2. Center Administration

Budget to support Center administration to assure: coordination and promotion of academic programs; interdisciplinary interaction; meeting of regional workforce needs; and evaluation of impact.

3. Continuing Education/Outreach Program

Budget to support outreach and continuing education activities to prepare, distribute, and conduct short courses, seminars, and workshops.

4. Hazardous Substance Training Programs

Budget to support the development and presentation of continuing education courses for professionals engaged in the management of hazardous substances.

5. Hazardous Substance Academic Training Programs

Budget to support the development and presentation of specialized academic programs in hazardous substance management.

6. Agricultural Safety and Health Academic Programs

Budget to support the development and presentation of specialized academic programs and continuing education courses in agricultural safety and health.

For Long-Term Training Project grants, the following factors will be considered in determining funding allocations.

Academic Programs

a. Budget to support programs primarily for personnel and other personnel-related costs. Advanced (doctoral and post-doctoral), specialty (master's), and baccalaureate/associate programs will be considered.

b. Budget to support programs based on program quality/technical merit.

c. Budget to support students based on the program level and the number of students supported.

Executive Order 12372 Review

Applications are not subject to review as governed by Executive Order 12372, Intergovernmental Review of Federal Programs.

Public Health System Reporting Requirement

This program is not subject to the Public Health System Reporting Requirements.

Catalog of Federal Domestic Assistance Number

The Catalog of Federal Domestic Assistance Number is 93.263.

Application Submission and Deadline

Applications should be clearly identified as an application for an Occupational Safety and Health Long-Term Training Project Grant or ERC Training Grant. The submission schedule is as follows:

New, Competing Continuation and Supplemental Application Receipt Date: July 1, 1998.

An original and two copies of new, competing continuation and supplemental applications (Form CDC 2.145A ERC or TPG) should be submitted to: Ron Van Duyne (ATTN: Patrick A. Smith), Grants Management Officer, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 255 East Paces Ferry Road, NE., Room 321, Mailstop E–13 Atlanta, GA 30305.

1. *Deadline:* Applications shall 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 independent review group. (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 shall not be acceptable as proof of timely mailing.)

2. *Late Applications:* Applications which 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 applicant.

Non-Competing Continuation Receipt Date: November 16, 1998.

An original and two copies of non-competing continuation applications (Form CDC 2.145B ERC or TPG) should be submitted to: Ron Van Duyne (ATTN: Patrick A. Smith), Grants Management Officer, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 255 East Paces Ferry Road, NE., Room 321, Mailstop E–13 Atlanta, GA 30305.

Where to Obtain Additional Information

To receive an application kit, call 1–888–GRANTS4. You will be asked your name, address, and telephone number and will need to refer to NIOSH Announcement 98045. In addition, this and other CDC announcements are available through the CDC Home page on the Internet. The address for the CDC Home Page is http://www.cdc.gov. If you have questions after reviewing the contents of all the documents, business management technical assistance may be obtained from Patrick A. Smith, Grants Management Specialist, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 255 East Paces Ferry Road, NE., Room 321, Mailstop E–13, Atlanta, GA 30305, telephone (404) 842–6803, or by Internet, phs3@cdc.gov. Programmatic technical assistance may be obtained from John T. Talty, Principal Engineer, Office of Extramural Coordination and Special Projects, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention (CDC), 4676 Columbia Parkway, Mailstop C–7, Cincinnati, OH 45226, telephone (513) 533–8241, or by Internet, jtt2@cdc.gov.

Please refer to Announcement Number 98045 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) through the Superintendent of Documents, Government Printing Office, Washington, DC 20402–9325, telephone (202) 512–1800.

Dated: April 14, 1998.

Diane D. Porter,

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

[FR Doc. 98–10471 Filed 4–20–98; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: ACF Annual Grantee Survey of the Low Income Home Energy Assistance Program (LIHEAP).

OMB No.: 0970–0076.

Description: ACF is required by law to provide Congress with fiscal and caseload estimates of the Grantee LIHEAP Programs. The Secretary is also required to submit a report to Congress each fiscal year for the prior fiscal year.

Respondents: States.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Survey	51	1	3.75	191.25

Estimated Total Annual Burden Hours: 191.25.

Additional Information: Copies of the proposed collection may be obtained by writing to The Administration for Children and Families, 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 60 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 Project, 725 17th Street, NW., Washington, DC 20503, Attn: Ms. Wendy Taylor.

Dated: April 14, 1998.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 98–10444 Filed 4–20–98; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0194]

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, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the voluntary registration of cosmetic product establishments with FDA.

DATES: Submit written comments on the collection of information by June 22, 1998.

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: Margaret R. 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 listed below.

With respect to the following collection 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.

Registration of Cosmetic Product Establishment—21 CFR Part 710 (OMB Control Number 0910–0027—Extension)

Under the Federal Food, Drug, and Cosmetic Act (the act), cosmetic products that are adulterated under section 601 of the act (21 U.S.C. 361) or misbranded under section 602 of the act (21 U.S.C. 362) may not be distributed in interstate commerce. To assist FDA in carrying out its responsibility to regulate cosmetics, FDA requests that establishments that manufacture or package cosmetic products register with the agency on Form FDA 2511 ''Registration of Cosmetic Product Establishment.'' Regulations providing procedures for the voluntary registration of cosmetic product establishments are found in 21 CFR part 710.

Because mandatory registration of cosmetic establishments is not authorized by statute, voluntary registration provides FDA with the best information available about the location, business trading names used, and the type of activity (manufacturing or packaging) of cosmetic product establishments that participate in this program. In addition, the registration information is an essential part of planning onsite inspections to determine the scope and extent of noncompliance with applicable provisions of the act. The registration information is used to estimate the size of the cosmetic industry regulated. Registration is permanent, although FDA requests that firms submit an amended registration on Form FDA 2511 if any of the information originally submitted changes.

FDA uses registration information as input for a computer data base of cosmetic product establishments. This data base is used for mailing lists to distribute regulatory information or to invite firms to participate in workshops on topics they may be interested in.