DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; Notice of Closed Meetings

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 meetings.

The meetings 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 contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Heart, Lung, and Blood Institute; Special Emphasis Panel, Science Moving TowArds Research Translation and Therapy (SMARTT), Pharmacology and Toxicology, Program Extension.

Date: August 25, 2015.
Time: 3:00 p.m. to 3:30 p.m.
Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, 6701 Rockledge Drive, Room 7185, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Kristen Page, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7185, Bethesda, MD 20892, kristen.page@nih.gov.

Name of Committee: National Heart, Lung, and Blood Institute, Special Emphasis Panel, Science Moving TowArds Research Translation and Therapy (SMARTT), Biologics Production Facility (PF), Program Extension.

Date: August 25, 2015.
Time: 3:30 p.m. to 4:00 p.m.
Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, 6701 Rockledge Drive, Room 7185, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Kristen Page, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7185, Bethesda, MD 20892, kristen.page@nih.gov.

Name of Committee: National Heart, Lung, and Blood Institute, Special Emphasis Panel, Science Moving TowArds Research Translation and Therapy (SMARTT), Non-Biological Production Facility (PF), Program Extension.

Date: August 25, 2015.
Time: 4:00 p.m. to 4:30 p.m.
Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, 6701 Rockledge Drive, Room 7185, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Kristen Page, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7185, Bethesda, MD 20892, kristen.page@nih.gov.

Name of Committee: National Heart, Lung, and Blood Institute, Special Emphasis Panel, Science Moving TowArds Research Translation and Therapy (SMARTT), Pharmacology and Toxicology, Program Extension.

Date: August 25, 2015.
Time: 4:30 p.m. to 5:00 p.m.
Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, 6701 Rockledge Drive, Room 7185, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Kristen Page, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7185, Bethesda, MD 20892, kristen.page@nih.gov.

(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: July 28, 2015.
Michelle Trout, Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015–18874 Filed 7–31–15; 8:45 am]
specimen validity tests on urine specimens for federal agencies.

To become certified, an applicant laboratory or IITF must undergo three rounds of performance testing plus an on-site inspection. To maintain that certification, a laboratory or IITF must participate in a quarterly performance testing program plus undergo periodic, on-site inspections.

Laboratories and IITFs in the applicant stage of certification are not to be considered as meeting the minimum requirements described in the HHS Mandatory Guidelines. A HHS-certified laboratory or IITF must have its letter of certification from HHS/SAMHSA (formerly: HHS/NIDA), which attests that it has met minimum standards.

In accordance with the Mandatory Guidelines dated November 25, 2008 (73 FR 71858), the following HHS-certified laboratories and IITFs meet the minimum standards to conduct drug and specimen validation tests on urine specimens:

**HHS-Certified Instrumented Initial Testing Facilities**

- Dynacare, 6628 50th Street NW, Edmonton, AB Canada T6B 2N7, 780–784–1190. (Formerly: Dynacare Medical Laboratories).
- Alere Toxicology Services, 1111 Newton St., Gretna, LA 70053, 504–361–8999/800–433–3823, (Formerly: Kroll Laboratory Specialists, Inc., Laboratory Specialists, Inc.).
- Baptist Medical Center-Toxicology Laboratory, 11401 I–30, Little Rock, AR 72209–7056, 501–202–2783, (Formerly: Forensic Toxicology Laboratory Baptist Medical Center).
- DrugScan, Inc., 200 Precision Road, Suite 200, Horsham, PA 19044, 800–239–2283.
- Fortes Laboratories, Inc., 25749 SW Canyon Creek Road, Suite 600, Wilsonville, OR 97070, 503–486–1023.
- Laboratory Corporation of America Holdings, 7207 N. Gessner Road, Houston, TX 77040, 713–856–8288/800–200–2387.
- Laboratory Corporation of America Holdings, 1220 Main Street, Southaven, MS 38671, 866–827–8042/800–233–6339, (Formerly: LabCorp Occupational Testing Services, Inc.; MedExpress/National Laboratory Center).
- LabOne, Inc. d/b/a Quest Diagnostics, 10101 Renner Blvd., Lenexa, KS 66219, 913–888–3927/800–873–8845, (Formerly: Quest Diagnostics Incorporated; LabOne, Inc.; Center for Laboratory Services, a Division of LabOne, Inc.).
- Minneapolis Veterans Affairs Medical Center-Forensic Toxicology Laboratory, 1 Veterans Drive, Minneapolis, MN 55417, 612–725–2088.
- One Source Toxicology Laboratory, Inc., 1213 Genoa-Red Bluff, Pasadena, TX 77504, 888–747–3774, (Formerly: University of Texas Medical Branch, Clinical Chemistry Division; UTMB Pathology-Toxicology Laboratory).
- Pacific Toxicology Laboratories, 9348 DeSoto Ave., Chatsworth, CA 91311, 800–328–6942, (Formerly: Continena Hospital Airport Toxicology Laboratory).


Quest Diagnostics Incorporated, 8401 Fallbrook Ave., West Hills, CA 91304, 818–737–6370, (Formerly: SmithKline Beecham Clinical Laboratories).

Redwood Toxicology Laboratory, 3700650 Westwind Blvd., Santa Rosa, CA 95403, 800–255–2159.


STERLING Reference Laboratories, 2617 East L Street, Tacoma, Washington 98421, 800–442–0438.

US Army Forensic Toxicology Drug Testing Laboratory, 2490 Wilson St., Fort George G. Meade, MD 20755–5235, 301–677–7085.

* The Standards Council of Canada (SCC) voted to end its Laboratory Accreditation Program for Substance Abuse (LASA) effective May 12, 1998. Laboratories certified through that program were accredited to conduct forensic urine drug testing as required by U.S. Department of Transportation (DOT) regulations. As of that date, the certification of those accredited Canadian laboratories will continue under DOT authority. The responsibility for conducting quarterly performance testing plus periodic on-site inspections of those LASA-accredited laboratories was transferred to the U.S. HHS, with the HHS’ NLCP contractor continuing to have an active role in the performance testing and laboratory inspection processes. Other Canadian laboratories wishing to be considered for the NLCP may apply directly to the NLCP contractor just as U.S. laboratories do.

Upon finding a Canadian laboratory to be qualified, HHS will recommend that DOT certify the laboratory (Federal Register, July 16, 1996) as meeting the minimum standards of the Mandatory Guidelines published in the Federal Register on April 30, 1990 (75 FR 22809). After receiving DOT certification, the laboratory will be
DEPARTMENT OF THE INTERIOR
Fish and Wildlife Service
Endangered Species; Marine Mammals; Receipt of Applications for Permit
AGENCY: Fish and Wildlife Service, Interior.
ACTION: Notice of receipt of applications for permit.
SUMMARY: We, the U.S. Fish and Wildlife Service, invite the public to comment on the following applications to conduct certain activities with endangered species, marine mammals, or both. With some exceptions, the Endangered Species Act (ESA) and Marine Mammal Protection Act (MMPA) prohibit activities with listed species unless Federal authorization is acquired that allows such activities.
DATES: We must receive comments or requests for documents on or before September 2, 2015. We must receive requests for marine mammal permit public hearings, in writing, at the address shown in the ADDRESSES section by September 2, 2015.
ADDRESSES: Brenda Tapia, U.S. Fish and Wildlife Service, Division of Management Authority, Branch of Permits, MS: IA, 5275 Leesburg Pike, Falls Church, VA 22041; fax (703) 358–2281; or email DMAFR@fws.gov.
FOR FURTHER INFORMATION CONTACT: Brenda Tapia, (703) 358–2104 (telephone); (703) 358–2281 (fax); DMAFR@fws.gov (email).
SUPPLEMENTARY INFORMATION:
I. Public Comment Procedures
A. How do I request copies of applications or comment on submitted applications?
Send your request for copies of applications or comments and materials concerning any of the applications to the contact listed under ADDRESSES. Please include the Federal Register notice publication date, the PRT-number, and the name of the applicant in your request or submission. We will not consider requests or comments sent to an email or address not listed under ADDRESSES. If you provide an email address in your request for copies of applications, we will attempt to respond to your request electronically. Please make your requests or comments as specific as possible. Please confine your comments to issues for which we seek comments in this notice, and explain the basis for your comments. Include sufficient information with your comments to allow us to authenticate any scientific or commercial data you include. The comments and recommendations that will be most useful and likely to influence agency decisions are: (1) Those supported by quantitative information or studies; and (2) Those that include citations to, and analyses of, the applicable laws and regulations. We will not consider or include in our administrative record comments we receive after the close of the comment period (see DATES) or comments delivered to an address other than those listed above (see ADDRESSES).
B. May I review comments submitted by others?
Comments, including names and street addresses of respondents, will be available for public review at the street address listed under ADDRESSES. The public may review documents and other information applicants have sent in support of the application unless our allowing viewing would violate the Privacy Act or Freedom of Information Act. Before including your address, phone number, email address, or other personal identifying information in your comment, you should be aware that your entire comment—including your personal identifying information—may be made publicly available at any time. While you can ask us in your comment to withhold your personal identifying information from public review, we cannot guarantee that we will be able to do so.
II. Background
To help us carry out our conservation responsibilities for affected species, and in consideration of section 10(a)(1)(A) of the Endangered Species Act of 1973, as amended (16 U.S.C. 1531 et seq.), and the Marine Mammal Protection Act of 1972, as amended (16 U.S.C. 1361 et seq.), along with Executive Order 13576, “Delivering an Efficient, Effective, and Accountable Government,” and the President’s Memorandum for the Heads of Executive Departments and Agencies of January 22, 2009—Transparency and Open Government (74 FR 4685; January 26, 2009), which call on all Federal agencies to promote openness and transparency in Government by disclosing information to the public, we invite public comment on these permit applications before final action is taken. Under the MMPA, you may request a hearing on any MMPA application received. If you request a hearing, give specific reasons why a hearing would be appropriate. The holding of such a hearing is at the discretion of the Service Director.
III. Permit Applications
A. Endangered Species
Applicant: KJC Holdings LP, Lohn, TX; PRT–200211
The applicant requests a permit for cull and take of excess barasingha (Rucervus duvaucellii) for the purpose of enhancement of the survival of the species. This notification covers activities to be conducted by the applicant over a 5-year period.
Applicant: Fleming Creative Concept LLC, Scottsdale, AZ; PRT–65772B
The applicant requests a permit for interstate transport of 10 jackass penguins (Spheniscus demersus) from the Six Flags Discovery Kingdom, Vallejo, California to Miami Seaquarium, Miami, Florida, for the purpose of enhancement of the survival of the species. This notification covers activities to be conducted by the applicant over a 5-year period.
Applicant: Earth Promise, Glen Rose, TX; PRT–59071B
The applicant requests a permit to export two scimitar-horned oryx (Oryx dammah) and two addaxs (Addax nasomaculatus) born in captivity to African Lion Safari, Ontario, Canada, for the purpose of enhancement of the survival of the species.
Applicant: Barefoot Zoological, dba Alligator Adventure, North Myrtle Beach, SC; PRT–141742
The applicant requests renewal of a captive-bred wildlife registration under 50 CFR 17.21(g) for the following species to enhance species propagation or survival: Ring-tailed lemur (Lemur catta), red ruffed lemur (Varecia rubra), black and white ruffed lemur (Varecia variegata), Chinese alligator (Alligator sinensis), Nile crocodile (Crocodylus niloticus), saltwater crocodile (Crocodylus porosus), common caiman (Caiman crocodilus crocodilus), brown caiman (Caiman crocodilus fuscus), Cuban crocodile (Crocodylus rhombifer), Siamese crocodile (Crocodylus siamensis), Yacare (Caiman yacare), dwarf crocodile (Osteolaemus