

Dated: December 27, 2010.,

Robert Hendricks,

Director, Division of Policy and Information Coordination.

[FR Doc. 2010-33062 Filed 12-30-10; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health & Human Development; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The concept meeting, proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Child Health and Human Development Special Emphasis Panel. Contraceptive Clinical Trials Network.

Date: January 12, 2011.

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

Agenda: To review and evaluate concept review.

Place: National Institutes of Health, 6100 Executive Boulevard, Room 2A01, Rockville, MD 20852. (Telephone Conference Call).

Contact Person: Sathasiva B. Kandasamy, PhD, Scientific Review Administrator, Division of Scientific Review, National Institute of Child Health and Human Development, 6100 Executive Boulevard, Rockville, Md 20892-9304. (301) 435-6680. skandasa@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and Infertility Loan Repayment program, National Institutes of Health, HHS)

Dated: December 27, 2010.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010-33067 Filed 12-30-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health & Human Development; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Child Health and Human Development Special Emphasis Panel.

Date: January 10, 2011.

Time: 12:45 p.m. to 3 p.m.

Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, 6100 Executive Boulevard, Room 2A01, Rockville, MD 20852. (Telephone Conference).

Contact Person: Sathasiva B. Kandasamy, PhD, Scientific Review Administrator, Division of Scientific Review, National Institute of Child Health and Human Development, 6100 Executive Boulevard, Rockville, MD 20892-9304. (301) 435-6680. skandasa@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and Infertility Loan Repayment Program, National Institutes of Health, HHS)

Dated: December 27, 2010.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010-33068 Filed 12-30-10; 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, SAMHSA/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, requires {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:

Instrumented Initial Testing Facilities (IITF)

None.

Laboratories

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. 585-429-2264.

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

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 Mearns Road, Warminster, PA 18974. 215-674-9310.

DynaLIFE Dx,* 10150-102 St., Suite 200, Edmonton, Alberta, Canada T5J 5E2. 780-451-3702/800-661-9876. (Formerly: Dynacare Kasper Medical Laboratories.)

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

Gamma-Dynacare Medical Laboratories,* A Division of the Gamma-Dynacare Laboratory Partnership, 245 Pall Mall Street, London, ONT, Canada N6A 1P4. 519-679-1630.

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.)

Maxxam Analytics,* 6740 Campobello Road, Mississauga, ON, Canada L5N

* 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-

2L8. 905-817-5700. (Formerly: Maxxam Analytics Inc., NOVAMANN (Ontario), 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.

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., 10151 Barnes Canyon Road, San Diego, CA 92121. 858-643-5555.

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. 800-877-2520. (Formerly: SmithKline Beecham Clinical Laboratories.)

S.E.D. Medical Laboratories, 5601 Office Blvd., Albuquerque, NM 87109. 505-727-6300/800-999-5227.

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.

South Bend Medical Foundation, Inc.,
530 N. Lafayette Blvd., South Bend,
IN 46601. 574-234-4176 x1276.

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

St. Anthony Hospital Toxicology
Laboratory, 1000 N. Lee St.,
Oklahoma City, OK 73101. 405-272-
7052.

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

Toxicology & Drug Monitoring
Laboratory, University of Missouri
Hospital & Clinics, 301 Business Loop
70 West, Suite 208, Columbia, MO
65203. 573-882-1273.

Toxicology Testing Service, Inc., 5426
N.W. 79th Ave., Miami, FL 33166.
305-593-2260.

U.S. Army Forensic Toxicology Drug
Testing Laboratory, 2490 Wilson St.,
Fort George G. Meade, MD 20755-
5235. 301-677-7085.

Dated: December 15, 2010.

Elaine Parry,

*Director, Office of Management, Technology,
and Operations, SAMHSA.*

[FR Doc. 2010-32908 Filed 12-30-10; 8:45 am]

BILLING CODE 4160-20-P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Agency Information Collection

Activities: CBP Regulations Pertaining to Customs Brokers

AGENCY: U.S. Customs and Border
Protection, Department of Homeland
Security.

ACTION: 30-Day notice and request for
comments; Extension of an existing
information collection: 1651-0034.

SUMMARY: U.S. Customs and Border
Protection (CBP) of the Department of
Homeland Security will be submitting
the following information collection
request to the Office of Management and
Budget (OMB) for review and approval
in accordance with the Paperwork
Reduction Act: CBP Regulations
Pertaining to Customs Brokers (19 CFR
part 111). This is a proposed extension
of an information collection that was
previously approved. CBP is proposing
that this information collection be
extended with no change to the burden
hours or to the information being
collected. This document is published
to obtain comments from the public and
affected agencies. This proposed
information collection was previously

published in the **Federal Register** (75
FR 67094) on November 1, 2010,
allowing for a 60-day comment period.
This notice allows for an additional 30
days for public comments. This process
is conducted in accordance with 5 CFR
1320.10.

DATES: Written comments should be
received on or before February 2, 2011.

ADDRESSES: Interested persons are
invited to submit written comments on
this proposed information collection to
the Office of Information and Regulatory
Affairs, Office of Management and
Budget. Comments should be addressed
to the OMB Desk Officer for Customs
and Border Protection, Department of
Homeland Security, and sent via
electronic mail to
oira_submission@omb.eop.gov or faxed
to (202) 395-5806.

SUPPLEMENTARY INFORMATION: U.S.
Customs and Border Protection (CBP)
encourages the general public and
affected Federal agencies to submit
written comments and suggestions on
proposed and/or continuing information
collection requests pursuant to the
Paperwork Reduction Act (Pub. L. 104-
13). Your comments should address one
of the following four points:

(1) Evaluate whether the proposed
collection of information is necessary
for the proper performance of the
functions of the agency/component,
including whether the information will
have practical utility;

(2) Evaluate the accuracy of the
agencies/components estimate of the
burden of the proposed collection of
information, including the validity of
the methodology and assumptions used;

(3) Enhance the quality, utility, and
clarity of the information to be
collected; and

(4) Minimize the burden of the
collections of information on those who
are to respond, including the use of
appropriate automated, electronic,
mechanical, or other technological
techniques or other forms of
information.

Title: CBP Regulations Pertaining to
Customs Brokers (19 CFR Part 111).

OMB Number: 1651-0034.

Form Numbers: CBP Forms 3124 and
3124E.

Abstract: The information contained
in part 111 of the CBP regulations
governs the licensing and conduct of
customs brokers. Specifically, an
individual who wishes to take the
broker exam would complete CBP Form
3124E, "Application for Customs Broker
License Exam"; or to apply for a broker
license, CBP Form 3124, "Application
for Customs Broker License" must be
completed. The procedures to request a

local or national broker permit can be
found in 19 CFR 111.19, and a triennial
report is required under 19 CFR 111.30.
The information collected from customs
brokers is provided for by 19 U.S.C.
1641. CBP Forms 3124 and 3124E may
be found at [http://www.cbp.gov/xp/
cgov/toolbox/forms/](http://www.cbp.gov/xp/cgov/toolbox/forms/). Further
information about the customs broker
exam and how to apply for it may be
found at [http://www.cbp.gov/xp/cgov/
trade/trade_programs/broker/
broker_exam/notice_of_exam.xml](http://www.cbp.gov/xp/cgov/trade/trade_programs/broker/broker_exam/notice_of_exam.xml).

Current Actions: This submission is
being made to extend the expiration
date with no change to the burden hours
or to this collection of information.

Type of Review: Extension (without
change).

Affected Public: Businesses,
Individuals.

**CBP Form 3124E, "Application for
Customs Broker License Exam
Estimated Number of Respondents:**
2,300.

**Total Number of Estimated Annual
Responses:** 2,300.

Estimated Time per Response: 1 hour.

Estimated Total Annual Burden

Hours: 2,300.

**Estimated Total Annual Cost to the
Public:** \$466,000.

**CBP Form 3124, "Application for
Customs Broker License"**

Estimated Number of Respondents:
300.

**Total Number of Estimated Annual
Responses:** 300.

Estimated Time per Response: 1 hour.

Estimated Total Annual Burden

Hours: 300.

Triennial Report (19 CFR 111.30)

Estimated Number of Respondents:
3,833.

**Total Number of Estimated Annual
Responses:** 3,833.

Estimated Time per Response: .5
hours.

Estimated Total Annual Burden

Hours: 1,917.

**Estimated Total Annual Cost to the
Public:** \$383,300.

**National Broker Permit Application
(19 CFR 111.19)**

Estimated Number of Respondents:
500.

**Total Number of Estimated Annual
Responses:** 500.

Estimated Time per Response: 1 hour.

Estimated Total Annual Burden

Hours: 500.

**Estimated Total Annual Cost to the
Public:** \$112,500.

If additional information is required
contact: Tracey Denning, U.S. Customs
and Border Protection, Regulations and
Rulings, Office of International Trade,
799 9th Street, NW., 5th Floor,