- (1) Contracting or subcontracting with any agency of the United States Government and from eligibility for, or involvement in, nonprocurement transactions (e.g., grants and cooperative agreements) of the United States Government as defined in 45 CFR Part 76 (Debarment Regulations);
- (2) Serving in any advisory capacity to the Public Health Service (PHS), including but not limited to service on any PHS advisory committee, board, and/or peer review committee, or as a consultant.

Mr. Hajra agreed to request or cooperate in requesting the retraction or correction of those research publications that have not already been corrected or retracted. He also agreed to notify the relevant editors of the affected review articles that the articles cannot be relied upon.

#### FOR FURTHER INFORMATION CONTACT:

Acting Director, Division of Research Investigations, Office of Research Integrity, 5515 Security Lane, Suite 700, Rockville, MD 20852, (301) 443–5330. Chris B. Pascal.

Acting Director, Office of Research Integrity. [FR Doc. 97–18453 Filed 7–14–97; 8:45 am] BILLING CODE 4160–17–P

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[INFO-97-16]

# 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) 639–7090.

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 for other forms of information technology. Send comments to Wilma Johnson, CDC Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

### **Proposed Projects**

1. Follow-Up Study of Children With Developmental Disabilities—New—In the mid-1980s, 10-year-old children were identified as having one or more of five developmental disabilities: mental retardation, cerebral palsy, epilepsy, hearing impairment, or vision impairment. These children were identified (mainly from special education records in the public schools) in the metro-Atlanta area as part of a study to develop surveillance methods for these conditions in school-age

children. A follow-up study is proposed to trace, locate, and interview these children, who are now in their early twenties, to assess their status with regard to educational attainment, employment, living arrangements, services received, functional limitations, adaptive behavior, social participation, health, and quality of life. Previous studies (published mostly in the mid-1980s) on the post-secondary school experiences of former recipients of special education services were either limited to one type of impairment (e.g., mild mental retardation) or were restricted to a narrow range of outcomes (e.g., employment and education) or did not incorporate a comparison group of persons who were not in special education. The proposed study is a onetime, in-person interview and includes a contemporaneous comparison group of persons who, at age 10 years, were in regular education classes in the same schools as were the persons with developmental disabilities. A base of 1,608 identified children and 650 comparison persons will be used to find a total of 1,600 who will be interviewed. The data generated from this study will be used to estimate the burden of secondary health conditions, limited social participation, and economic disadvantage among young adults with long-standing developmental impairments. This information will be helpful to efforts aimed at the prevention of various secondary problems in this population. The total cost to respondents is \$0.

Respondents	Number of respondents	Number of responses/ respondent	Avg. bur- den/re- sponses (in hrs.)	Total bur- den (in hrs.)
Initial Location Call Contact Call Scheduling Call Telephone Interview	2,258 1,900 1,600 1,600	1 1 1 1	.08 .17 .08 1	180 323 128 1600
Total				2231

Dated: July 9, 1997.

### Wilma G. Johnson,

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

[FR Doc. 97–18498 Filed 7–14–97; 8:45 am] BILLING CODE 4163–18–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### Meeting

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: Task Group Session of the Safety and Occupational Health Study Section, National Institute for Occupational Safety and Health (NIOSH). *Time and Date:* 12:30 p.m.–5:30 p.m., July 30, 1997.

Place: Teleconference originating at the NIOSH Grants Office, 1095 Willowdale Road, Morgantown, West Virginia 26505–2888.

Status: The meeting will be closed in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), title 5 U.S.C., and the Determination of the Associate Director for Management and Operations, CDC, pursuant to Pub. L. 92–463. Application(s) and/or proposal(s) and the discussions could reveal confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the application(s) and/or proposal(s), the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Purpose: The Task Group Session of the Safety and Occupational Health Study Section will review, discuss, and evaluate grant application(s) in response to the Institute's Request for Application Number 722, entitled "Intervention Studies for Construction Safety and Health."

It is the intent of NIOSH to support broad-based research endeavors which will lead to the prevention of workrelated diseases and injuries in the construction industry by designing, implementing, and evaluating measures to reduce occupational hazards. If prevention measures are not currently available, new technologies should be developed for controlling hazardous exposures. Such new technologies must be evaluated to determine that the prevention measures are feasible, even for smaller businesses. Intervention research, of which control technology is a part, examines the utility and impact of new and existing preventive measures in the workplace. It is anticipated that research funded will promote these goals.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Pervis C. Major, Ph.D., Scientific Review Administrator, Office of Extramural Coordination and Special Projects, Office of the Director, NIOSH, 1095 Willowdale Road, Morgantown, West Virginia 26505–2888, telephone 304/ 285–5979.

Dated: July 9, 1997.

### Carolyn J. Russell,

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

[FR Doc. 97–18501 Filed 7–14–97; 8:45 am] BILLING CODE 4163–19–P

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### Meeting

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: Task Group Session of the Safety and Occupational Health Study Section, National Institute for Occupational Safety and Health (NIOSH).

Times and Dates: 8 a.m.-5:30 p.m., August 13, 1997. 8 a.m.-5:30 p.m., August 14, 1997.

*Place:* Hyatt Regency Washington on Capitol Hill, 400 New Jersey Avenue, Washington, DC, 20001.

Status: The meeting will be closed in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), title 5 U.S.C., and the Determination of the Associate Director for Management and Operations, CDC, pursuant to Pub. L. 92–463. Application(s) and/or proposal(s) and the discussions could reveal confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the application(s) and/or proposal(s), the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Purpose: The Task Group Session of the Safety and Occupational Health Study Section will review, discuss, and evaluate grant application(s) in response to the Institute's Request for Application Number 725, entitled "Childhood Agricultural Safety and Health Research."

It is the intent of NIOSH to support broad-based research endeavors which will maximize the safety and health of children and adolescents exposed to agricultural production hazards by expanding the knowledge base regarding etiology; outcomes; intervention strategies; and the effectiveness of commonly utilized educational materials and methods.

Research may address children directly involved in work tasks and/or other children exposed to agricultural production hazards. The funded research projects should cover a variety of types of agricultural production in different geographical regions (e.g. tomato harvesting in California, dairy farms in Wisconsin, and blueberry picking in Maine). It is anticipated that

research funded will promote these goals.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Pervis C. Major, Ph.D., Scientific Review Administrator, Office of Extramural Coordination and Special Projects, Office of the Director, NIOSH, 1095 Willowdale Road, Morgantown, West Virginia 26505–2888, telephone 304/ 285–5979.

Dated: July 9, 1997.

#### Carolyn J. Russell,

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

[FR Doc. 97–18500 Filed 7–14–97; 8:45 am]

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Food and Drug Administration

[Docket No. 97N-0260]

### Agency Information Collection Activities: Proposed Collection; Comment Request

**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 voluntary customer/partner service surveys to implement Executive Order 12862.

**DATES:** Submit written comments on the collection of information by (*insert date 60 days after date of publication in the* **Federal Register**.)

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

### FOR FURTHER INFORMATION CONTACT: Mark L. Pincus, Office of Information Resources Management (HFA–80), Food and Drug Administration, 5600 Fishers