Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Jason D. Hoffert, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 7343, 6707 Democracy Boulevard, Bethesda, MD 20817, 301–496–9010, hoffertj@niddk.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: May 30, 2017.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017-11496 Filed 6-2-17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Current List of HHS-Certified Laboratories and Instrumented Initial Testing Facilities 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 and Instrumented Initial Testing Facilities (IITF) currently certified to meet the standards of the Mandatory Guidelines for Federal Workplace Drug Testing Programs (Mandatory Guidelines).

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

If any laboratory or IITF has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end and will be omitted from the monthly listing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://www.samhsa.gov/workplace.

FOR FURTHER INFORMATION CONTACT: Giselle Hersh, Division of Workplace

Programs, SAMHSA/CSAP, 5600 Fishers Lane, Room 16N03A, Rockville, Maryland 20857; 240–276–2600 (voice).

SUPPLEMENTARY INFORMATION: The Department of Health and Human Services (HHS) notifies federal agencies of the laboratories and Instrumented Initial Testing Facilities (IITF) currently certified to meet the standards 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); September 30, 1997 (62 FR 51118); April 13, 2004 (69 FR 19644); November 25, 2008 (73 FR 71858); December 10, 2008 (73 FR 75122); and on April 30, 2010 (75 FR 22809).

The Mandatory Guidelines were initially developed in accordance with Executive Order 12564 and section 503 of Public Law 100–71. The "Mandatory Guidelines for Federal Workplace Drug Testing Programs," as amended in the revisions listed above, requires strict standards that laboratories and IITFs must meet in order to conduct drug and 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 HHScertified laboratories and IITFs meet the minimum standards to conduct drug and specimen validity tests on urine specimens:

HHS-Certified Instrumented Initial Testing Facilities

Dynacare, 6628 50th Street NW., Edmonton, AB Canada T6B 2N7, 780– 784–1190 (Formerly: Gamma-Dynacare Medical Laboratories)

HHS-Certified Laboratories

ACM Medical Laboratory, Inc., 160 Elmgrove Park, Rochester, NY 14624, 844–486–9226 Aegis Analytical Laboratories, Inc., 345 Hill Ave., Nashville, TN 37210, 615– 255–2400 (Formerly: Aegis Sciences Corporation, Aegis Analytical Laboratories, Inc., Aegis Analytical Laboratories)

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

Alere Toxicology Services, 450 Southlake Blvd., Richmond, VA 23236, 804–378–9130 (Formerly: Kroll Laboratory Specialists, Inc., Scientific Testing Laboratories, Inc.; Kroll Scientific Testing Laboratories, Inc.)

Baptist Medical Center–Toxicology Laboratory, 11401 I–30, Little Rock, AR 72209–7056, 501–202–2783 (Formerly: Forensic Toxicology Laboratory Baptist Medical Center)

Clinical Reference Laboratory, Inc., 8433 Quivira Road, Lenexa, KS 66215– 2802, 800–445–6917,

DrugScan, Inc., 200 Precision Road, Suite 200, Horsham, PA 19044, 800– 235–4890

Dynacare *, 245 Pall Mall Street, London, ONT, Canada N6A 1P4, 519– 679–1630 (Formerly: Gamma-Dynacare Medical Laboratories)

ElSohly Laboratories, Inc., 5 Industrial Park Drive, Oxford, MS 38655, 662– 236–2609

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, 1120 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.)

MedTox Laboratories, Inc., 402 W. County Road 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, MN 55417, 612–725– 2088, Testing for Veterans Affairs (VA) Employees Only

National Toxicology Laboratories, Inc., 1100 California Ave., Bakersfield, CA 93304, 661–322–4250/800–350–3515

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: Centinela
Hospital Airport Toxicology
Laboratory)

Pathology Associates Medical Laboratories, 110 West Cliff Dr., Spokane, WA 99204, 509–755–8991/ 800–541–7891 x7

Phamatech, Inc., 15175 Innovation Drive, San Diego, CA 92128, 888– 635–5840

Quest Diagnostics Incorporated, 1777 Montreal Circle, Tucker, GA 30084, 800–729–6432 (Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories)

Quest Diagnostics Incorporated, 400 Egypt Road, Norristown, PA 19403, 610–631–4600/877–642–2216 (Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories)

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

Redwood Toxicology Laboratory, 3700 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, Testing for Department of Defense (DoD) Employees Only

Charles LoDico,

Chemist.

[FR Doc. 2017–11512 Filed 6–2–17; 8:45 am]

BILLING CODE 4160-20-P

DEPARTMENT OF THE INTERIOR

Bureau of Land Management

[17X.LLAK942000.L54200000.FR0000. LVDIL0440000; AA086373]

Notice of Application for a Recordable Disclaimer of Interest for Lands Underlying the George River in Alaska

AGENCY: Bureau of Land Management, Interior.

ACTION: Notice.

SUMMARY: The State of Alaska (State) has filed an application with the Bureau of Land Management (BLM) for a Recordable Disclaimer of Interest (RDI) from the United States in those lands underlying the George River from its mouth to Julian Creek. The State asserts that the George River, a major tributary of the Kuskokwim River in southwestern Alaska, was navigable and unreserved at the time of Alaska Statehood in 1959.

DATES: Comments on this action are due on or before September 5, 2017.

ADDRESSES: You may submit comments on the State of Alaska's Application for an RDI or the BLM Draft Summary Report for the State's Application for a Recordable Disclaimer of Interest by mail or email. To file by mail, send to: RDI Program Manager (AK–942), Division of Lands and Cadastral, BLM Alaska State Office, 222 West 7th Avenue, #13, Anchorage, AK 99513. To submit by email, send to: anichols@blm.gov.

FOR FURTHER INFORMATION CONTACT:

Angie Nichols, RDI Program Manager, at 222 West 7th Avenue, #13, Anchorage, AK 99513; 907–271–3359; or anichols@blm.gov; or visit the BLM Recordable Disclaimer of Interest Web site at

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. 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 November 25, 2008 (73 FR 71858). 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.

https://www.blm.gov/programs/lands-and-realty/regional-information/alaska/RDI/kuskokwim. Persons who use a telecommunications device for the deaf (TDD) may call the Federal Relay System (FRS) at 1–800–877–8339 to contact the individual identified in this section during normal business hours. The FRS is available 24 hours a day, seven days a week, to leave a message or a question with that individual. You will receive a reply during normal business hours.

SUPPLEMENTARY INFORMATION: On March 10, 2006, as modified on September 16, 2015, the State of Alaska filed an application (AA-86373) for an RDI pursuant to section 315 of the Federal Land Policy and Management Act of 1976 and the regulations contained in 43 CFR subpart 1864 for the lands underlying the George River. The State asserts that this river was navigable at the time of Alaska Statehood. As such, the State contends that ownership of the lands underlying this river automatically passed from the United States to the State of Alaska in 1959 at the time of Statehood under the Equal Footing Doctrine; the Submerged Lands Act of 1953; the Alaska Statehood Act; and other title navigability law.

The State's application is for all submerged lands underlying the portion of the George River from its mouth to Julian Creek. Specifically, these are the submerged lands within the bed of the George River between the ordinary high water mark of the left and right banks, beginning at the confluence of Julian Creek in Township 24 North, Range 44 West, Section 4, Seward Meridian, Alaska, U.S. Geological Survey (USGS) 1:63,360 series topographic map Iditarod A-3 (1954). Thence southerly to its confluence with the Kuskokwim River in Township 21 North, Range 46 West, Section 21, Seward Meridian, Alaska, USGS 1:63,360 series topographic map Sleetmute D-5 (1954, minor revisions 1975). The applied section of the George River flows through the following Townships and Ranges:

Seward Meridian:

Township 24 North, Ranges 44–45

Township 23 North, Ranges 45–46 West;

Township 22 North, Ranges 45–46

Township 21 North, Range 46 West. The precise location may be within other townships due to the ambulatory nature of these water bodies.

An RDI is a legal document through which the United States disavows ownership of specified land, but it does

^{*} The Standards Council of Canada (SCC) voted to end its Laboratory Accreditation Program for