Subject, city, state	Effective date
PLANO, TX	
DEFAULT ON HEAL LOAN	
BARAHEMI, MANSOUREH ..... ANAHEIM HILLS, CA	09/20/1999
BARCELO, JAIME V ..... MIAMI, FL	09/20/1999
BARNO, MICHAEL D ..... ORANGEVALE, CA	09/20/1999
GOODWIN, RANDALL J ..... ATWOOD, KS	09/20/1999
JACKSON, STEPHEN C ..... MILLVILLE, NJ	09/20/1999
JOHNSTON, DAVID K ..... HOUSTON, TX	09/20/1999
KAHRS, JEFFREY B ..... TACOMA, WA	09/20/1999
KINCY, GARY W ..... BIRMINGHAM, AL	09/20/1999
KNAPKE, VICKI L ..... INDIANAPOLIS, IN	09/20/1999
KRUPP, MICHAEL D ..... PORTLAND, OR	09/20/1999
MCCOMBS, MARTIN B ..... LONG BEACH, CA	09/20/1999
MCGHEE, ORSEL S III ..... ANAHEIM, CA	09/20/1999
MCKENZIE, LAWRENCE G ..... THIEF RIVER FALLS, MN	09/20/1999
MONGALO, VIRGILIO J ..... MIAMI, FL	09/20/1999
O'BRIEN, DENNIS E ..... LA CENTER, WA	09/20/1999
ORNELAS, MANUEL E ..... MOBILE, AL	09/20/1999
ORTIZ, ERNEST E ..... VALENCIA, CA	09/20/1999
PINSON, JEFFREY R ..... NATALIA, TX	09/20/1999
RAZAVIAN, FAHIMEH ..... MISSION VIEJO, CA	09/20/1999
ROUTLEY, DAVID B ..... BIG RAPIDS, MI	09/20/1999
SEFLA, TODD S ..... PAWTUCKETT, RI	09/20/1999
SOKOL, LOUIS J ..... HOBE SOUND, FL	09/20/1999
THOMPSON, BENJAMIN F ..... MADISON, AL	09/20/1999
WHITTAKER, ROBIN D ..... NASHVILLE, TN	08/25/1999
SETTLEMENT AGREEMENT	
MODERN MEDICAL CENTER, INC ..... MIAMI, FL	09/20/1999

Dated: September 2, 1999.

Joanne Lanahan,

Director, Health Care Administrative Sanctions, Office of Inspector General.

[FR Doc. 99–24429 Filed 9–20–99; 8:45 am]

BILLING CODE 4150–04–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request: Request for Generic Clearance To Conduct Voluntary Customer/Partner Surveys

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Library of Medicine (NLM), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection

Title: Voluntary Customer Satisfaction Surveys. *Type of Information Collection Request:* New. *New and Use of Information Collection:* Executive Order 12962 directs agencies that provide significant services directly to the public to survey customers to determine the kind and quality of services they want and their level of satisfaction with existing services. Additionally, since 1994, the NLM has been a ''Federal Reinvention Laboratory'' with a goal of improving its methods of delivering information to the public. An essential strategy in accomplishing reinvention goals is the ability to periodically receive input and feedback from customers about the design and quality of the services they receive.

The NLM provides significant services directly to the public including health providers, researchers, universities, other federal agencies, state and local governments, and to others through a range of mechanisms, including publications, technical assistance, and web sites. These services are primarily focused on health and medical information dissemination activities. The purpose of this submission is to obtain OMB's generic approval to conduct satisfaction surveys of NLM's customers. The NLM will use the information provided by individuals and institutions to identify strengths and weaknesses in current services and to make improvements where feasible. The ability to periodically survey NLM's customers is essential to continually update and upgrade methods of providing high quality service. *Frequency of Response:* Annually or biennially. *Affected Public:* Individuals or households; businesses or other for profit; state or local governments; Federal agencies; non-profit institutions; small businesses or organizations. *Type of Respondents:* Organizations, medical researchers, physicians and other health care providers, librarians, students, and the general public. Annual reporting burden is an follows:

Title of Survey	Type of survey	Number of respondents	Estimated response time	Burden hours
Evaluation of Clinical Studies Database	Web-based	1,000	.167	167
Visible Human Project—Image Processing Tools	Electronic Mail	1,000	.25	250
PubMed	Web-based	5,000	.0835	418
Entrez	Web-based	2,000	.0835	167
GeneMap	Web-based	2,000	.0835	167
NCBI Web Site	Web-based	2,000	.0835	167
NLM Service Desk Survey	Interactive Voice Response telephone.	400	.0835	33
NLM Onsite Reading Room Use	Exit Interview	500	.167	84
NLM Electronic Mail Customer Survey	Electronic Mail	1,000	.0835	84
MEDLINEplus User Survey	Web-based	500	.0835	59
Survey of Unified Medical Language System (UMLS) Use	Mail Survey	1,000	.5	500
NLM Services Satisfaction Survey	Web-based	2,000	.0835	167
Total				2,163

There are no capital costs to report. There are no operating or maintenance costs to report.

Request for Comments

Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information; (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 those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

FOR FURTHER INFORMATION: To request additional information on the proposed collection of information contact Ronald F. Stewart, National Library of Medicine, Building 38, Room 2N07, 8600 Rockville Pike, Bethesda, MD 20894, or call non-toll free number (301) 496–6491. You may also e-mail your request to: ron__stewart@nlm.nih.gov.

Comments Due Date

Comments regarding this information collection are best assured of having their full effect if received on of before November 22, 1999.

Dated: September 13, 1999.

Donald C. Poppke,

Associate Director for Administrative Management, National Library of Medicine.

[FR Doc. 99–24520 Filed 9–20–99; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Opportunity for a Cooperative Research and Development Agreement (CRADA) To Develop Live Attenuated Dengue Viruses for Use as Vaccines in Humans

AGENCY: National Institutes of Health, Public Health Service, DHHS.

ACTION: Notice.

SUMMARY: The National Institute of Allergy and Infectious Diseases (NIAID) of the National Institutes of Health (NIH) is seeking capability statements from parties interested in entering into a Cooperative Research and Development Agreement (CRADA) on a project to develop live attenuated dengue viruses for use as vaccines to prevent dengue hemorrhagic fever and dengue shock syndrome in humans. This project is part of ongoing vaccine development activities in the Laboratory of Infectious Diseases (LID), Division of Intramural Research, NIAID.

DATES: Only written CRADA capability statements received by the NIAID on or before November 2, 1999 will be considered.

ADDRESSES: Capability statements should be submitted to Dr. Michael R. Mowatt, Office of Technology Development, National Institute of Allergy and Infectious Diseases, National Institutes of Health, 31 Center Drive MSC 2137, Building 31, Room 3B62, Bethesda, MD 20892–2137; Tel: 301/496–2644, Fax: 301/402–7123; Electronic mail: mmowatt@nih.gov.

SUPPLEMENTARY INFORMATION: The CRADA will employ attenuated dengue virus strains (types 1 through 4) developed in LID using recombinant DNA methodologies to (1) identify and characterize the mutations responsible for attenuation, (2) engineer viral strains suitably attenuated for use as human vaccines, and (3) evaluate the attenuated viruses as live vaccines in animals and humans. The Public Health Service (PHS) has filed patent applications both in the U.S. and internationally related to these technologies.

The LID has extensive experience in evaluating the safety, antigenicity, immunogenicity and efficacy of various human viral pathogens and vaccines thereof both in experimental animals and human volunteers. The Collaborator in this endeavor is expected to commit several scientists off-site to support the activities defined by the CRADA Research Plan. These scientists, in collaboration with investigators in the LID, would coordinate the production and release testing of the candidate vaccines, generate monoclonal antibodies needed for manufacture of clinical lots and for their clinical evaluation, and use molecular virologic techniques to generate attenuating mutations suitable for use in live vaccine candidates. In addition, it is expected that the Collaborator will provide funds to supplement LID's research budget for the project and would make a major funding commitment to support the safety, immunogenicity and efficacy studies for candidate vaccines developed under the CRADA.

The capability statement must address, with specificity, each of the following selection criteria: (1) The technical expertise of the Collaborator's Principal Investigator and laboratory group in molecular virology, (2) Ability of Collaborator to manufacture experimental vaccine lots for parenteral administration under Good Manufacturing Practices (GMP) conditions, and (3) Ability to provide adequate and sustained funding to support the requisite vaccine safety and efficacy studies.

Dated: September 13, 1999.

Mark L. Rohrbaugh,

Director, Office of Technology Development, NIAID.

[FR Doc. 99–24516 Filed 9–20–99; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Fogarty International Center; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of a meeting of the Fogarty International Center Advisory Board.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Fogarty International Center Advisory Board, Research Awards Subcommittee Meeting.

Date: September 27, 1999.

Time: 3:00 PM. to 5:00 PM.

Agenda: To review and evaluate grant applications.

Place: Lawton Chiles International House, 16 Center Drive, (Building 16), Bethesda, MD 20892.

Contact Person: Irene W. Edwards, Information Officer, Fogarty International Center, National Institutes of Health, Building 31, Room B2C08, 31 Center Drive MSC 2220, Bethesda, MD 20892, 301–496–2075.

This notice is being published less than 15 days prior to the meeting due to the timing limitation imposed by the review and funding cycle.

(Catalogue of Federal Domestic Assistance Program Nos. 93.106, Minority International