used to support the expiration date in order to ensure that the expiration date is accurate.

The respondents to this collection of information are domestic and foreign condom manufacturers.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
801.435	45	1	45	96	4,320

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

The number of domestic establishments was estimated by reviewing the FDA data base of registered medical device manufacturers to arrive at 5 domestic and 40 foreign condom manufacturers. Based upon conversations with condom manufacturers, FDA field personnel, and comments received from the public during this collections initial approval, FDA determined the number hours to complete labeling and testing of condoms to be 96 hours per respondent.

Dated: June 15, 2000.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 00–15865 Filed 6–22–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Children's Hospitals Graduate Medical Education (CHGME) Program Conference

On June 19, 2000, the Health Resources and Services Administration (HRSA) published a notice in the **Federal Register** announcing the Children's Hospitals Graduate Medical Education (CHGME) Program (65 FR 37985). Interested parties are invited to attend a briefing conference on July 7, 2000, from 1 p.m. to 3 p.m. EDT in conference room D in the Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857. Parties may also participate in the conference by telephone. To do so, dial: 800–545–4387 or 700–991–1738 (for Federal Government employees), then enter the access code ID# 28353. Telephone participants should call by 12:45 p.m.

The conference is to provide information on the topics contained in the CHGME notice: proposed eligibility criteria, funding factors and methodology, and performance measures. It will include a question and answer period.

For additional information call or write to: F. Lawrence Clare, telephone: (301) 443–7334; Division of Medicine and Dentistry, Bureau of Health Professions, Room 9–A–27, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857; *lclare@hrsa.gov.*

Dated: June 19, 2000.

Claude Earl Fox,

Administrator.

[FR Doc. 00–15901 Filed 6–22–00; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute: Development of Innovative Idiotype Tumor Vaccines

Multiple opportunities are available for collaboration with the National Cancer Institute (NCI), Division of Clinical Sciences, for the pre-clinical and clinical development of protein and/or DNA-based idiotypic vaccines using novel formulations, adjuvants or delivery systems and directed against low-grade and intermediate B-cell lymphomas, mantle cell lymphomas or chronic lymphocytic leukemias (CLL). It is anticipated that because of the magnitude and diversity of these projects the collaboration(s) will take the form of multiple Cooperative Research and Development Agreements (CRADAs). The collaboration(s) may involve any aspect of the therapeutic development of these tumor vaccines from basic scientific inquiry to late stage clinical trials.

AGENCY: National Institutes of Health, PHS, DHHS.

ACTION: Notice of opportunities for Cooperative Research and Development Agreements.

SUMMARY: Pursuant to the Federal Technology Transfer Act of 1986 (FTTA, 15 U.S.C. 3710; Executive Order 12591 of April 10, 1987 as amended by the National Technology Transfer and Advancement Act of 1995), the National Cancer Institute (NCI) of the National Institutes of Health (NIH) of the Public Health Service (PHS) of the Department of Health and Human Services (DHHS) is seeking pharmaceutical or biotechnology companies which can effectively collaborate on the scientific and commercial development of idiotypic tumor vaccines for treatment of low-grade and intermediate B-cell lymphomas, mantle cell lymphoma or chronic lymphocytic leukemia (CLL). The goal of the collaboration(s) will be the development of novel vaccine strategies to elicit an immune response directed against autologous idiotypic surface immunoglobulin derived from these tumors. Any CRADA for further development of this technology that focuses on preclinical or clinical studies of idiotypic vaccines for treatment of the indicated diseases will be considered. The CRADA would have an expected duration of three (3) to five (5) years. The goals of the CRADA will include the rapid publication of research results and timely commercialization of products, diagnostics, and treatments that result from the research. The CRADA Collaborators will have an option to negotiate the terms of an exclusive or nonexclusive commercialization license to subject inventions arising under the CRADA.

ADDRESSES: Proposals and questions about this CRADA opportunity may be addressed to Dr. Thomas M. Stackhouse, Technology Development & Commercialization Branch, National Cancer Institute—Frederick Cancer Research and Development Center, Fairview Center, 1003 West Seventh Street, Room 502, Frederick, MD 20852, Telephone: (301) 846–5222; Facsimile: (301) 846–6820. Scientific Inquiries may be directed to Dr. Larry Kwak, M.D., Ph.D., Senior Investigator, Division of Clinical Sciences, National Cancer Institute, Bldg. 567, Room 205, Frederick, MD 21702–1201, Telephone: (301) 846–1607; Facsimile: (301) 846–6107.

EFFECTIVE DATE: Organizations must submit a proposal summary preferably two pages or less, to NCI within 90 days from date of this publication. Guidelines