irrelevant. The right to contest records is limited to information which is incomplete, irrelevant, incorrect, or untimely (obsolete).

RECORD SOURCE CATEGORIES:
Inventors and other collaborating persons, grantees, fellowship recipients and contractors; other Federal agencies; scientific experts from non-Government organizations; contract patent counsel and their employees and foreign contract personnel; United States and foreign patent offices; prospective licensees; HHS Technology Development Coordinators, Internet and commercial databases, and third parties whom HHS contacts to determine individual invention ownership or Government ownership.

SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:
None.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health
Notice of Listing of Members of the National Institutes of Health's Senior Executive Service Performance Review Board (PRB)

The National Institutes of Health (NIH) announces the persons who will serve on the National Institutes of Health’s Senior Executive Service Performance Review Board. This action is being taken in accordance with Title 5, U.S.C., Section 3141(c)(4), which requires that members of performance review boards be appointed in a manner to ensure consistency, stability, and objectivity in performance appraisals and requires that notice of the appointment of an individual to serve as a member be published in the Federal Register. The following persons will serve on the NIH Performance Review Board, which oversees the evaluation of performance appraisals of NIH Senior Executive Service (SES) members: Ms. Colleen Barros (Chair); Dr. Norka Ruiz Bravo; Dr. Michael Gottesman; Dr. John Hallenbeck; Ms. Lynn Hellinger; Dr. Raynard Kington; Dr. Lore Anne McNicol.

For further information about the NIH Performance Review Board, contact the Office of Human Resources, Workforce Relations Division, National Institutes of Health, Building 31, Room B3C07, Bethesda, Maryland 20892, telephone 301–402–9203 (not a toll-free number). Dated: August 1, 2006.

Elias A. Zerhouni,
Director, National Institutes of Health.
[FR Doc. E6–13209 Filed 8–11–06; 8:45 am]
BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Substance Abuse and Mental Health Services Administration
Current List of Laboratories Which Meet Minimum Standards To Engage in Urine Drug Testing for Federal Agencies

AGENCY: Substance Abuse and Mental Health Services Administration, HHS.
ACTION: Notice.
SUMMARY: The Department of Health and Human Services (HHS) notifies Federal agencies of the laboratories currently certified to meet the standards of Subpart C of the Mandatory Guidelines for Federal Workplace Drug Testing Programs (Mandatory Guidelines). The Mandatory Guidelines were first published in the Federal Register on April 11, 1988 (53 FR 11970), and subsequently revised in the Federal Register on June 9, 1994 (59 FR 29908), on September 30, 1997 (62 FR 51118), and on April 13, 2004 (69 FR 19644).

A notice listing all currently certified laboratories is published in the Federal Register during the first week of each month. If any laboratory’s certification is suspended or revoked, the laboratory will be omitted from subsequent lists until such time as it is restored to full certification under the Mandatory Guidelines.

If any laboratory has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end, and will be omitted from the monthly listing thereafter. This notice is also available on the Internet at http://workplace.samhsa.gov and http://www.drugfreenetwork.gov.

FOR FURTHER INFORMATION CONTACT: Mrs. Giselle Hersh or Dr. Walter Vogl, Division of Workplace Programs, SAMHSA/CSAP, Room 2–1035, 1 Choke Cherry Road, Rockville, Maryland 20857; 240–276–2600 (voice), 240–276–2610 (fax).

SUPPLEMENTARY INFORMATION: The Mandatory Guidelines were developed in accordance with Executive Order 12564 and section 503 of Pub. L. 100–71. Subpart C of the Mandatory Guidelines, “Certification of Laboratories Engaged in Urine Drug Testing for Federal Agencies,” sets strict standards that laboratories must meet in order to conduct drug and specimen validity tests on urine specimens for Federal agencies. To become certified, an applicant laboratory must undergo three rounds of performance testing plus an on-site inspection. To maintain that certification, a laboratory must participate in a quarterly performance testing program plus undergo periodic, on-site inspections.

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

In accordance with Subpart C of the Mandatory Guidelines dated April 13, 2004 (69 FR 19644), the following laboratories meet the minimum standards to conduct drug and specimen validity tests on urine specimens:


Baptist Medical Center-Toxicology Laboratory, 9601 I–630, Exit 7, Little Rock, AR 72205–7299, 501–202–2783, (Formerly: Forensic Toxicology Laboratory Baptist Medical Center).

Clinical Reference Lab, 8433 Quivira Road, Lenexa, KS 66215–2902, 800–445–6017.


Doctors Laboratory, Inc., 2906 Julia Drive, Valdosta, GA 31602, 229–671–2281.

DrugScan, Inc., P.O. Box 2969, 1119 Mearns Road, Warminster, PA 18974, 215–674–9310.


General Medical Laboratories, 36 South Brooks St., Madison, WI 53715, 608–267–6225.

Kroll Laboratory Specialists, Inc., 1111 Newton St., Gretta, LA 70053, 504–361–8989 / 800–433–3823, (Formerly: Laboratory Specialists, Inc.).


Laboratory Corporation of America Holdings, 7207 N. Gessner Road, Houston, TX 77040, 713–856–8288 / 800–800–2387.

Laboratory Corporation of America Holdings, 69 First Ave., Raritan, NJ 08869, 908–502–2400 / 800–437–4986, (Formerly: Roche Biomedical Laboratories, Inc.).


Laboratory Corporation of America Holdings, 10788 Roselle St., San Diego, CA 92121, 800–882–7272, (Formerly: Forestalab, Inc.).

Laboratory Corporation of America Holdings, 550 17th Ave., Suite 300, Seattle, WA 98122, 206–923–7020 / 800–898–0180, (Formerly: DrugProof, Division of Dynacare/Laboratory of Pathology, LLC; Laboratory of Pathology of Seattle, Inc.; DrugProof, Division of Laboratory of Pathology of Seattle, Inc.).

Laboratory Corporation of America Holdings, 1120 Main Street, Southaven, MS 38671, 666–827–8042 / 800–233–6339, (Formerly: LabCorp Occupational Testing Services, Inc.; MedExpress/National Laboratory Center).

Marshfield Laboratories, Forensic Toxicology Laboratory, 1000 North Oak Ave., Marshfield, WI 54449, 715–389–3734 / 800–331–3734.

MAXXAM Analytics Inc.*, 6740 Campobello Road, Mississauga, ON, Canada L5N 2L8, 905–817–5700, (Formerly: NOVAMANN (Ontario), Inc.).


MetroLab-Legacy Laboratory Services, 1225 NE 2nd Ave., Portland, OR 97232, 503–413–5295 / 800–950–5295.

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, 800–326–6942, (Formerly: Centinela Hospital Airport Toxicology Laboratory).


Quest Diagnostics Incorporated, 4770 Regent Blvd., Irving, TX 75063, 800–824–6152, (Moved from the Dallas location on 03/31/01; Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories).


Quest Diagnostics Incorporated, 10101 Renner Blvd., Lenexa, KS 66219, 913–888–3927 / 800–873–8845, (Formerly: LabOne, Inc.; Center for Laboratory Services, a Division of LabOne, Inc.).


Quest Diagnostics Incorporated, 2282 South Presidents Drive, Suite C, West Valley City, UT 84120, 801–606–6301 / 800–322–3361, (Formerly: Northwest Toxicology, a LabOne Company; LabOne, Inc., dba Northwest Toxicology; NWT Drug Testing, NorthWest Toxicology, Inc.; Northwest Drug Testing, a division of NWI Inc.).


South Bend Medical Foundation, Inc., 530 N. Lafayette Blvd., South Bend, IN 46601, 574–234–4176 x276.


Sparrow Health System, Toxicology Testing Center, St. Lawrence Campus, 1210 W. Saginaw, Lansing, MI 48915, 517–364–7400, (Formerly: St. Lawrence Hospital & Healthcare System).

St. Anthony Hospital Toxicology Laboratory, 1000 N. Lee St., Oklahoma City, OK 73101, 405–272–7052.

Toxicology & Drug Monitoring Laboratory, University of Missouri Hospital & Clinics, 301 Business Loop 70 West, Suite 208, Columbia, MO 65203, 573–882–1273.


U.S. 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 (LAPSA) 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 LAPSA-accredited laboratories

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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 13, 2004 (69 FR 19644). After receiving DOT certification, the laboratory will be included in the monthly list of HHS-certified laboratories and participate in the NLCP certification maintenance program.

Anna Marsh,
Director, Office Program Services, SAMHSA.

[FR Doc. E6–13237 Filed 8–11–06; 8:45 am]

DEPARTMENT OF THE INTERIOR
Fish and Wildlife Service

Receipt of Applications for Permit

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice of receipt of applications for permit.

SUMMARY: The public is invited to comment on the following applications to conduct certain activities with endangered species.

DATES: Written data, comments or requests must be received by September 13, 2006.

ADDRESSES: Documents and other information submitted with these applications are available for review, subject to the requirements of the Privacy Act and Freedom of Information Act, by any party who submits a written request for a copy of such documents within 30 days of the date of publication of this notice to: U.S. Fish and Wildlife Service, Division of Management Authority, 4401 North Fairfax Drive, Room 700, Arlington, Virginia 22203; fax 703/358–2281.

FOR FURTHER INFORMATION CONTACT: Division of Management Authority, telephone 703/358–2104.

SUPPLEMENTARY INFORMATION:

Endangered Species

The public is invited to comment on the following applications for a permit to conduct certain activities with endangered species. This notice is provided pursuant to Section 10(c) of the Endangered Species Act of 1973, as amended (16 U.S.C. 1531 et seq.). Written data, comments, or requests for copies of these complete applications should be submitted to the Director (address above).

Applicant: University of Texas at Austin, Austin, TX, PRT–124346

The applicant requests a permit to import biological samples from Verreaux’s sifaka (Propithecus verreauxi) collected in the wild in Madagascar, for scientific research. This notification covers activities to be conducted by the applicant over a five-year period.

Applicant: Virginia Polytechnic Institute and State University, Blacksburg, VA, PRT–132043

The applicant requests a permit to import biological samples from chimpanzees (Pan troglodytes) in Tanzania for the purpose of scientific research. This action was authorized under Section 10(c) of the Endangered Species Act of 1973, as amended (16 U.S.C. 1531 et seq.). The Service determined that an emergency affecting the health and life of the chimpanzees at the Mahale Mountains National Park and Rubondo Island National Park in Tanzania existed and that no reasonable alternative was available to the applicant for the following reasons.

Virginia Polytechnic Institute and State University requested a permit to import biological samples (bodily tissues and organs, hair, saliva, and other body parts) from the forest floor and from deceased animals found in the Mahale Mountains National Park in Kigoma, Tanzania, and Rubondo Island National Park in Mwanza, Tanzania, for emergency and ongoing health and disease evaluation purposes. Samples will be utilized exclusively for diagnostic and scientific purposes. The specimens will be used to run diagnostics to determine the cause of death. The necessary diagnostic testing is not available in Africa. The results of health and disease testing from these chimpanzees will help determine why the animals died in order to develop interventions to help prevent reoccurrence.


Michael S. Moore,
Senior Permit Biologist, Branch of Permits, Division of Management Authority.

[FR Doc. E6–13239 Filed 8–11–06; 8:45 am]

DEPARTMENT OF THE INTERIOR
Fish and Wildlife Service

Emergency Exemption: Issuance of Permit for Endangered Species

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice of emergency issuance of permit for endangered species.

SUMMARY: The following permit was issued.

ADDRESSES: Documents and other information submitted for this application are available for review, subject to the requirements of the Privacy Act and Freedom of Information Act, by any party who submits a written request for a copy of such documents to: U.S. Fish and Wildlife Service, Division of Management Authority, 4401 North Fairfax Drive, Room 700, Arlington, Virginia 22203; telephone 703/358–2104 or fax 703/358–2281.

FOR FURTHER INFORMATION CONTACT: Division of Management Authority, telephone 703/358–2104.

SUPPLEMENTARY INFORMATION: On July 21, 2006, the U.S. Fish and Wildlife Service (Service) issued a permit (PRT–108841) to the Virginia Polytechnic Institute and State University, Blacksburg, Virginia, to import biological samples from wild chimpanzees (Pan troglodytes) in Tanzania for the purpose of scientific research. This action was authorized under Section 10(c) of the Endangered Species Act of 1973, as amended (16 U.S.C. 1531 et seq.). The Service determined that an emergency affecting the health and life of the chimpanzees at the Mahale Mountains National Park and Rubondo Island National Park in Tanzania existed and that no reasonable alternative was available to the applicant for the following reasons.

Virginia Polytechnic Institute and State University requested a permit to import biological samples (bodily tissues and organs, hair, saliva, and other body parts) from the forest floor and from deceased animals found in the Mahale Mountains National Park in Kigoma, Tanzania, and Rubondo Island National Park in Mwanza, Tanzania, for emergency and ongoing health and disease evaluation purposes. Samples will be utilized exclusively for diagnostic and scientific purposes. The specimens will be used to run diagnostics to determine the cause of death. The necessary diagnostic testing is not available in Africa. The results of health and disease testing from these chimpanzees will help determine why the animals died in order to develop interventions to help prevent reoccurrence.


Michael S. Moore,
Senior Permit Biologist, Branch of Permits, Division of Management Authority.

[FR Doc. E6–13243 Filed 8–11–06; 8:45 am]