. The purpose of the workshop is to gather information on recruiting blood donors and to develop recommendations on the best practices in donor recruitment in the United States.

Date and Time: The public workshop will be held on July 6, 2000, 8:30 a.m. to 5 p.m; and on July 7, 2000, 8:30 a.m. to 2 p.m.

Locations: The July 6, 2000, workshop will be held at the National Institutes of Health, Lister Hill Center, 8600 Rockville Pike, Bldg. 38A, Bethesda, MD. The July 7, 2000, workshop will be held at the same location, and then will move to the Natcher Conference Center, 45 Center Dr., Bldg. 45, for breakout sessions.

Contact: Joseph Wilczek, Center for Biologics Evaluation and Research (HFM–302), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852; 301–827–6129; FAX: 301–827–2843; e-mail: wilczek@cber.fda.gov.

Registration: Early registration is recommended on or before June 23, 2000. Mail or fax registration information (including name, title, firm name, address, telephone, and fax number) to Joseph Wilczek (address above). Registration at the site will be on a space-available basis on the day of the workshop, beginning at 7:30 a.m. There is no registration fee for the workshop. If you need special accommodations due to a disability, please contact Joseph Wilczek at least 7 days in advance.

Agenda: During the first day of the workshop, speakers from the blood bank industry will describe successful recruitment practices. The topics of the presentations will include methods used in successful programs, donor perception, donor retention, telephone recruiting and scheduling, cooperative recruiting in a competitive environment, advertising, education, incentives, and coordinating blood collection with anticipated needs. During the second day, attendees will break into small groups to further discuss key donor recruitment issues. The group discussions will be developed into recommendations of the best practices most likely to increase blood collection to levels sufficient to meet future transfusion needs. At the close of the second day, the attendees will reconvene to share the group recommendations. The information gathered at the workshop may provide the basis for an FDA document on best practices in donor recruitment.

Transcripts: Transcripts of the meeting may be requested in writing from the Freedom of Information Office (HFI–35), Food and Drug Administration, 5600 Fishers Lane, rm. 12A–16, Rockville, MD 20857, approximately 15 working days after the meeting at a cost of 10 cents per page. In addition, the transcript will be placed on the FDA Internet site at www.fda.gov/cber/minutes/workshop-min.htm.

Dated: June 7, 2000.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 00–14904 Filed 6–14–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Circulatory System Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.