information to the committee for their consideration. Background information for each day's topic, including the agenda and questions for the committee, will be available to the public 1-business day before the meeting on the Internet at http://www.fda.gov/cdrh/panelmtg.html. Material for the July 8, 2002, session will be posted on July 5, 2002; material for the July 9, 2002, session will be posted on July 8, 2002.

Procedure: On July 8, 2002, from 1:30 p.m. to 5 p.m., and on July 9, 2002, from 8 a.m. to 5 p.m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by July 5, 2002. Oral presentations from the public will be scheduled between approximately 1:45 p.m. and 2:15 p.m. and 4 p.m. and 4:30 p.m. on July 8, 2002; and between approximately 8:30 a.m. and 10:30 a.m. on July 9, 2002. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person by July 5, 2002, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Closed Committee Deliberations: On July 8, 2002, from 1 p.m. to 1:30 p.m., the meeting will be closed to permit discussion and review of trade secret and/or confidential commercial information (5 U.S.C. 552b(c)(4)) relating to pending issues and applications.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams, Conference Management Staff, at 301–594–1283, ext. 113, by July 5, 2002.

FDA regrets that it was unable to publish this notice 15 days prior to the General and Plastic Surgery Devices Panel of the Medical Devices Advisory Committee meeting. Because the agency believes there is some urgency to bring these issues to public discussion and qualified members of the General and Plastic Surgery Devices Panel of the Medical Devices Advisory Committee were available at this time, the Commissioner of Food and Drugs concluded that it was in the public interest to hold this meeting even if there was not sufficient time for the customary 15-day public notice.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: June 25, 2002.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 02–16734 Filed 6–28–02; 3:10 pm]

BILLING CODE 4160–01–S

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 notifies Federal agencies of the laboratories currently certified to meet standards of Subpart C of Mandatory Guidelines for Federal Workplace Drug Testing Programs (59 FR 29916, 29925). 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 Guidelines.

If any laboratory has withdrawn from the National Laboratory Certification Program 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 the following Web sites: *http://workplace.samhsa.gov and http://www.drugfreeworkplace.gov.*

FOR FURTHER INFORMATION CONTACT: Mrs. Giselle Hersh or Dr. Walter Vogl, Division of Workplace Programs, 5600 Fishers Lane, Rockwall 2 Building, Room 815, Rockville, Maryland 20857; Tel.: (301) 443–6014, Fax: (301) 443–3031.

SUPPLEMENTARY INFORMATION: Mandatory Guidelines for Federal Workplace Drug Testing were developed in accordance with Executive Order 12564 and section 503 of Pub. L. 100–71. Subpart C of the Guidelines, ''Certification of Laboratories Engaged in Urine Drug Testing for Federal Agencies,'' sets strict standards which laboratories must meet in order to conduct urine drug testing 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 periodic, on-site inspections.

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

In accordance with Subpart C of the Guidelines, the following laboratories meet the minimum standards set forth in the Guidelines:

ACL Laboratories, 8901 W. Lincoln Ave., West Allis, WI 53227, 414–328–7840/800–877–7016, (Formerly: Bayshore Clinical Laboratory)

ACM Medical Laboratory, Inc., 160 Elmgrove Park, Rochester, NY 14624, 716–429–2264

Advanced Toxicology Network, 3560 Air Center Cove, Suite 101, Memphis, TN 38118, 901–794–5770/888–290–1150

Aegis Analytical Laboratories, Inc., 345 Hill Ave., Nashville, TN 37210, 615–255–2400

Alliance Laboratory Services, 3200 Burnet Ave., Cincinnati, OH 45229, 513–585–9000, (Formerly: Jewish Hospital of Cincinnati, Inc.)

American Medical Laboratories, Inc., 14225 Newbrook Dr., Chantilly, VA 20151, 703–802–6900

Associated Pathologists Laboratories, Inc., 4230 South Burnham Ave., Suite 250, Las Vegas, NV 89119–5412, 702–733–7866 / 800–433–2750

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 Laboratory Partners, LLC, 129 East Cedar St., Newington, CT 06111, 860–696–8115, (Formerly: Hartford Hospital Toxicology Laboratory)

Clinical Reference Lab, 8433 Quivira Rd., Lenexa, KS 66215–2802, 800–445–6917

Cox Health Systems, Department of Toxicology, 1423 North Jefferson Ave., Springfield, MO 65802, 800–876–3652/417–269–3093, (Formerly: Cox Medical Centers)

Diagnostic Services Inc., dba DSI, 12700 Westlinks Drive, Fort Myers, FL 33913, 941–561–8200 / 800–735–5416
Doctors Laboratory, Inc., P.O. Box 2658, 2906 Julia Dr., Valdosta, GA 31602, 912–244–4468
DrugProof, Divison of Dynacare, 543 South Hull St., Montgomery, AL 36103, 888–777–9497/334–241–0522, (Formerly: Alabama Reference Laboratories, Inc.)
DrugProof, Division of Dynacare/ Laboratory of Pathology, LLC, 1229 Madison St., Suite 500, Nordstrom Medical Tower, Seattle, WA 98104, 206–386–2672/800–898–0180, (Formerly: Laboratory of Pathology of Seattle, Inc., DrugProof, Division of Laboratory of Pathology of Seattle, Inc.)
DrugScan, Inc., P.O. Box 2969, 1119 Mearns Rd., Warminster, PA 18974, 215–674–9310
Dynacare Kasper Medical Laboratories*, 14940–123 Ave., Edmonton, Alberta Canada T5V 1B4, 780–451–3702/800–661–9876
ElSohly Laboratories, Inc., 5 Industrial Park Dr., Oxford, MS 38655, 662–236–2609
Express Analytical Labs, 3405 7th Avenue, Suite 106, Marion, IA 52302, 319–377–0500
Gamma-Dynacare Medical Laboratories*, A Division of the Gamma-Dynacare Laboratory Partnership, 245 Pall Mall St., London, ONT Canada N6A 1P4, 519–679–1630
General Medical Laboratories, 36 South Brooks St., Madison, WI 53715, 608–267–6267
Kroll Laboratory Specialists, Inc., 1111 Newton St., Gretna, LA 70053, 504–361–8989/800–433–3823, (Formerly: Laboratory Specialists, Inc.)
LabOne, Inc., 10101 Renner Blvd., Lenexa, KS 66219, 913–888–3927/ 800–728–4064, (Formerly: Center for Laboratory Services, a Division of LabOne, 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, 908–526–2400/800–437–4986, (Formerly: Roche Biomedical Laboratories, Inc.)
Laboratory Corporation of America Holdings, 1904 Alexander Drive, Research Triangle Park, NC 27709, 919–572–6900/800–833–3984, (Formerly: LabCorp Occupational Testing Services, Inc., CompuChem Laboratories, Inc.; CompuChem Laboratories, Inc., A Subsidiary of Roche Biomedical Laboratory; Roche CompuChem Laboratories, Inc., A Member of the Roche Group)
Laboratory Corporation of America Holdings, 10788 Roselle Street, San Diego, CA 92121, 800–882–7272, (Formerly: Poisonlab, Inc.)
Laboratory Corporation of America Holdings, 1120 Stateline Road West, Southaven, MS 38671, 866–827–8042/ 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.*, 5540 McAdam Rd., Mississauga, ON Canada L4Z 1P1, 905–890–2555, (Formerly: NOVAMANN (Ontario) Inc.)
Medical College Hospitals Toxicology Laboratory, Department of Pathology, 3000 Arlington Ave., Toledo, OH 43699, 419–383–5213
MedTox Laboratories, Inc., 402 W. County Rd. D, St. Paul, MN 55112, 651–636–7466/800–832–3244
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, Minnesota 55417, 612–725–2088
National Toxicology Laboratories, Inc., 1100 California Ave., Bakersfield, CA 93304, 661–322–4250/800–350–3515
Northwest Drug Testing, a division of NWT Inc., 1141 E. 3900 South, Salt Lake City, UT 84124, 801–293–2300/ 800–322–3361, (Formerly: NWT Drug Testing, NorthWest Toxicology, Inc.)
One Source Toxicology Laboratory, Inc., 1705 Center Street, Deer Park, TX 77536, 713–920–2559, (Formerly: University of Texas Medical Branch, Clinical Chemistry Division; UTMB Pathology-Toxicology Laboratory)
Oregon Medical Laboratories, P.O. Box 972, 722 East 11th Ave., Eugene, OR 97440–0972, 541–687–2134
Pacific Toxicology Laboratories, 6160 Variel Ave., Woodland Hills, CA 91367, 818–598–3110/800–328–6942, (Formerly: Centinela Hospital Airport Toxicology Laboratory
Pathology Associates Medical Laboratories, 110 West Cliff Drive, Spokane, WA 99204, 509–755–8991/ 800–541–7891x8991
PharmChem Laboratories, Inc., 4600 N. Beach, Haltom City, TX 76137, 817–605–5300, (Formerly: PharmChem Laboratories, Inc., Texas Division; Harris Medical Laboratory)
Physicians Reference Laboratory, 7800 West 110th St., Overland Park, KS 66210, 913–339–0372/800–821–3627
Quest Diagnostics Incorporated, 3175 Presidential Dr., Atlanta, GA 30340, 770–452–1590, (Formerly: SmithKline Beecham Clinical Laboratories, SmithKline Bio-Science Laboratories)
Quest Diagnostics Incorporated, 4770 Regent Blvd., Irving, TX 75063, 800–842–6152, (Moved from the Dallas location on 033101; Formerly: SmithKline Beecham Clinical Laboratories, SmithKline Bio-Science Laboratories)
Quest Diagnostics Incorporated, 400 Egypt Rd., Norristown, PA 19403, 610–631–4600/877–642–2216, (Formerly: SmithKline Beecham Clinical Laboratories, SmithKline Bio-Science Laboratories)
Quest Diagnostics Incorporated, 506 E. State Pkwy., Schaumburg, IL 60173, 800–669–6995847–885–2010, (Formerly: SmithKline Beecham Clinical Laboratories, International Toxicology Laboratories)
Quest Diagnostics Incorporated, 7600 Tyrone Ave., Van Nuys, CA 91405, 818–989–2520/800–877–2520, (Formerly: SmithKline qBeecham Clinical Laboratories)
Scientific Testing Laboratories, Inc., 463 Southlake Blvd., Richmond, VA 23236, 804–378–9130
S.E.D. Medical Laboratories, 5601 Office Blvd., Albuquerque, NM 87109, 505–727–6300/800–999–5227
South Bend Medical Foundation, Inc., 530 N. Lafayette Blvd., South Bend, IN 46601, 219–234–4176
Southwest Laboratories, 2727 W. Baseline Rd., Tempe, AZ 85283, 602–438–8507/800–279–0027
Sparrow Health System, Toxicology Testing Center, St. Lawrence Campus, 1210 W. Saginaw, Lansing, MI 48915, 517–377–0520, (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, 2703 Clark Lane, Suite B, Lower Level, Columbia, MO 65202, 573–882–1273
Toxicology Testing Service, Inc., 5426 N.W. 79th Ave., Miami, FL 33166, 305–593–2260
US Army Forensic Toxicology Drug Testing Laboratory, Fort Meade, Building 2490, Wilson Street, 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 was transferred to the U.S. DHHS, with the DHHS' National Laboratory Certification Program (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, the DHHS will recommend that DOT certify the laboratory (Federal Register, 16 July 1996) as meeting the minimum standards of the ''Mandatory Guidelines for Workplace Drug Testing'' (59 Federal Register, 9 June 1994, Pages After receiving the DOT certification, the laboratory will be included in the monthly list of DHHS certified laboratories and participate in the NLCP certification maintenance program.

Patricia Bransford,

Acting Executive Officer, SAMHSA.

[FR Doc. 02–16696 Filed 7–2–02; 8:45 am]

BILLING CODE 4160–20–P

DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

Endangered and Threatened Species Permit Applications

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice of receipt of applications.

SUMMARY: The following applicants have applied for a scientific research permit to conduct certain activities with endangered species pursuant to section 10(a)(1)(A) of the Endangered Species Act of 1973, as amended (16 U.S.C. 1531, *et seq.*).

DATES: Written comments on these permit applications must be received within 30 days of the date of publication.

ADDRESSES: Written data or comments should be submitted to the Chief, Endangered Species Division, Ecological Services, PO Box 1306, Room 4102, Albuquerque, New Mexico 87103; (505) 248–6649; Fax (505) 248–6788. Documents will be available for public inspection by written request, by appointment only, during normal business hours (8:00 to 4:30) at the U.S. Fish and Wildlife Service, 500 Gold Ave. SW., Room 4102, Albuquerque, New Mexico. Please refer to the respective permit number for each application when submitting comments. All comments received, including names and addresses, will become part of the official administrative record and may be made available to the public.

FOR FURTHER INFORMATION CONTACT: Chief, Endangered Species Division, PO Box 1306, Room 4102, Albuquerque, New Mexico 87103. 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 to the address above for a copy of such documents within 30 days of the date of publication of this notice.

SUPPLEMENTARY INFORMATION:

Permit No. TE–056457

Applicant: Peter Stacey, Albuquerque, New Mexico

Applicant requests a permit for recovery purposes to conduct presence/absence and monitoring surveys for southwestern willow flycatcher (Empidonax traillii extimus) within New Mexico.

Permit No. TE–055107

Applicant: Adam Crateau, Santa Fe, New Mexico

Applicant requests a permit for recovery purposes to conduct presence/absence surveys for southwestern willow flycatcher (Empidonax traillii extimus) within San Juan Counties of New Mexico and Utah.

Permit No. TE–055111

Applicant: Animas Environmental, Farmington, New Mexico

Applicant requests a permit for recovery purposes to conduct presence/absence surveys for southwestern willow flycatcher (Empidonax traillii extimus) within San Juan County, New Mexico.

Permit No. TE–055339

Applicant: USDA, ARS, Grassland, Soil and Water Research Laboratory, Temple, Texas

Applicant requests a permit for recovery purposes to collect live plants and stem cuttings of Johnston's Frankenia (Frankenia johnstonii) within Texas.

Permit No. TE–055419

Applicant: Turner Biological Consulting, LLC, Tuscola, Texas

Applicant requests a permit for recovery purposes to conduct presence/absence surveys and assess habitat for black-capped vireo (Vireo atricapillus) and to assess habitat for northern aplomado falcon (Falco femoralis septentrionalis), golden-cheeked warbler (Dendroica chrysoparia), and interior least tern (Sterna antillarum). All activities are to occur within Texas.

Permit No. TE–035885

Applicant: Melvin J. Wilhelm, Vernon, Arizona

Applicant requests an amendment to an existing permit to allow presence/absence surveys for southwestern willow flycatcher (Empidonax traillii extimus) within Arizona and New Mexico.

Permit No. TE–056119

Applicant: Marlin B. Sawyer, San Antonio, Texas

Applicant requests a permit for recovery purposes to conduct presence/absence surveys within Texas for the following species: black-capped vireo (Vireo atricapillus), golden-cheeked warbler (Dendroica chrysoparia), red-cockaded woodpecker (Picoides borealis), ocelot (Leopardus pardalis), jaguarundi (Herpailurus yagouaroundi cacomitli), and Houston toad (Bufo houstonensis).

Permit No. TE–056118

Applicant: Charles L. Black, Albuquerque, New Mexico

Applicant requests a permit for recovery purposes to conduct presence/absence and monitoring surveys for southwestern willow flycatcher (Empidonax traillii extimus) within New Mexico.

Permit No. TE–056471

Applicant: Gregory Tickle, Santa Fe, New Mexico

Applicant requests a permit for recovery purposes to conduct presence/absence surveys within Texas for the following species: black-capped vireo (Vireo atricapillus), golden-cheeked warbler (Dendroica chrysoparia), and southwestern willow flycatcher (Empidonax traillii extimus).

Permit No. TE–038050

Applicant: Trevor Hare, Tucson, Arizona

Applicant requests an amendment to an existing permit to allow presence/absence surveys for desert pupfish (Cyprinodon macularius) and Gila