[FR Doc. 02–28985 Filed 11–14–02; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY–04–03]

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) 498–1210. 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: Impact of Community Coordinated Response for the Prevention of Intimate Partner Violence: A Random Digital Dial Survey—NEW—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

A random digit dial survey will be conducted with 12,000 male and female adults in the communities of ten experimental sites and ten control sites (600 per site). The survey will determine whether adding resources to a community to develop a coordinated community response to intimate partner violence (IPV), leads to increased knowledge about IPV such as where to go for help and how to assist a victim, child witness and/or perpetrator of IPV. A base survey instrument will be administered along with an addendum from the sites that wish to address other research needs in their experiment and control communities.

While previous surveys such as the National Violence Against Women Survey (1996) have collected information on intimate partner violence, no previous survey has explored the effects of a coordinated community response, enhanced services, and public awareness campaigns between experimental and control sites.

Interviews will be conducted with persons at residential phone numbers selected using random digit dialing. No more than one respondent per household will be selected, and each sample member will complete just one interview. Non-residential numbers are ineligible for the sample and will not be interviewed. Female interviewers will be used and bi-lingual Spanish interviewers will conduct interviews in Spanish to reduce language barriers to participation. The estimated annualized burden is 3813 hours.

Respondents	Number of respondents	Number of responses/ respondent	Avg. burden/ response (in hours)
Pretest	50	1	15/60
Contacted but not eligible or refused	15,000	1	2/60
Core questionnaire (7 sites and comparison communities)*	8,400	1	15/60
Core questionnaire plus addendums*	3,600	1	20/60

Dated: November 6, 2002.

Nancy E. Cheal,

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

[FR Doc. 02–29000 Filed 11–14–02; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Contaminants and Natural Toxicants Subcommittee of the Food 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). The meeting will be open to the public.

Name of Committee: Contaminants and Natural Toxicants Subcommittee of the Food Advisory Committee.

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

Date and Time: The meeting will be held on December 4 and 5, 2002, from 8:30 a.m. to 6 p.m.

Location: The Inn and Conference Center, University of Maryland University College, 3501 University Boulevard East, Adelphi, MD 20783, 301–985–7300.

Contact Person: Henry Kim, Center for Food Safety and Applied Nutrition (HFS–306), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 301–436–2023, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 10564. Please call the Information Line for up-to-date information on this meeting.

Agenda: The subcommittee will discuss FDA's action plan for addressing the issue of acrylamide in food. An agenda for the meeting will be available on the Internet at http://www.cfsan.fda.gov/list.html and at the meeting location on the day of the meeting.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by November 20, 2002. Oral presentations from the public will be scheduled between approximately 4:30 p.m. and 6 p.m. on December 4, 2002, on issues related to acrylamide in food. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before November 20, 2002, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Henry Kim at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: November 7, 2002.

Linda Arey Skladany,

Senior Associate Commissioner for External Relations.

[FR Doc. 02–28941 Filed 11–14–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Vaccines and Related Biological Products 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). The meeting will be open to the public.

Name of Committee: Vaccines and Related Biological Products Advisory Committee.

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

Date and Time: The meeting will be held on December 17, 2002, from 8:30 a.m. to 5 p.m.

Location: Bethesda Marriott Hotel, 5151 Pooks Hill Rd., Bethesda, MD.

Contact Person: Jody G. Sachs or Denise H. Royster, Center for Biologics Evaluation and Research (HFM–71), Food and Drug Administration, 301–827–0314, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12391. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will review and discuss safety and efficacy and proposed indications for the product, FluMist, a cold-adapted, live attenuated, trivalent influenza vaccine for the prevention of influenza sponsored by MedImmune Vaccines, Inc.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by December 1, 2002. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before December 1, 2002, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Jody G. Sachs or Denise H. Royster at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: November 7, 2002.

Linda Arey Skladany,

Senior Associate Commissioner for External Relations.

[FR Doc. 02–28940 Filed 11–14–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Pub. L. 104–13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to OMB under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, call the HRSA Reports Clearance Officer on (301) 443–1129.

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.

Proposed Project: Voluntary Partner Surveys To Implement Executive Order 12862 in the Health Resources and Services Administration—(OMB No. 0915–0212)—Extension

In response to Executive Order 12862, the Health Resources and Services Administration (HRSA) is proposing to conduct voluntary customer surveys of its ''partners'' to assess strengths and weaknesses in program services. A generic approval is being requested from OMB to conduct the partner surveys. HRSA partners are typically State or local governments, health care facilities, health care consortia, health care providers, and researchers.

Partner surveys to be conducted by HRSA might include, for example, mail or telephone surveys of grantees to determine satisfaction with a technical assistance contractor, or in-class evaluation forms completed by providers who receive training from HRSA grantees, to measure satisfaction with the training experience. Results of these surveys will be used to plan and redirect resources and efforts as needed to improve service. Focus groups may also be used to gain partner input into the design of mail and telephone surveys. Focus groups in-class evaluation forms, mail surveys, and telephone surveys are expected to be the preferred methodologies.

A generic approval will permit HRSA to conduct a limited number of partner surveys without a full-scale OMB review of each survey. If generic approval is granted, information on each individual partner survey will not be published in the **Federal Register**.

The estimated response burden is as follows:

Type of survey	Number of respondents	Responses per response	Hours per response	Total hour burden
In-class evaluations	40,000	1	.05	2,000
Mail/Telephone surveys	12,000	1	.25	3,000