

Short-Term Research Education to Increase Diversity.

Date: May 23, 2016.

Time: 12:00 p.m. to 2:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Room 7189, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Stephanie L. Constant, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7189, Bethesda, MD 20892, 301-443-8784, [constants@nhlbi.nih.gov](mailto:constants@nhlbi.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: April 27, 2016.

**Michelle Trout,**

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016-10261 Filed 5-2-16; 8:45 am]

BILLING CODE 4140-01-P

## 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). 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).

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If any laboratory or IITF has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end and will be omitted from the monthly listing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 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 HHS-certified 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  
585-429-2264

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 Lab  
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

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  
713-856-8288/800-800-2387

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  
509-755-8991/800-541-7891x7

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  
3700650 Westwind Blvd.  
Santa Rosa, CA 95403  
800-255-2159

Southwest Laboratories  
4625 E. Cotton Center Boulevard  
Suite 177  
Phoenix, AZ 85040  
602-438-8507/800-279-0027

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

U.S. 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  
\* 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. 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, 2010 (75 FR 22809). 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.

**Summer King**,  
*Statistician*.

[FR Doc. 2016-10260 Filed 5-2-16; 8:45 am]

**BILLING CODE 4160-20-P**

## DEPARTMENT OF HOMELAND SECURITY

### Coast Guard

[Docket No. USCG-2015-0757]

### Collection of Information Under Review by Office of Management and Budget; OMB Control Number: 1625-0041

**AGENCY:** Coast Guard, DHS.

**ACTION:** Thirty-day notice requesting comments.

**SUMMARY:** In compliance with the Paperwork Reduction Act of 1995 the U.S. Coast Guard is forwarding an Information Collection Request (ICR), abstracted below, to the Office of Management and Budget (OMB), Office of Information and Regulatory Affairs (OIRA), requesting approval for reinstatement, with change, of the following collection of information: 1625-0041, Various International Agreement Pollution Prevention Certificates and Documents, and Equivalency Certificates. Our ICR describes the information we seek to collect from the public. Review and comments by OIRA ensure we only impose paperwork burdens commensurate with our performance of duties.

**DATES:** Comments must reach the Coast Guard and OIRA on or before June 2, 2016.

**ADDRESSES:** You may submit comments identified by Coast Guard docket number [USCG-2015-0757] to the Coast Guard using the Federal eRulemaking Portal at <http://www.regulations.gov>. Alternatively, you may submit comments to OIRA using one of the following means:

(1) *Email:* [OIRA-submission@omb.eop.gov](mailto:OIRA-submission@omb.eop.gov).

(2) *Mail:* OIRA, 725 17th Street NW., Washington, DC 20503, attention Desk Officer for the Coast Guard.