

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### National Cancer Institute; Notice of Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of meeting of the National Cancer Advisory Board.

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.

A portion of the meeting will be closed to the public as indicated below in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual intramural programs and projects conducted by the National Cancer Institute, including consideration of personnel qualifications and performance and the competence of individual investigators. Additionally for the review of grant applications and the discussions, that could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* National Cancer Advisory Board; *Ad hoc* Subcommittee on Global Cancer Research.

*Open:* December 9, 2013, 6:00 p.m. to 7:30 p.m.

*Agenda:* Discussion on Global Cancer Research.

*Place:* Hyatt Regency Bethesda, One Bethesda Metro Center, Bethesda, Maryland 20814.

*Contact Person:* Dr. Edward Trimble, Executive Secretary, NCAB Ad hoc Subcommittee on Global Cancer Research, National Cancer Institute, National Institutes of Health, 9609 Medical Center Drive, Room 3W-562, Bethesda, MD 20892, (240) 276-5796, [trimble@mail.nih.gov](mailto:trimble@mail.nih.gov).

*Name of Committee:* National Cancer Advisory Board; *Ad hoc* Subcommittee on Communications.

*Open:* December 9, 2013, 7:45 p.m. to 9:15 p.m.

*Agenda:* Discussion on Communications.

*Place:* Hyatt Regency Bethesda, One Bethesda Metro Center, Bethesda, Maryland 20814.

*Contact Person:* Dr. Lenora Johnson, Executive Secretary, NCAB *Ad hoc*

Subcommittee on Communications, National Cancer Institute, National Institutes of Health, 9606 Medical Center Drive, Room 2E-454, Bethesda, MD 20892, (240) 276-6680, [johnslen@mail.nih.gov](mailto:johnslen@mail.nih.gov).

*Name of Committee:* National Cancer Advisory Board.

*Open:* December 10, 2013, 9:00 a.m. to 3:30 p.m.

*Agenda:* Program reports and presentations; business of the Board.

*Place:* National Institutes of Health, 9000 Rockville Pike, Building 31, C Wing, 6th Floor, Conference Room 10, Bethesda, MD 20892.

*Contact Person:* Dr. Paulette S. Gray, Executive Secretary, National Cancer Institute, National Institutes of Health, 9606 Medical Center Drive, Room 7W-444, Bethesda, MD 20892, (240) 276-6340.

*Name of Committee:* National Cancer Advisory Board.

*Closed:* December 10, 2013, 3:30 p.m. to 5:00 p.m.

*Agenda:* Review intramural program site visit outcomes. Discussion of confidential and personnel issues. Review and evaluate grant applications.

*Place:* National Institutes of Health, 9000 Rockville Pike, Building 31, C Wing, 6th Floor, Conference Room 10, Bethesda, MD 20892.

*Contact Person:* Dr. Paulette S. Gray, Executive Secretary, National Cancer Institute, National Institutes of Health, 9606 Medical Center Drive, Room 7W-444, Bethesda, MD 20892, (240) 276-6340.

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

In the interest of security, NIH has instituted stringent procedures for entrance onto the NIH campus. All visitor vehicles, including taxicabs, hotel, and airport shuttles will be inspected before being allowed on campus. Visitors will be asked to show one form of identification (for example, a government-issued photo ID, driver's license, or passport) and to state the purpose of their visit.

Information is also available on the Institute's/Center's home page: NCAB: [deainfo.nci.nih.gov/advisory/ncab.htm](http://deainfo.nci.nih.gov/advisory/ncab.htm) where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS).

Dated: October 29, 2013.

**Melanie Gray,**

*Program Analyst, Office of Federal Advisory Committee Policy.*

[FR Doc. 2013-26253 Filed 11-1-13; 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 IITF 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.workplace.samhsa.gov>.

**FOR FURTHER INFORMATION CONTACT:** Giselle Hersh, Division of Workplace Programs, SAMHSA/CSAP, Room 7-1051, 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 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 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 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 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 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
- 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., 200 Precision Road, Suite 200, Horsham, PA 19044, 800-235-4890
- 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
- 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.)
- 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, 818-737-6370, (Formerly: SmithKline Beecham Clinical Laboratories)
- Redwood Toxicology Laboratory, 3650 Westwind Blvd., Santa Rosa, CA 95403, 707-570-4434
- 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
- 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
- U.S. Army Forensic Toxicology Drug Testing Laboratory, 2490 Wilson St., Fort George G. Meade, MD 20755-5235, 301-677-7085

\* 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.

**Janine Denis Cook,**

*Chemist, Division of Workplace Programs, Center for Substance Abuse Prevention, SAMHSA.*

[FR Doc. 2013-26276 Filed 11-1-13; 8:45 am]

**BILLING CODE 4160-20-P**

## DEPARTMENT OF HOMELAND SECURITY

[Docket No. DHS-2013-0028]

### Agency Information Collection Activities: Submission for Review; Information Collection Extension Request for the DHS S&T First Responders Community of Practice Program

**AGENCY:** Science and Technology Directorate, DHS.

**ACTION:** 30-day Notice and request for comment.

**SUMMARY:** The Department of Homeland Security (DHS) invites the general public to comment on the data collection form for the DHS Science & Technology (S&T) First Responders Community of Practice (FRCoP): User Registration Page (DHS Form 10059 (9/09)). The FRCoP web based tool collects profile information from first responders and select authorized non-first responder users to facilitate networking and formation of online communities. All users are required to authenticate prior to entering the site. In addition, the tool provides members the capability to create wikis, discussion threads, blogs, documents, etc., allowing them to enter and upload content in accordance with the site's Rules of Behavior. Members are able to participate in threaded discussions and comment on other member's content. The DHS S&T FRCoP program is responsible for providing a collaborative environment for the first responder community to share information, best practices, and lessons learned. Section 313 of the Homeland Security Act of

2002 (Pub. L. 107-296) established this requirement. This notice and request for comments is required by the Paperwork Reduction Act of 1995 (Pub. L. 104-13, 44 U.S.C. Chapter 35).

**DATES:** Comments are encouraged and will be accepted until December 4, 2013.

**ADDRESSES:** Interested persons are invited to submit comments, identified by docket number DHS-2013-0028, by one of the following methods:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Please follow the instructions for submitting comments.

- *Email:* [Kathy.Higgins@hq.dhs.gov](mailto:Kathy.Higgins@hq.dhs.gov). Please include docket number DHS-2013-0028 in the subject line of the message.

- *Fax:* (202) 254-6171. (Not a toll-free number).

- *Mail:* Science and Technology Directorate, ATTN: Chief Information Officer—Rick Stevens, 1120 Vermont Ave, Mail Stop 0202, Washington, DC 20005.

**FOR FURTHER INFORMATION CONTACT:** DHS FRCoP Contact Kathy Higgins (202) 254-2293 (Not a toll free number).

**SUPPLEMENTARY INFORMATION:** DHS S&T currently has approval to collect information utilizing the User Registration Form until September 30, 2012 with OMB approval number 1640-0016. The User Registration Form will be available on the First Responders Community of Practice Web site found at [<https://communities.firstresponder.gov/>]. The user will complete the form online and submit it through the Web site.

The Department is committed to improving its information collection and urges all interested parties to suggest how these materials can further reduce burden while seeking necessary information under the Paper Reduction Act.

DHS is particularly interested in comments that:

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

- (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

- (3) Suggest ways to enhance the quality, utility, and clarity of the information to be collected; and

- (4) Suggest ways to minimize the burden of the collection of information on those who are to respond, including

through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submissions of responses.

### Overview of This Information Collection

(1) *Type of Information Collection:* Renewal of Information Collection.

(2) *Title of the Form/Collection:* First Responders Community of Practice: User Registration Form.

(3) *Agency Form Number, if any, and the applicable component of the Department of Homeland Security sponsoring the collection:* DHS Science & Technology Directorate, R-Tech (RTD), DHS Form 10059 (09/09).

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Individuals; the data will be gathered from individual first responders who wish to participate in the First Responders Community of Practice.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:*

- a. *Estimate of the total number of respondents:* 2000.

- b. *An estimate of the time for an average respondent to respond:* 0.5 burden hours.

- c. *An estimate of the total public burden (in hours) associated with the collection:* 1000 burden hours.

Dated: September 27, 2013.

**Rick Stevens,**

*Chief Information Officer for Science and Technology.*

[FR Doc. 2013-26259 Filed 11-1-13; 8:45 am]

**BILLING CODE 9110-9F-P**

## DEPARTMENT OF HOMELAND SECURITY

### Office of the Secretary

#### Exercise of Authority Under the Immigration and Nationality Act

**AGENCY:** Office of the Secretary, DHS.

**ACTION:** Notice of determination.

**Authority:** 8 U.S.C. 1182(d)(3)(B)(i).

Following consultations with the Secretary of State and the Attorney General, I hereby conclude, as a matter of discretion in accordance with the authority granted to me by section 212(d)(3)(B)(i) of the Immigration and Nationality Act (INA), 8 U.S.C. 1182(d)(3)(B)(i), as amended, as well as the foreign policy and national security