Contact Person: Chhanda L. Ganguly, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4120, MSC 7802, Bethesda, MD 20892, (301) 435– 1739, gangulyc@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Neurobiology of HPA Axis Hormones and

Transmitters ZRG1 IFCN D (02). Date: November 16, 2004.

*Time:* 2 p.m. to 4 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: Gamil C Debbas, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5170, MSC 7844, Bethesda, MD 20892, (301) 435– 1018, debbasg@csr.nih.gov.

*Name of Committee:* Center for Scientific Review Special Emphasis Panel, Platelet Microparticles in Hemostasis.

*Date:* November 16, 2004.

*Time:* 2 p.m. to 3 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:* Delia Tang, MD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4126, MSC 7802, Bethesda, MD 20892, (301) 435–2506, *tangd@csr.nih.gov.* 

*Name of Committee:* Center for Scientific Review Special Emphasis Panel, ZRG1 MOSS–G 01S: Musculoskeletal Tissue Engineering.

Date: November 16-17, 2004.

*Time:* 4 p.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

<sup>^</sup>*Place:* Jurys Doyle Hotel, 1500 New Hampshire Avenue, NW., Washington, DC 20036.

*Contact Person:* Jean D. Sipe, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4106, MSC 7814, Bethesda, MD 20892, (301) 435– 1743, *sipej@csr.nih.gov.* 

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

Date: November 16, 2004.

*Time:* 12 p.m. to 4 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* One Washington Circle Hotel, One Washington Circle, Washington, DC 20037.

Contact Person: Stephen M. Nigida, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4212, MSC 7812, Bethesda, MD 20892, (301) 435– 1222, nigidas@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: October 27, 2004. LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy. [FR Doc. 04–24484 Filed 11–2–04; 8:45 am] BILLING CODE 4140–01–M

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

## National Institutes of Health

# Statement of Organization, Functions, and Delegations of Authority

Part N, National Institutes of Health, of the Statement of Organization, Functions, and Delegations of Authority for the Department of Health and Human Services (40 FR 22859, May 27, 1975, as amended most recently at 68 FR 10743, March 6, 2003, and redesignated from Part HN as Part N at 60 FR 56605, November 9, 1995), is amended as set forth below to reflect the elevation of the NIH Ethics Office from the Office of Management, Office of the Director, NIH, to the Office of the Director, NIH.

Section N–B, Organization and Functions, under the heading Office of the Director (NA, formerly HNA) is amended as follows:

Under the heading Office of the Director (NA, formerly HNA), insert the following:

NIH Ethics Office (NAT, formerly HNAT). (1) Develops and administers the NIH policies for implementing the government-wide conflict of interest statutes and regulations, the HHS supplemental conflict of interest regulations, and HHS and NIH policies; (2) provides ethics policy guidance, training, and advice to: (a) The ICs ethics staffs, (b) Office of the Director, NIH, staff, and (c) employees whose Deputy Ethics Counselor (DEC) is the NIH DEC; and (3) coordinates the NIH response to requests from Congress, the Inspector General, DHHS and/or the Office of Government Ethics.

#### **Delegations of Authority Statement**

All delegations and redelegations of authority to officers and employees of NIH that were in effect immediately prior to the effective date of this establishment and are consistent with this amendment shall continue in effect, pending further redelegation.

Dated: September 14, 2004.

#### Elias A. Zerhouni,

Director, National Institutes of Health. [FR Doc. 04–24480 Filed 11–2–04; 8:45 am] BILLING CODE 4140–01–P

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

### **Public Health Service**

National Toxicology Program; National Institute of Environmental Health Sciences (NIEHS); National Institutes of Health (NIH) Notice of an Expert Panel Meeting To Assess the Current Validation Status of In Vitro Testing Methods for Identifying Potential Ocular Irritants; Request for Comments

#### Summary

Notice is hereby given of a meeting sponsored by the NIEHS and the National Toxicology Program (NTP), and organized by the NTP Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM) in collaboration with the Interagency Coordinating Committee on the Validation of Alternative Methods (ICCVAM). At this meeting, an expert panel ("Panel") will assess the current validation status and develop recommendations for further validation of in vitro test methods proposed for identifying substances that may cause serious eye damage. The meeting will take place on January 11-12, 2005, from 8:30 a.m. to 5 p.m., at the National Institutes of Health (NIH), Natcher Center, Bethesda, MD. The meeting is open to the public with attendance limited only by the space available.

#### Evaluation of In Vitro Ocular Test Methods Background

In August 2003, the Scientific Advisory Committee on Alternative Toxicological Methods (SACATM) recommended that ICCVAM review the validation status of screening test methods that could be used to identify severe and irreversible ocular effects and carry out appropriate reviews of these test methods. In October 2003, the U.S. Environmental Protection Agency nominated several ocular-related activities to ICCVAM including evaluation of the validation status of four in vitro ocular toxicity test methods for screening for severe/irreversible ocular effects: (1) The Bovine Corneal Opacity and Permeability (BCOP) test; (2) the Hen's Egg Test—Chorion Allantoic Membrane (HET–CAM); (3) the Isolated Rabbit Eye (IRE) test; and (4) the Isolated Chicken Eye (ICE) test. ICCVAM endorsed the review of the methods as a high priority and recommended that Background Review Documents be developed for each method by NICEATM in collaboration with the ICCVAM Ocular Toxicity Working Group. ICCVAM also