

*Name of Committee:* Immunology Integrated Review Group, Vaccines Against Microbial Diseases.

*Date:* June 29–30, 2006.

*Time:* 8:30 a.m. to 5 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

*Contact Person:* Jian Wang, MD, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4198, MSC 7812, Bethesda, MD 20892. (301) 435–2778. [wangjia@csr.nih.gov](mailto:wangjia@csr.nih.gov).

*Name of Committee:* Center for Scientific Review Special Emphasis Panel, Biophysical and Biochemical Sciences Fellowship Panel.

*Date:* June 29–30, 2006.

*Time:* 8:30 a.m. to 5 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Churchill Hotel, 1914 Connecticut Avenue, NW., Washington, DC 20009.

*Contact Person:* James W. Mack, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892. (301) 435–1747. [mackj2@csr.nih.gov](mailto:mackj2@csr.nih.gov).

*Name of Committee:* Center for Scientific Review Special Emphasis Panel, Member Conflict-Behavioral Pharmacology.

*Date:* June 29, 2006.

*Time:* 1 p.m. to 2:30 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892. (Telephone Conference Call).

*Contact Person:* Maribeth Champoux, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3146, MSC 7759, Bethesda, MD 20892. (301) 594–3163. [champoum@csr.nih.gov](mailto:champoum@csr.nih.gov).

*Name of Committee:* Center for Scientific Review Special Emphasis Panel, Technology Development.

*Date:* June 29–30, 2006.

*Time:* 6 p.m. to 4 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Embassy Suites at the Chevy Chase Pavillon, 4300 Military Road, NW., Washington, DC 20015.

*Contact Person:* Sally Ann Amero, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4190, MSC 7849, Bethesda, MD 20892. (301) 435–1159. [ameros@csr.nih.gov](mailto:ameros@csr.nih.gov).

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: May 25, 2006.

**Anna Snouffer,**

*Acting Director, Office of Federal Advisory Committee Policy.*

[FR Doc. 06–5067 Filed 6–1–06; 8:45 am]

**BILLING CODE 4140–01–M**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Prospective Grant of Exclusive License: Human Monoclonal Antibodies, Their Fragments and Derivatives as Biotherapeutics for the Treatment of HIV Infections

**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 license to practice the inventions embodied in: U.S. Provisional Patent Application S/N 60/378,408, filed May 6, 2002 (E–144–2002/0–US–01), PCT Application, PCT/US03/14292, filed May 6, 2003, (E–144–2002/0–PCT–02), converted into 03736557.4 (E–144–2002/0–EP–04) filed in Europe on December 3, 2004, and 2003237187 (E–144–2002/0–AU–05) filed in Australia on November 3, 2004, 10/513,725 (E–144–2002/0–US–03) filed in USA on November 5, 2004, as well as 2,484,930 (E–144–2002/0–CA–06) filed in Canada on November 5, 2004, entitled “Novel broadly cross-reactive HIV neutralizing human monoclonal antibodies selected from Fab phage display libraries using a novel strategy based on alternative antigen panning,” Inventors: Dimiter S. Dimitrov (NCI) and Mei-Yun Zhang (SAIC), to Profectus Biosciences, Inc., having a place of business in Baltimore, Maryland. The patent rights in these inventions have been assigned to the United States of America.

**DATES:** Only written comments and/or application for a license, which are received by the NIH Office of Technology Transfer on or before August 1, 2006 will be considered.

**ADDRESSES:** Requests for a copy of the patent application, inquiries, comments and other materials relating to the contemplated license should be directed to: Sally Hu, Ph.D., M.B.A., Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, MD 20852–3804; E-mail: [hus@od.nih.gov](mailto:hus@od.nih.gov); Telephone: (301) 435–5606; Facsimile: (301) 402–0220.

**SUPPLEMENTARY INFORMATION:** The subject invention (E–144–2002/0) identifies four antibodies, designed 4B1–4, 4B1–10, 4H4, and 5H22 (M12, M14, M16, and M18). These four antibodies were isolated from a human

Fab phage display library using alternating antigen panning (AAP). All four antibodies bind to recombinant HIV envelope glycoproteins (Env) gp<sub>120</sub><sup>2089.6</sup>, gp<sub>120JR-FL</sub> and gp<sub>120IIB</sub> with high affinity. Moreover, 4B1–10 binding to gp 120 or gp 140 is significantly enhanced in the presence of the receptor CD4.

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 60 days from the date of this published Notice, 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.

The field of use may be limited to the development of human monoclonal antibodies for use as a therapeutic or preventative in HIV infection either alone or in combination with other compounds.

Properly filed competing applications for a license filed in response to this notice will be treated as objections to the contemplated license. Comments and objections submitted in response 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: May 25, 2006.

**David R. Sadowski,**

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

[FR Doc. E6–8628 Filed 6–1–06; 8:45 am]

**BILLING CODE 4140–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### First-Generation Guidelines for NCI-Supported Biorepositories

**AGENCY:** National Institutes of Health (NIH), National Cancer Institute (NCI), HHS.

**ACTION:** Notice.

**SUMMARY:** The public comment period for the First Generation Guidelines for NCI-Supported Biorepositories (**Federal Register**, Vol. 71, Number 82, Page 25814, April 28, 2006) will be extended an additional 30 days beyond publication of this notice.

**DATES:** *Effective Date:* July 3, 2006.

**FOR FURTHER INFORMATION CONTACT:** Implementation assistance and inquiries should be directed to senior staff of the