Traumatic Injuries

Work Environment and Workforce

Emerging Technologies
Indoor Environment
Mixed Exposures
Organization of Work
Special Populations at Risk

Research Tools and Approaches

Cancer Research Methods
Control Technology and Personal Protective Equipment
Exposure Assessment Methods
Health Services Research
Intervention Effectiveness Research
Risk Assessment Methods
Social and Economic Consequences of Workplace Illness and Injury
Surveillance Research Methods

Application Submission and Deadline

The original and two copies of the application PHS Form 5161–1 (Revised 7/92, OMB Number 0937–0189) must be submitted to Ron Van Duyne, Grants Management Officer, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), Mailstop E–13, 255 East Paces Ferry Road, NE., Room 300, Atlanta, GA 30305, on or before July 10, 1996.

1. Deadline: Applications will be considered as meeting the deadline if they are either:

(a) Received on or before the deadline date, or

(b) Sent on or before the deadline date and received in time for submission to the objective review group. (The applicants must request a legibly dated U.S. Postal Service postmark or obtain a legibly dated receipt from a commercial carrier or the U.S. Postal Service. Private metered postmarks will not be acceptable as proof of timely mailing.)

2. Late Applications: Applications that do not meet the criteria in 1.(a) or 1.(b) above are considered late applications. Late applications will not be considered in the current competition and will be returned to the applicants.

Where To Obtain Additional Information

To receive additional written information call (404) 332–4561. You will be asked to leave your name, address, and telephone number and will need to refer to Announcement 647. You will receive a complete program description and information on application procedures and application forms. If you have questions after reviewing the contents of all the documents, business management technical assistance may be obtained from Oppie Byrd, Grants Management Specialist, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 255 East Paces Ferry Road, NE., Room 300, Mailstop E–13, Atlanta, GA 30305, telephone (404) 842–6546, Internet: oxb3@opspgo1.em.cdc.gov, fax (404) 842–6513.

Programmatic technical assistance may be obtained from Greg Kullman, Ph.D., Division of Respiratory Disease Studies, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention (CDC), 1095 Willowdale Road, Morgantown, WV 26505–2888, telephone (304) 285–5711, Internet: gjkl@niords1.em.cdc.gov, fax (304) 285–5796.

There may be delays in mail delivery as well as difficulty in reaching the CDC Atlanta offices during the 1996 Summer Olympics (July 19–August 4). Therefore, CDC suggests the following to get more timely responses to any questions: use Internet/email; follow all instructions in this announcement; and leave messages on the contact person's voice mail.

Please refer to Announcement 647 when requesting information and submitting an application.

Potential applicants may obtain a copy of Healthy People 2000 (Full Report, Stock No. 017–001–00474–0) or Healthy People 2000 (Summary Report, Stock No. 017–001–00473–1) referenced in the ''INTRODUCTION'' Section through the Superintendent of Documents, Government Printing Office, Washington, DC 20402–9325, telephone (202) 512–1800.

Dated: May 22, 1996.

Diane D. Porter,

Acting Director, National Institute for Occupational Safety and Health Centers for Disease Control and Prevention (CDC).

[FR Doc. 96–13472 Filed 5–29–96; 8:45 am]

BILLING CODE 4163–19–P

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Evaluation of Family Support Programs.

OMB Number: New collection.

Description: This study, conducted under a contract to Abt Associates, Inc., responds to the requirement of Subpart 2, Section 435 of OBRA 1993, which directs the Secretary of Health and Human Services to evaluate the effectiveness of family support programs. The information collected will provide descriptive information about family support programs, including detailed information about program operations and variation among programs, and will address the question of the effectiveness of such programs in achieving their goals. The data collected will complement a previous review of existing evaluations of family support programs, and will provide prospective information on eight programs, including information about the operation of such programs and outcomes for families and children who participate. Information will be collected beginning in Fall, 1996, through interviews with parents, children, and teachers of children who are participants in family support programs. Domains of interest include adult and child strengths, home environment, child development, children's school success, development of children's social responsibility, family resources, family social support networks, adoption of healthy lifestyles, community environment, community resources, and community networks.

Respondents: Individuals or households, not-for-profit institutions.

ANNUAL BURDEN ESTIMATE

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Family Interview	1,085	3.1	1	3,340
Child Interview	845	3.4	25	715
Student Interview	245	2	.25	125
Teacher Questionnaire	825	2.8	.17	395

Estimated Total Annual Burden Hours: 4,575.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, Division of Information Resource Management Services, 370 L'Enfant Promenade, SW., Washington, DC. 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by title.

In addition, requests for copies may be made and comments forwarded to the Reports Clearance Office over the Internet by sending message to rsargis@acf.dhhs.gov. Internet message must be submitted as an ASCII file without special characters or encryption.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the function 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 of other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: May 23, 1996.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 96–13526 Filed 5–29–96; 8:45 am]

BILLING CODE 4184–01–M

Food and Drug Administration

[Docket No. 96N–0122]

Agency Information Collection Activities: Proposed Collections; Comment Request; Extension/ Reinstatement

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, Federal agencies are required to publish a notice in the Federal Register concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on reporting requirements contained in existing FDA regulations governing temporary marketing permit applications, State petitions for exemption from preemption, State enforcement notifications, and reference amount petitions.

DATES: Submit written comments on the collections of information by July 29, 1996.

ADDRESSES: Submit written comments on the collections 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: Charity B. Smith, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 16B–19, Rockville, MD 20857, 301–827–1686.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 (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 or reinstatement of an existing collection of information. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c). To comply with this requirement, FDA is publishing notice of the proposed collections of information listed below.

With respect to each of the following collections of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

1. Temporary Marketing Permit Applications (21 CFR 130.17(c) and (i)) (OMB Control Number 0910–0133—Extension)

Section 401 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 341) directs FDA to issue regulations establishing definitions and standards of identity for food ''whenever * * * such action will promote honesty and fair dealing in the interest of consumers.'' Under section 403(g) of the act (21 U.S.C. 343(g)), a food that is subject to a definition and standard of identity prescribed by regulation is misbranded if it does not conform to such definition and standard of identity. Section 130.17 (21 CFR 130.17) provides for the issuance by FDA of temporary marketing permits that enable the food industry to test consumer acceptance and measure the technological and commercial feasibility in interstate commerce of experimental packs of food that deviate from applicable definitions and standards of identity. Section 130.17(c) specifies the information that a firm must submit to FDA to obtain a temporary marketing permit. The information required in a temporary marketing permit application under § 130.17(c) enables the agency to monitor the manufacture, labeling, and distribution of experimental packs of food that deviate from applicable definitions or standards of identity. The information so obtained can be used in support of a petition to establish or amend the applicable definition or standard of identity to provide for the variations. Section 130.17(i) specifies the information that a firm must submit to FDA to obtain an extension of a temporary marketing permit.

FDA estimates the burden of the temporary marketing permit application requirements as follows: