*Time:* 8 a.m. to 1 p.m. *Agenda:* To review and evaluate grant

applications. *Place* Crystal Gateway Marriott, 1700 Jefferson Davis Highway, Arlington, VA 22202.

Contact Person: Keith A. Mintzer, PhD, Scientific Review Officer, Review Branch/ DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7186, Bethesda, MD 20892–7924, 301–435–0280, mintzerk@nhlbi.nih.gov.

Name of Committee: National Heart, Lung, and Blood Institute Special Emphasis Panel, Center for Gene Transfer.

Date:

*Date:* June 20, 2011.

Time: 12 p.m. to 3 p.m.

*Agenca:* To review and evaluate grant applications.

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

Contact Person: William J Johnson, PhD, Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7178, Bethesda, MD 20892–7924, 301–435– 0725, johnsonwj@nhlbi.nih.gov.q2 (Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: May 17, 2011.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy. [FR Doc. 2011–12630 Filed 5–20–11; 8:45 am]

BILLING CODE 4140-01-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

# National Center for Complementary & Alternative Medicine; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meeting.

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:* National Center for Complementary and Alternative Medicine

Special Emphasis Panel; NCCAM Education Panel.

*Date:* June 23–24, 2011.

*Time:* 8 a.m. to 12 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Marriott Courtyard Gaithersburg Washingtonian Ctr, 204 Boardwalk Place, Gaithersburg, MD 20878.

Contact Person: Peter Kozel, PhD, Scientific Review Officer, NCCAM, 6707 Democracy Boulevard, SUITE 401, Bethesda, MD 20892–5475, 301–496–8004, kozelp@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.213, Research and Training in Complementary and Alternative Medicine, National Institutes of Health, HHS)

Dated: May 17, 2011.

### Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2011–12640 Filed 5–20–11; 8:45 am] BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

# National Institute of General Medical Sciences; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meeting.

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:* National Institute of General Medical Sciences Initial Review Group, Biomedical Research and Research Training Review Subcommittee B.

Date: June 17, 2011.

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

*Agenda:* To review and evaluate grant applications.

*Place:* Hilton Washington DC–Silver Spring, 8727 Colesville Road, Silver Spring, MD 20910.

*Contact Person:* Arthur L. Zachary, PhD, Scientific Review Officer, Office of Scientific Review, National Institute of General Medical Sciences, National Institutes of Health, Natcher Building, Room 3AN–18, Bethesda, MD 20892, 301–594–2886, zacharya@nigms.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.375, Minority Biomedical Research Support; 93.821, Cell Biology and Biophysics Research; 93.859, Pharmacology, Physiology, and Biological Chemistry Research; 93.862, Genetics and Developmental Biology Research; 93.88, Minority Access to Research Careers; 93.96, Special Minority Initiatives, National Institutes of Health, HHS)

Dated: May 17, 2011.

#### Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy. [FR Doc. 2011–12639 Filed 5–20–11; 8:45 am] BILLING CODE 4140–01–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

# Call for Participation in Pillbox Patient-Safety Initiative

### ACTION: Notice.

SUMMARY: The National Library of Medicine (NLM) invites the participation of manufacturers, including repackagers, and private label distributors of solid oral dosage form medications in the development of Pillbox, a publicly accessible online repository of digital images and descriptive information for solid oral dosage form medications. This project seeks to promote utilization of the SPLIMAGE element of the Food and Drug Administration (FDA) Structured Product Label (SPL) through development and use of imaging standards and methodologies. Through this Call for Participation, NLM seeks to evaluate the photography methodology and procedures it has developed for creating standardized high-resolution images of solid oral dosage form medications that are appropriate for inclusion in the SPL. Participating organizations will be invited to submit samples of their solid oral dosage form medications to NLM for imaging. Resulting image files will be provided to participants, who may choose to voluntarily include them in their subsequent SPL submissions to FDA. Image files that are voluntarily submitted to FDA as part of an SPL listing submission will be included in the publicly accessible, production version of Pillbox. This initiative is an important element of ongoing efforts to enhance patient safety, reduce adverse drug events, and improve the quality and availability of drug information.

**SUPPLEMENTARY INFORMATION:** NLM has established Pillbox, an initiative to enhance patient safety, by making available via a publicly accessible resource (*http://pillbox.nlm.nih.gov*) digital images and descriptive data of