mailed to veterinary clinics in the United States. The American Veterinary Medical Association has volunteered to collaborate on the survey and will provide a list of clinics through their membership mailing list. The study objectives are to describe current knowledge, attitudes, and practices of veterinarians regarding zoonotic disease risks and protection of veterinary clinic staff, and to determine what types of national guidelines on infection control practices in veterinary settings are needed. There is no cost to respondents.

| Respondents | Number of respondents | Number of responses/respondent | Average burden/response (in hours) | Total burden (in hours) |
|---|---|---|---|---|
| Written surveys | 5000 | 1 | 20/60 | 1667 |
| Total | | | | 1667 |

Dated: November 7, 2003.

**Gaylon D. Morris,**

*M.P.Aff, Acting Executive Secretariat, Centers for Disease Control and Prevention.*

[FR Doc. 03–28604 Filed 11–14–03; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[60Day–04–06]**

### 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:* Potential Reproductive and Neurological Effects of Exposure to Acrylamide—NEW—The National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

The mission of the National Institute for Occupational Safety and Health (NIOSH) is to promote safety and health at work for all people through research and prevention. Consistent with this mission, NIOSH is undertaking a study of the reproductive and neurobehavioral effects of the occupational exposure to acrylamide. Acrylamide workers and control workers (N = 100 per group) will be recruited from manufacturing, end-user and non-exposed settings. Exposure will be characterized by acrylamide hemoglobin, adduct and urinary metabolite levels, ambient area, personal air, and dermal sampling. Reproductive effects will be evaluated by examining semen quality, sperm DNA integrity, reproductive hormone levels, and prostate specific antigen (PSA) levels.

Neurobehavioral effects will be assessed using sensation-tactile, postual stability, grooved pegboard, and simple reaction time tests. Two questionnaires will be administered on one occasion. Questionnaire information will be collected concurrently to augment test interpretation, adjust for potential confounders and covariates during regression analysis, correlate specific jobs and job activities with exposure measurements, and for validation purposes. Findings from this study will clarify if the adverse reproductive effects observed in animal studies are also present in acrylamide-exposed workers, and if preclinical neurobehavioral deficits are present at acrylamide doses currently considered to be within safe limits.

This study is scheduled for implementation in late 2003 and 2004. There are no costs to respondents.

| Survey questionnaire | Number of respondents | Number of responses/respondent | Average burden/response (in hours) | Total burden (in hrs.) |
|---|---|---|---|---|
| Medical & Reproductive History Questionnaire | 200 | 1 | 13/60 | 43 |
| Occupational History Questionnaire | 200 | 1 | 34/60 | 113 |
| Non-participant Questionnaire | 50 | 1 | 2/60 | 2 |
| Total | | | | 158 |

Dated: November 10, 2003.

**James D. Seligman,**

*Chief Information Officer, Office of the Chief Operations Officer, Centers for Disease Control and Prevention.*

[FR Doc. 03–28605 Filed 11–14–03; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### Advisory Board on Radiation and Worker Health

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 committee meeting:

*Name:* Advisory Board on Radiation and Worker Health (ABRWH), National Institute for Occupational Safety and Health (NIOSH).

*Times and Dates:* 8 a.m.–4 p.m., December 9, 2003, 8 a.m.–5 p.m., December 10, 2003.

*Place:* The Westin Casuarina, 160 East Flamingo Road, Las Vegas, Nevada 89109, telephone 702/836–9775, fax 702/836–9776.

*Status:* Open 8 a.m.–4 p.m., December 9, 2003. Open 8 a.m.–12:30 p.m., December 10, 2003. Closed 2 p.m.–5 p.m., December 10, 2003.

*Background:* The Advisory Board on Radiation and Worker Health (''the Board'') was established under the Energy Employees Occupational Illness Compensation Program Act (EEOICPA) of 2000 to advise the President, through the Secretary of Health and Human Services (HHS), on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Board include providing advice on the development of probability of causation guidelines which have been promulgated by HHS as a final rule, advice on methods of dose reconstruction which have also been promulgated by HHS as a final rule, evaluation of the scientific validity and quality of dose reconstructions conducted by the National Institute for Occupational Safety and Health (NIOSH) for qualified cancer claimants, and advice on the addition of classes of workers to the Special Exposure Cohort.

In December 2000 the President delegated responsibility for funding, staffing, and operating the Board to HHS, which subsequently delegated this authority to the CDC. NIOSH implements this responsibility for CDC. The charter was renewed on August 3, 2003 and the President has completed the appointment of members to the Board to ensure a balanced representation on the Board.

*Purpose:* This board is charged with (a) providing advice to the Secretary, HHS on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS on the scientific validity and quality of dose reconstruction efforts performed for this Program; and (c) upon request by the Secretary, HHS, advise the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class.

*Matters to be Discussed:* The meeting will convene in open session from 8 a.m.–4 p.m. on December 9, 2003 and 8 a.m.–12:30 p.m. on December 10, 2003, to address matters related to NIOSH and Department of Labor updates, an Integrated Module Bioassay Analysis (IMBA) Update, site profile status and roll-out, a Sanford Cohen and Associates brief, reports from the Workgroup on Options for Evaluating Interviews and the Research Issues Workgroup, as well as Board discussion. The remainder of the meeting will proceed in closed session.

The purpose of the closed sessions is to include development, review, and discussion of a proposed Independent Government Cost Estimate (IGCE) for a technical support contract intended to assist the Board in fulfilling its statutory duty to advise the Secretary, HHS regarding dose reconstruction efforts under the EEOICPA. The IGCE will include contract cost estimates, the disclosure of which would adversely impact the Government's negotiating position and strategy in regards to this contract by giving potential bidders an undue advantage in determining the price associated with their bids. The information being discussed will include information of a confidential nature.

This portion of the meeting will be closed to the public in accordance with provisions set forth regarding subject matter considered confidential under the terms of 5 U.S.C. 552b(c)(9)(B), 48 CFR 5.401(b)(1) and (4), and 48 CFR 7.304(d), and the Determination of the Director of the Management Analysis and Services Office, Centers for Disease Control and Prevention, pursuant to Pub. L. 92–463.

A summary of this meeting will be prepared and submitted with 14 days of the close of the meeting.

Agenda items are subject to change as priorities dictate.

**FOR FURTHER INFORMATION CONTACT:** Larry Elliott, Executive Secretary, ABRWH, NIOSH, CDC, 4676 Columbia Parkway, Cincinnati, Ohio 45226, telephone 513/533–6825, fax 513/533–6826.

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: November 10, 2003.

**Betsey Dunaway,**

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

[FR Doc. 03–28600 Filed 11–14–03; 8:45 am]

**BILLING CODE 4163–19–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. 2003N–0502]**

### Agency Information Collection Activities; Proposed Collection; Comment Request; Study to Measure the Compliance of Prescribers With the Contraindication of the Use of Triptans in Migraine Headache Patients With Vascular Disease

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on FDA's burden estimates to distribute an Internet-based questionnaire to measure the compliance of prescribers with the contraindication of the use of triptans in migraine headache patients with vascular disease.

**DATES:** Submit written or electronic comments on the collection of information by January 16, 2004.