full Committee meeting will convene at 1 p.m. ACOT will hear presentations on the Organ Procurement and Transplantation Network (OPTN) Kidney and Liver Allocation Policies, Reports on Living Donor Studies, Report on Donor Potential Study, Kidney Paired Donor Exchange Pilot Project, Report and Conclusions of the Expert Panel on Circulatory Death Criteria, the National Kidney Foundation’s “End the Wait” Initiative, and Summary of Breakthrough Collaborative Issues/Donation and Transplantation Community of Practice. Agenda items are subject to change as priorities indicate.

After the presentations and Committee discussions, members of the public will have an opportunity to provide comments. Because of the Committee’s full agenda and the timeframe in which to cover the agenda topics, public comments will be limited. All public comments will be included in the record of the ACOT meeting. Meeting summary notes will be made available on the Department’s donation Web site at http://www.organdonor.gov/acot.html.

The draft meeting agenda is available on the Department’s donation Web site at http://www.organdonor.gov/acot.html and at https://www.team-psa.com/dot/summer2010/ACOT/splash.asp. Registration can be completed electronically at https://www.team-psa.com/dot/summer2010/ACOT/splash.asp or submitted by facsimile to HRM/Professional and Scientific Associates (PSA), the logistical support contractor for the meeting, at fax number (703) 234–1701 ATTN: Brittany Carey. Individuals without access to the Internet who wish to register may call Brittany Carey with HRM/PSA at (703) 889–9033.

FOR FURTHER INFORMATION CONTACT:
Patricia Stroup, Executive Secretary, Healthcare Systems Bureau, Health Resources and Services Administration, 5600 Fishers Lane, Room 12–105, Rockville, Maryland 20857; telephone (301) 443–1127.

Dated: July 2, 2010.

Sahira Rafiullah,
Director, Office of Information Coordination.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Neurological Disorders and Stroke; 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 Neurological Disorders and Stroke Special Emphasis Panel; Translational Stroke

Place: National Institutes of Health, 6001 Executive Boulevard, Rockville, MD 20852 (Telephone Conference Call).

Contact Person: Shanta Rajaram, PhD, Scientific Review Administrator, Scientific Review Branch, Division of Extramural Research, NINDS/NIH/DHHS/Neuroscience Center, 6001 Executive Blvd., Suite 3208. Ms: 9529, Bethesda, MD 20852, 301–435–6033, rajarams@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.853, Clinical Research Related to Neurological Disorders; 93.854, Biological Basis Research in the Neurosciences, National Institutes of Health, HHS)

Dated: July 1, 2010.

Jennifer Spaeth,
Director, Office of Federal Advisory Committee Policy.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; 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 Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; Autoimmune Microbiome in Diabetes Ancillary Studies

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

Contact Person: Robert C Elliott, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3130, MSC 7850, Bethesda, MD 20892, 301–435–3009, elliottro@nih.gov.

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel;
DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Accreditation and Approval of Intertek USA, Inc., as a Commercial Gauger and Laboratory


ACTION: Notice of accreditation and approval of Intertek USA, Inc., as a commercial gauger and laboratory.

SUMMARY: Notice is hereby given that, pursuant to 19 CFR 151.12 and 19 CFR 151.13, Intertek USA, Inc., 1000 Port Carteret Drive Building C, Carteret, NJ 07008, has been approved to gauge and accredited to test petroleum and petroleum products for customs purposes, in accordance with the provisions of 19 CFR 151.12 and 19 CFR 151.13. Anyone wishing to employ this entity to conduct laboratory analyses and gauger services should request and receive written assurances from the entity that it is accredited or approved by the U.S. Customs and Border Protection to conduct the specific test or gauger service requested. Alternatively, inquiries regarding the specific test or gauger service this entity is accredited or approved to perform may be directed to the U.S. Customs and Border Protection by calling (202) 344-1060.

The inquiry may also be sent to cbp.labhq@dhs.gov. Please reference the Web site listed below for a complete listing of CBP approved gaugers and accredited laboratories: http://cbp.gov/xp/cgov/import/operations_support/labsscientific_svcs/commercial_gaugers/.

DATES: The accreditation and approval of Intertek USA, Inc., as commercial gauger and laboratory became effective on April 16, 2010. The next triennial inspection date will be scheduled for April 2013.