DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center For Scientific Review; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the Surgery, Anesthesiology and Trauma Study Section, June 14, 2006, 1 p.m. to June 15, 2006, 3 p.m., Holiday Inn Select Bethesda, 8120 Wisconsin Ave., Bethesda, MD 20814 which was published in the Federal Register on May 3, 2006, 71 FR 26105-26106.

The meeting will be held at the DoubleTree Hotel, 8120 Wisconsin Avenue, Bethesda, MD 20814. The meeting dates and time remain the same. The meeting is closed to the public.


Anna Snouffer,
Acting Director, Office of Federal Advisory Committee Policy.

BILLING CODE 4140-01-M

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the Synthetic and Biological Chemistry B Study Section, June 8, 2006, 8:30 a.m. to June 9, 2006, 6 p.m., Holiday Inn Select Bethesda, 8120 Wisconsin Ave, Bethesda, MD 20814 which was published in the Federal Register on April 25, 2006, 71 FR 23929-23931.

The meeting will be held at the Double Tree Hotel, 8120 Wisconsin Avenue Bethesda, MD 20814. The meeting dates and time remain the same. The meeting is closed to the public.


Anna Snouffer,
Acting Director, Office of Federal Advisory Committee Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Current List of Laboratories 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 currently certified to meet the standards of Subpart C 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), on September 30, 1997 (62 FR 51118), and on April 13, 2004 (69 FR 19644).

A notice listing all currently certified laboratories is published in the Federal Register during the first week of each month. If any laboratory’s certification is suspended or revoked, the laboratory will be omitted from subsequent lists until such time as it is restored to full certification under the Mandatory Guidelines. If any laboratory 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://workplace.samhsa.gov and http://www.drugfreeworkplace.gov.

FOR FURTHER INFORMATION CONTACT: Mrs. Giselle Hersh or Dr. Walter Vogl, Division of Workplace Programs, SAMHSA/CSAP, Room 2–1035, 1 Choke Cherry Road, Rockville, Maryland 20857; 240–276–2600 (voice), 240–276–2610 (fax).

SUPPLEMENTARY INFORMATION: The Mandatory Guidelines were developed in accordance with Executive Order 12564 and section 503 of Public Law 100–71. Subpart C of the Mandatory Guidelines, “Certification of Laboratories Engaged in Urine Drug Testing for Federal Agencies,” sets strict standards that laboratories must meet in order to conduct drug and specimen validity tests on urine specimens for Federal agencies. To become certified, an applicant laboratory must undergo three rounds of performance testing plus an on-site inspection. To maintain that
certification, a laboratory must participate in a quarterly performance testing program plus undergo periodic, on-site inspections.

Laboratories which claim to be in the applicant stage of certification are not to be considered as meeting the minimum requirements described in the HHS Mandatory Guidelines. A laboratory must have its letter of certification from HHS/SAMHSA (formerly: HHS/NIDA) which attests that it has met minimum standards.

In accordance with Subpart C of the Mandatory Guidelines dated April 13, 2004 (69 FR 19644), the following laboratories meet the minimum standards to conduct drug and specimen validity tests on urine specimens:

- Baptist Medical Center-Toxicology Laboratory, 9601 1–630, Exit 7, Little Rock, AR 72205–7299, 501–202–2783, (Formerly: Forensic Toxicology Laboratory Baptist Medical Center)
- Doctors Laboratory, Inc., 2096 Julia Drive, Valdosta, GA 31602, 229–671–2281.
- DrugScan, Inc., P.O. Box 2969, 1119 Mears Road, Warmington, PA 18974, 215–674–9310.
- General Medical Laboratories, 36 South Brooks St., Madison, WI 53715, 608–267–6225.
- Kroll Laboratory Specialists, Inc., 1111 Newton St., Greta, LA 70053, 504–361–8089/800–433–3823. (Formerly: Laboratory Specialists, Inc.)
- Laboratory Corporation of America Holdings, 7207 N. Gessner Road, Houston, TX 77040, 713–856–8288/800–800–2387.
- Laboratory Corporation of America Holdings, 10788 Roselle St., San Diego, CA 92121, 800–882–7272, (Formerly: Poisonlab, Inc.).
- Laboratory Corporation of America Holdings, 550 17th Ave., Suite 300, Seattle, WA 98121, 206–923–7020/800–898–0180, (Formerly: DrugProof, Division of Dyncare/Laboratory of Pathology, LLC; Laboratory of Pathology of Seattle, Inc.; DrugProof Division of Laboratory of Pathology of Seattle, Inc.).
- 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)
- MAXXAM Analytics Inc.*, 6740 Campobello Road, Mississauga, ON, Canada L5N 2L8, 905–687–3244.
- Minnesota Veterans Affairs Medical Center, Forensic Toxicology Laboratory, 1 Veterans Drive, Minneapolis, MN 55417, 612–725–2088.
- One Source Toxicology Laboratory, Inc., 1213 Geona-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).
- Quest Diagnostics Incorporated, 4770 Regent Blvd., Irving, TX 75063, 800–824–6152, (Moved from the Dallas location on 03/31/01; Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories).
- Quest Diagnostics Incorporated, 10101 Renner Blvd., Lenexa, KS 66219, 913–888–3927/800–873–8845, (Formerly: LabOne, Inc.; Center for Laboratory Services, a Division of LabOne, Inc.).
- Quest Diagnostics Incorporated, 2282 South Presidents Drive, West Valley City, UT 84120, 801–606–6301/800–322–3361, (Formerly: Northwest Toxicology, a LabOne Company; LabOne, Inc., dba Northwest Toxicology; NWT Drug Testing, Northwest Toxicology, Inc.; Northwest Drug Testing, a division of NWT Inc.).
- South Bend Medical Foundation, Inc., 530 N. Lafayette Blvd., South Bend, IN 46601, 574–234–4176 x276.
- Sparrow Health System, Toxicology Testing Center, St. Lawrence Hospital & Pathology of Seattle, Inc.; UTMB Pathology-Toxicology Laboratory).
US Army Forensic Toxicology Drug Testing Laboratory, 2490 Wilson St., Fort George G. Meade, MD 20755–5235, 301–677–7085.

The following laboratory’s certification was suspended on November 14, 2005, with an effective date of November 15, 2005, and then revoked on February 8, 2006:

“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 13, 2004 (69 FR 19644). 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.

Anna Marsh,
Director, Office Program Services, SAMHSA.
[FR Doc. 06–4318 Filed 5–11–06; 8:45 am]
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INTERNATIONAL TRADE COMMISSION
[Inv. No. 337–TA–568]
In the Matter of Certain Products and Pharmaceutical Compositions Containing Recombinant Human Erythropoietin; Notice of Investigation
ACTION: Institution of investigation pursuant to 19 U.S.C. 1337.
The complainant requests that the Commission institute an investigation and, after the investigation, issue a permanent exclusion order and permanent cease and desist orders.
ADDRESSES: The amended complaint, except for any confidential information contained therein, is available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street, SW., Room 112, Washington, DC 20436, telephone 202–205–2000. Hearing impaired individuals are advised that information on this matter can be obtained by contacting the Commission’s TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its Internet server at http://www.usitc.gov. The public record for this investigation may be viewed on the Commission’s electronic docket (EDIS) at http://edis.usitc.gov.


Scope of Investigation: Having considered the complaint, the U.S. International Trade Commission, on May 8, 2006, ordered that—
(1) Pursuant to subsection (b) of section 337 of the Tariff Act of 1930, as amended, an investigation be instituted to determine whether there is a violation of subsection (a)(1)(B) of section 337 in the importation into the United States, the sale for importation, or the sale within the United States after importation of certain products and pharmaceutical compositions containing recombinant human