IV. Comments

Interested persons may submit to the Division of Dockets Management (see ADDRESSES) written or electronic notices of participation and comments for consideration at the hearing (see DATES). Submit a single copy of written or electronic notices of participation and comments, or two paper copies of any mailed notices of participation and comments, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.


Jeffrey Shuren,
Assistant Commissioner for Policy.

[FR Doc. 05–2094 Filed 1–31–05; 3:37 pm]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Committee on Infant Mortality; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), notice is hereby given of the following meeting:

Name: Advisory Committee on Infant Mortality (ACIM).

Dates and Times: March 1, 2005, 9 a.m.–5 p.m. March 2, 2005, 8:30 a.m.–3 p.m.

Place: Sheraton National Hotel, 900 South Orme Street, Arlington, Virginia 22204, (703) 521–1900.

Status: The meeting is open to the public with attendance limited to space availability.

Purpose: The Committee provides advice and recommendations to the Secretary of Health and Human Services on the following: Department programs that are directed at reducing infant mortality and improving the health status of pregnant women and infants; factors affecting the continuum of care with respect to maternal and child health care, including outcomes following childbirth; strategies to coordinate the variety of Federal, State, local and private programs and efforts that are designed to deal with the health and social problems impacting on infant mortality; and the implementation of the Healthy Start program and Healthy People 2010 infant mortality objectives.

Agenda: Topics that will be discussed include the following: Improving Perinatal Data; Neonatal Intensive Care and Ethical Issues; and Provider Reimbursement Issues. Substantial time will be spent in small group and full Committee discussions aimed at formulating the ACIM issues agenda. Proposed agenda items are subject to change as priorities indicate.

Time will be provided for public comments limited to five minutes each; comments are to be submitted no later than February 15, 2005.

FOR FURTHER INFORMATION CONTACT:
Anyone requiring information regarding the Committee should contact Peter C. van Dyck, M.D., M.P.H., Executive Secretary, ACIM, Health Resources and Services Administration (HRSA), Room 18–05, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857, Telephone: (301) 443–2170.

Individuals who are submitting public comments or who have questions regarding the meeting should contact Ann M. Koonz, C.N.M., Dr. P.H., HRSA, Maternal and Child Health Bureau, telephone: (301) 443–6327, e-mail: akoontz@hrsa.gov.


Steven A. Pelovitz,
Associate Administrator for Administration and Financial Management.

[FR Doc. 05–2102 Filed 2–2–05; 8:45 am]

BILLING CODE 4165–15–P

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, that 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 Pub. L. 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: ACL Laboratories, 8901 W. Lincoln Ave., West Allis, WI 53227, 414–319–7840 / 800–877–7016 (Formerly: Bayshore Clinical Laboratory).


Baptist Medical Center-Toxicology Laboratory, 9601 I–630, Exit 7, Little Rock, AR 72205–7299, 501–202–2783 (Formerly: Forensic Science Laboratory Baptist Medical Center).


General Medical Laboratories, 36 South Brooks St., Madison, WI 53715, 608–287–6225.

Kroll Laboratory Specialists, Inc., 1111 Newton St., Gretna, LA 70053, 504–361–8989 / 800–433–3823 (Formerly: Laboratory Specialists, Inc.).

LabOne, Inc., 10101 Renner Blvd., Lenexa, KS 66219, 913–888–3927 / 800–873–8845 (Formerly: Center for Laboratory Services, a Division of LabOne, Inc.).


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 Dr., Research Triangle Park, NC 27709, 919–572–6990 / 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, 10788 Roselle St., San Diego, CA 92121, 800–862–7272 (Formerly: Poisonlab, 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).

Marshfield Laboratories, Forensic Toxicology Laboratory, 1000 North Oak Ave., Marshfield, WI 54449, 715–389–3734 / 800–331–3734.

MAXXAM Analytics Inc.*, 6740 Campobello Road, Mississauga, ON, Canada L5N 2L8, 905–817–5700 (Formerly: NOVAMANN (Ontario) Inc.).


MetroLab-Legacy Laboratory Services, 1225 NE 2nd Ave., Portland, OR 97232, 503–413–5295 / 800–950–5295.

Minnesota Veterans Affairs Medical Center, Forensic Toxicology Laboratory, 1 Veterans Dr., Minneapolis, MN 55417, 612–725–2088.

Northwest Toxicology Laboratories, a LabOne Company, 2282 South Presidents Drive, Suite C, West Valley City, UT 84120, 801–293–2300 / 800–322–3361 (Formerly: LabOne, Inc., d/b/a Northwest Toxicology; NWT Drug Testing, Northwest Toxicology, Inc.; Northwest Drug Testing, a division of NWT Inc.).

One Source Toxicology Laboratory, Inc., 1213 Genoa-Red Bluff, Pasadena, TX 77504, 887–347–3774 (Formerly: University of Texas Medical Branch, Clinical Chemistry Division; UTMB Pathology-Toxicology Laboratory).

Oregon Medical Laboratories, P.O. Box 972, 722 East 11th Ave., Eugene, OR 97440–0972, 541–687–2134.

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).


DEPARTMENT OF HOMELAND SECURITY

Information Analysis and Infrastructure Protection; Telecommunications Service Priority System Oversight Committee

AGENCY: National Communications System (NCS), Department of Homeland Security.

ACTION: Committee management; notice of advisory committee renewal.

SUMMARY: The Department of Homeland Security (DHS) has renewed the charter for the Telecommunications Service Priority (TSP) System Oversight Committee.

SUPPLEMENTARY INFORMATION: Pursuant to the Federal Advisory Committee Act, Public Law 92–463, as amended (5 U.S.C. App. 2, et seq.) the Secretary of Homeland Security has renewed the charter for the TSP System Oversight Committee. This renewal follows consultation with the Committee Management Secretariat, General Services Administration and has been determined by the Secretary to be in the public interest in connection with the performance of duties imposed on DHS by law.

The TSP System Oversight Committee identifies and reviews any problems developing in the TSP System and recommends actions to correct them or prevent recurrence. The TSP System Oversight Committee Designated Federal Officer is Lt. Col. Joanne Sechrest, USAF.

FOR FURTHER INFORMATION CONTACT: Susan Flint, NCS Office of Priority Telecommunications, 703–607–4932. Media or press should contact Mr. Steve Barrett at 703–607–6211.

Peter M. Fonash, Acting Deputy Manager, National Communications System. [FR Doc. 05–2093 Filed 2–2–05; 8:45 am]

BILLING CODE 4160–20–P

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

[USCG–2004–19977]

Inspection of Towing Vessels

AGENCY: Coast Guard, DHS.

ACTION: Notice; request for comments, and notice of public meeting; change of location.

SUMMARY: The location of the upcoming public meeting being held in New Orleans, Louisiana, is changed. Instead of the Hale Boggs Federal Building, as previously announced in the Federal Register, the meeting will take place at the Hyatt Regency New Orleans. The date of the meeting, February 10, and the hours, from 9 a.m. to 12 p.m. remain the same. In the recently enacted Coast Guard and Maritime Transportation Act of 2004, the Congress directed the Coast Guard to add towing vessels to the list of vessels subject to inspections, and to consider the establishment of a safety management system appropriate for towing vessels. Through public meetings, we are seeking public and industry involvement as we consider how to proceed.

DATES: Comments and related material must reach the Docket Management Facility on or before March 23, 2005. A public meeting will be held on February 10, 2005, in New Orleans, LA. Meetings in Oakland, CA, and St. Louis, MO, remain unchanged as previously announced in the Federal Register [69 FR 78471].

ADDRESSES: Comments. You may submit comments identified by Coast Guard docket number USCG–2004–19977 to the Docket Management Facility at the U.S. Department of Transportation. To avoid duplication, please use only one of the following methods:

(3) Fax: 202–493–2251.
(4) Delivery: Room PL–401 on the Plaza level of the Nassif Building, 400 Seventh Street SW., Washington, DC, between 9 a.m. and 5 p.m., Monday through Friday, except Federal holidays. The telephone number is 202–366–9329.


Meeting. The meeting in New Orleans will be held at the following location: Hyatt Regency New Orleans, Cabildo Room, Poydras at Loyola Avenue, New Orleans, LA 70113.