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, *constantsl@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If any laboratory or IITF has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end and will be omitted from the monthly listing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.*

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