DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Arthritis and Musculoskeletal and Skin Diseases; Notice of Closed 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 a meeting of the Division of Intramural Research Board of Scientific Counselors, NIAID. 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. The grant applications and the discussions 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: Division of Intramural Research Board of Scientific Counselors, NIAID.

Date: June 1–3, 2009.

Time: June 1, 2009, 8 a.m. to 6:15 p.m.

Agenda: To review and evaluate personal qualifications and performance, and competence of individual investigators.

Place: National Institutes of Health, Building 50, 50 Center Drive, Conference Rooms 1227/1233, Bethesda, MD 20892.

Time: June 2, 2009, 7:30 a.m. to 7:05 p.m.

Agenda: To review and evaluate personal qualifications and performance, and competence of individual investigators.

Place: National Institutes of Health, Building 50, 50 Center Drive, Conference Rooms 1227/1233, Bethesda, MD 20892.

Time: June 3, 2009, 7:30 a.m. to 12:45 p.m.

Agenda: To review and evaluate personal qualifications and performance, and competence of individual investigators.

Place: National Institutes of Health, Building 50, 50 Center Drive, Conference Rooms 1227/1233, Bethesda, MD 20892.

Contact Person: Kathryn C. Zoon, PhD., Director, Division of Intramural Research, National Institute of Allergy and Infectious Diseases, NIH, Building 31, Room 4A30, Bethesda, MD 20892, 301–496–3006, kzoorn@niaid.nih.gov.

Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS


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

[FR Doc. E9–5137 Filed 3–10–09; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

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BILLING CODE 4140–01–P
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://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 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 I–630, Exit 7, Little Rock, AR 72205–7299. 501–202–2783. (Formerly: Forensic Toxicology Laboratory Baptist Medical Center).
- Clendo Reference Laboratory, Avenue Santa Cruz #58, Bayamon, Puerto Rico 00959. 787–620–9095.
- Diagnostic Services, Inc., dba DSI, 12700 Westlinks Drive, Fort Myers, FL 33913. 239–561–8200/800–735–5416.
- Doctors Laboratory, Inc., 2906 Julia Drive, Valdosta, GA 31602. 229–671–2281.
- DrugScan, Inc., P.O. Box 2969, 1119 Mears Road, Warminster, PA 18974. 215–674–9310.
- Kroll Laboratory Specialists, Inc., 1111 Newton St., Cretin, LA 70053. 504–361–8989/800–433–3823. (Formerly: Laboratory Specialists, Inc.).
- Laboratory Corporation of America Holdings, 7207 N. Gossner Road, Houston, TX 77040. 713–856–8288/800–800–1946.
- Laboratory Corporation of America Holdings, 69 First Ave., Raritan, NJ 08869. 909–526–2400/800–437–4986. (Formerly: Roche Biomedical Laboratories, Inc.).
- Laboratory Corporation of America Holdings, 1120 Main Street, Southaven, MS 38671. 866–827–8042/800–233–6399. (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 2L8. 905–817–5700. (Formerly: Maxxam Analytics Inc., NOVAMANN (Ontario), Inc.).
- Minneapolis Veterans Affairs Medical Center, Forensic Toxicology Laboratory, 11 Veterans Drive, Minneapolis, MN 55417. 612–725–2088.
- 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).
- Phamatech, Inc., 1015 Barnes Canyon Road, San Diego, CA 92121. 858–643–5553.
- Quest Diagnostics Incorporated, 3175 Presidential Dr., Atlanta, GA 30340. 770–452–1500/800–729–6432. (Formerly: SmithKline Beecham)
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.

Elaine Parry,
Director, Office of Program Services, SAMHSA.

| BILLING CODE 4160–20–P |

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency


Oklahoma; Amendment No. 1 to Notice of a Major Disaster Declaration

AGENCY: Federal Emergency Management Agency, DHS.

ACTION: Notice.

SUMMARY: This notice amends the notice of a major disaster declaration for the State of Oklahoma (FEMA–1823–DR), dated February 17, 2009, and related determinations.

DATES: Effective Date: February 25, 2009.


SUPPLEMENTARY INFORMATION: Notice is hereby given that the incident period for this emergency is closed effective February 8, 2009.

The following Catalog of Federal Domestic Assistance Numbers (CFDA) are to be used for reporting and drawing funds: 97.030, Community Disaster Loans; 97.031, Cora Brown Fund; 97.032, Crisis Counseling; 97.033, Disaster Legal Services; 97.034, Disaster Unemployment Assistance (DUA); 97.046, Fire Management Assistance Grant; 97.048, Disaster Housing Assistance to Individuals and Households In Presidentially Declared Disaster Areas; 97.049, Presidentially Declared Disaster Assistance—Disaster Housing Operations for Individuals and Households; 97.050, Presidentially Declared Disaster Assistance to Individuals and Households—Other Needs; 97.056, Disaster Grants—Public Assistance (Presidentially Declared Disasters); 97.039, Hazard Mitigation Grant.)

Nancy Ward,
Acting Administrator, Federal Emergency Management Agency.

[FR Doc. E9–5153 Filed 3–10–09; 8:45 am]