on AHRQ's Web site at *http://www.ahrq.gov/news/foiaindx.htm.*

Dated: August 9, 2002.

Carolyn M. Clancy,

Acting Director.

[FR Doc. 02–20920 Filed 8–6–02; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY–43–02]

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: Evaluation of Worker Notification Program—New—The National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC). The mission of NIOSH is to promote safety and health at work for all people through research and prevention. NIOSH routinely notifies subjects about the results of epidemiologic studies and the implications of the results. The overall purpose of the proposed project is to gain insight into the effectiveness of NIOSH worker notification, in order to improve the quality and usefulness of the Institute's worker notification activities. Researchers from the NIOSH Division of Surveillance, Hazard Evaluations and Field Studies (DSHEFS) propose to provide notified workers with a Reader Response Form as an evaluation instrument for routinely assessing individual letter notification materials sent to them by NIOSH.

The results of this ongoing evaluation activity will be used to refine notification activities by standardizing and streamlining written notification materials, and to develop materials which are more readable, understandable, and informative to notified workers, their families, and other stakeholders. The findings from these evaluations may also allow the NIOSH worker notification program to help alleviate any negative impacts and enhance any positive impacts of risk communications.

The objective of the Reader Response Form, therefore, is to provide a structured reporting form which will capture the recipients' responses concerning the effectiveness of the NIOSH notification efforts and their impact on workers and other stakeholders.

The average number of letter-type notifications is estimated at 8,000 per year. Each form is estimated to take less than 10 minutes to complete. The annual burden for this data collection is 1,333 hours.

Respondents	Number of respondents	Number of responses/ respondent	Avg. burden per response (in hours)
Reader Response Form	8000	1	10/60

Dated: August 9, 2002.

Nancy E. Cheal,

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

[FR Doc. 02–20930 Filed 8–16–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Program Announcement 02158]

University of Georgia Center for Leadership in Education and Applied Research in Mass Destruction Defense; Notice of Award of Funds

A. Purpose

The Centers for Disease Control and Prevention (CDC) announces the award of fiscal year (FY) 2002 funds for a cooperative agreement program for the University of Georgia Center for Leadership in Education and Applied Research in Mass Destruction Defense (CLEARMADD).

The purpose of the program is to facilitate the development of an integrated national system of Centers for Public Health Preparedness focused on improving the capacity of the front-line public health worker to respond to current, new and emerging public health threats. This program addresses the ''Healthy People 2010'' focus areas of Public Health Infrastructure.

B. Eligible Applicant

Assistance is provided only to the University of Georgia Center for Leadership in Education and Applied Research in Mass Destruction Defense. No other applications were solicited. The House of Representatives Conference Report accompanying the Departments of Labor, Health and Human Services, and Education and Related Agencies Appropriation Bill ending September 30, 2002, and For Other Purposes (H.R. 3061, 107th Congress), recognized the University of Georgia's unique qualifications for carrying out the activities specified in this grant (H.R. Rep. 107–342).

C. Funds

Approximately $642,842 is being awarded in FY 2002. The award will begin on or about August 1, 2002 and will be made for a 12-month budget period within a one year project period.

D. Where To Obtain Additional Information

Business management technical assistance may be obtained from: Sharon Robertson, Grants Management Specialist, Acquisition and Assistance, Branch B, Procurement and Grants Office, Centers for Disease Control and Prevention, 2920 Brandywine Road, Room 3000, Atlanta, GA 30341–4146, Telephone number: 770–488–2748, e-mail address: *SRobertson@cdc.gov*.

For program technical assistance, contact: Gail Williams, MPH, CHES, Public Health Practice Program Office, Centers for Disease Control and Prevention (CDC), 4770 Buford Hwy., NE., Mailstop K–38, Atlanta, GA 30341–3717, Telephone: (770) 488–8166.

Dated: August 12, 2002.

Sandra R. Manning,

Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 02–20947 Filed 8–16–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Proposed Projects

Title: National Survey of Child and Adolescent Well-Being.

OMB No.: 0970–0202.

Description: This longitudinal survey provides national estimates on the characteristics related to children and fami8lies who enter the child welfare system. It has collected data from a cohort of 6,100 children who entered the child welfare system as a result of a CPS investigation between October 1999 and April 2001. Data were collected from the children themselves, their caregivers, their teachers, and their caseworkers at baseline, with follow-ups at 12 and 18 months post-baseline. The current request is to pursue a 36-month follow-up, essentially replicating the measure that were used at baseline and at the 18-month follow-up.

Respondents: Children who are clients of the child welfare system, their primary caregivers, caseworkers, and teachers.

Annual Burden Estimates:

Instrument	Responses	Number of responses per respondent	Average burden hours per respondent	Total burden hours
Child interview	5,491	1	1.63	8,950
Caregiver interview	5,491	1	1.50	8,237
Caseworker Interview	2,366	1	0.80	1,893
Caseworker Interview	2,491	1	0.75	1,868

Estimated Total Annual Burden Hours: 20,948.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 650 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Act, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF.

Dated: August 13, 2002.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 02–20936 Filed 8–16–02; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0355]

Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Device Recall Authority

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 including each proposed extension of an existing information collection, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection requirements for medical device recall authority.

DATES: Submit written or electronic comments on the collection of information by October 18, 2002.

ADDRESSES: Submit electronic comments on the collection of information to http://www.accessdata.fda.gov/scripts/oc/dockets/edockethome.cfm. Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary