pre- and post-natally, and for other conditions.

The prospective exclusive license will be royalty-bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR 404.7. The prospective exclusive license may be granted unless within sixty (60) days from the date of this published notice, the NIH receives written evidence and argument that establish that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR 404.7.

Applications for a license in the field of use filed in response to this notice will be treated as objections to the grant of the contemplated exclusive license. Comments and objections submitted to this notice will not be made available for public inspection and, to the extent permitted by law, will not be released under the Freedom of Information Act, 5 U.S.C. 552.

Dated: April 6, 2006.

Steven M. Ferguson,

Director, Division of Technology Development and Transfer, Office of Technology Transfer, National Institutes of Health.

[FR Doc. E6–5530 Filed 4–13–06; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Exclusive License: Use of Single Chain T-Cell Receptors To Diagnose and Treat Cancer

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

ACTION: Notice.

SUMMARY: This is notice, in accordance with 35 U.S.C. 209(c)(1) and 37 CFR 404.7(a)(1)(i), that the National Institutes of Health (NIH), Department of Health and Human Services (HHS), is contemplating the grant of an exclusive patent license to practice the inventions embodied in PCT Application Serial No. PCT/US04/29608, filed September 13, 2004 [HHS Ref. No. E–106–2004/0–PCT–01], entitled ''Compositions Comprising T-Cell Receptors and Methods of Use Thereof,'' to Altor Bioscience Corporation, which is located in Miramar, Florida. The patent rights in these inventions have been assigned to the United States of America.

The prospective exclusive license territory may be worldwide and the field of use may be limited to the use of single chain, soluble T-Cell Receptors that recognize NY–ESO, MART–1, and gp100 for diagnosis, prophylaxis, and treatment of melanoma, myeloma, sarcoma, head and neck cancer, bladder cancer, esophageal cancer; lung cancer; stomach cancer; breast cancer; ovarian cancer, colorectal cancer, prostate cancer or liver.

DATES: Only written comments and/or applications for a license which are received by the NIH Office of Technology Transfer on or before June 13, 2006 will be considered.

ADDRESSES: Requests for copies of the patent application, inquiries, comments, and other materials relating to the contemplated exclusive license should be directed to: Michelle A. Booden, PhD., Technology Licensing Specialist, Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852–3804; Telephone: (301) 451–7337; Facsimile: (301) 402–0220; E-mail: *boodenm@mail.nih.gov*.

SUPPLEMENTARY INFORMATION: The technology describes the composition and use of nucleic acid sequences that encode polypeptides capable of forming a T-Cell Receptor (TCR) in a genetically engineered cell. Specifically, these nucleic acid sequences will encode TCRs specific to tumor associated antigens (TAA), gp100, NY–ESO–1, and MART–1. T-Cells engineered with these tumor associated antigen specific TCRs show specific immune responses against TAA expressing cancer cells. Additionally, a method of treating or preventing cancer by administrating the above described TCRs is also disclosed.

The prospective exclusive license will be royalty bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR 404.7. The prospective exclusive license may be granted unless within sixty (60) days from the date of this published notice, the NIH receives written evidence and argument that establishes that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR 404.7.

Applications for a license in the field of use filed in response to this notice will be treated as objections to the grant of the contemplated exclusive license. Comments and objections submitted to this notice will not be made available for public inspection and, to the extent permitted by law, will not be released under the Freedom of Information Act, 5 U.S.C. 552.

Dated: April 6, 2006.

Steven M. Ferguson,

Director, Division of Technology Development and Transfer, Office of Technology Transfer, National Institutes of Health.

[FR Doc. E6–5531 Filed 4–13–06; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Federal Emergency Management Agency, Department of Homeland Security.

ACTION: Notice and request for comments.

SUMMARY: The Federal Emergency Management Agency (FEMA) has submitted the following information collection to the Office of Management and Budget (OMB) for review and clearance in accordance with the requirements of the Paperwork Reduction Act of 1995. The submission describes the nature of the information collection, the categories of respondents, the estimated burden (i.e., the time, effort and resources used by respondents to respond) and cost, and includes the actual data collection instruments FEMA will use.

Title: Application for Community Disaster Loan Program (CDL)/Special Community Disaster Loan Program (SCDL).

OMB Number: 1660–0083.

Abstract: The Application for Community Disaster Loan (CDL) Program and the Special Community Disaster Loan Program (SCDL) provide States, Local and Tribal governments that have suffered substantial loss of tax or other revenues as a result of a major disaster or emergency, the opportunity to obtain financial assistance in order to perform their governmental functions.

Affected Public: State, local, or tribal government.

Number of Respondents: 103.

Estimated Time per Respondent: Burden for this collection is 15.2 hours for the Traditional CDL and 13.4 hours for the Special CDL.

Estimated Total Annual Burden Hours: 1,812.

Frequency of Response: Once.

Comments: Interested persons are invited to submit written comments on the proposed information collection to the Office of Information and Regulatory