individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: AIDS/HIV Molecular Biology.

Date: December 8, 2011.

Time: 12 p.m. to 3:30 p.m.

Agenda: To review and evaluate grant applications.

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

Contact Person: Eduardo A Montalvo, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5108, MSC 7852, Bethesda, MD 20892, (301) 435–1168, montalve@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: AIDS/HIV Drug Development.

Date: December 12, 2011.

Time: 1 p.m. to 3 p.m.

Agenda: To review and evaluate grant applications.

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

Contact Person: Eduardo A Montalvo, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5108, MSC 7852, Bethesda, MD 20892, (301) 435–1168, montalve@csr.nih.gov.


Jennifer S. Spaeth, Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2011–28552 Filed 11–2–11; 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 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 certified Laboratories and Instrumented Initial Testing Facilities (IITF) is published in the Federal Register during the first week of each month. If any Laboratory/ IITF’s certification is suspended or revoked, the Laboratory/IITF will be omitted from subsequent lists until such time as it is restored to full certification under the Mandatory Guidelines.

If any Laboratory/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.workplace.samhsa.gov and http://www.drugfreeworkplace.gov.

FOR FURTHER INFORMATION CONTACT: Mrs. Giselle Hersh, Division of Workplace Programs, SAMSHA/CSAP, Room 2–1042, One Choke Cherry Road, Rockville, Maryland 20857; (240) 276–2600 (voice), (240) 276–2610 (fax).

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, require [or set] strict standards that Laboratories and Instrumented Initial Testing Facilities (IITF) must meet in order to conduct drug and specimen validity tests on urine specimens for Federal agencies.

To become certified, an applicant Laboratory/IITF must undergo three rounds of performance testing plus an on-site inspection. To maintain that certification, a Laboratory/IITF must participate in a quarterly performance testing program plus undergo periodic, on-site inspections.

Laboratories and Instrumented Initial Testing Facilities (IITF) in the applicant stage of certification are not to be considered as meeting the minimum requirements described in the HHS Mandatory Guidelines. A Laboratory/ 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 Laboratories and Instrumented Initial Testing Facilities (IITF) meet the minimum standards to conduct drug and specimen validity tests on urine specimens:


Laboratories:

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


Aegis Analytical Laboratories, 345 Hill Ave., Nashville, TN 37210, (615) 255–2400, (Formerly: Aegis Sciences Corporation, Aegis Analytical Laboratories, Inc.)

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.

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

DrugScan, Inc., P.O. Box 2969, 1119 Doctors Laboratory, Inc.)

Kroll Scientific Testing Laboratories, Inc.)

Laboratories, (Formerly: Forensic Toxicology Laboratory Baptist Medical Center.)

Clinical Reference Lab, 8433 Quivira Road, Lenexa, KS 66215–2802, (800) 445–6917.


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
SUMMARY: The Coast Guard announces the availability of recommendations from the Merchant Marine Personnel Advisory Committee in response to Task Statement 75, in which the Coast Guard requested review of the Supplemental Notice of Proposed Rulemaking entitled, “Implementation of the Amendments to the International Convention on Standards of Training, Certification and Watchkeeping for Seafarers, 1978, and Changes to Domestic Endorsements” (STCW SNPRM). The Coast Guard also announces the availability of recommendations from the Merchant Mariner Medical Advisory Committee after its review of the STCW SNPRM.

DATES: Comments and related material must either be submitted to our online docket via http://www.regulations.gov on or before December 5, 2011 or reach the Docket Management Facility by that date.

ADDRESSES: You may submit comments identified by docket number USCG–2004–17914 using any one of the following methods:


(2) Fax: (202) 493–2251.


(4) Hand delivery: Same as mail address above, between 9 a.m. and 5 p.m., Monday through Friday, except Federal holidays. The telephone number is (202) 366–9329.