DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention (CDC)

Advisory Board on Radiation and Worker Health: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Advisory Board on Radiation and Worker Health, Department of Health and Human Services, has been renewed for a 2-year period through August 3, 2011.

For information, contact Mr. Theodore Katz, Executive Secretary, Advisory Board on Radiation and Worker Health, Department of Health and Human Services, 1600 Clifton Road, M/S E20, Atlanta, Georgia, 30341, telephone 404/498–2533, or fax 404/498–2570.

For further information contact: The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities, for both the CDC and the Agency for Toxic Substances and Disease Registry.


Elaine L. Baker,
Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Katharine Kripke, Assistant Director, Vaccine Research Program, Division of AIDS, NIAID, NIH, 6700B Rockledge Dr., Bethesda, MD 20892–7628, or call non-toll-free number 301–402–0846, or E-mail your request, including your address to NIAIDsurvey@NIH.gov.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.


J. J. McGowan,
Executive Officer, NIAID, National Institutes of Health.

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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 1644). 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: For information on the NLCP, contact 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).

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Mandatory Guidelines were developed in accordance with Executive Order 12554 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 1644), the following laboratories meet the minimum standards to conduct drug and specimen validity tests on urine specimens:

ACI Laboratories, 8901 W. Lincoln Ave., West Allis, WI 53227. 414–328–7840/800–877–2075. (Formerly: Bayshore Clinical Laboratory.)


Aegis Analytical Laboratories, 345 Hill Ave., Nashville, TN 37210. 615–255–2400. (Formerly: Aegis Sciences Corporation, Aegis Analytical Laboratories, Inc.)

Baptist Medical Center–Toxicology Laboratory, 9601 I–465, 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. 767–820–9095.

Clinical Reference Lab, 8433 Quivira Road, Lenexa, KS 66215–2802. 800–445–6917. (Formerly: Doctors Laboratory, Inc. 2906 Julia Drive, Valdosta, GA 31602. 229–671–2281.

DrugScan, Inc., P.O. Box 2969, 1119 Mearns Road, Warminster, PA 18974. 215–474–9310.


Ellisby Laboratories, Inc., 5 Industrial Park Drive, Oxford, MS 38655. 662–236–2900. (Formerly: Gamma-Dynacare Medical Laboratories, Inc.)

ElSohy Laboratories, Inc., 5 Industrial Park Drive, Oxford, MS 38655. 662–236–2900. (Formerly: Gamma-Dynacare Medical Laboratories, Inc.)

Grosky Laboratory Specialists, Inc., 1111 Newton St., Gretna, NE 68053. 402–361–
DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, Coordinating Center for Health Promotion (BSC, CCHP or the BSC)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting of the aforementioned committee:

**Time and Date:** 1 p.m.–5 p.m., September 15, 2009.

The call may end before 5 p.m. if business is completed. Someone will remain on the line until that time to notify callers that the call is ended and business complete. **Place:** Teleconference originating from CDC, 1825 Century Boulevard, NE., Room 4066, Atlanta, Georgia 30345. **Call-in number:** (800) 779–9076. **Participant pass code:** 39780.

If you have a problem in accessing the call, call (404) 498–6700.

**Status:** This meeting is open to the public via the conference line above which will accommodate approximately 100 callers.

**Purpose:** This BSC is charged with providing advice and guidance to the Secretary of Health and Human Services, the Director of CDC, and the Director of CCHP concerning strategies and goals for the programs and research within the National Center on Birth Defects and Developmental Disabilities and the National Center for Chronic Disease Prevention and Health Promotion.

**Matters To Be Discussed:** The agenda will include a continuation of the discussion and finalization of recommendations to CDC leadership as prescribed in the Coordinating Center for Health Promotion Charter. Those recommendations relate to strategic planning for the National Center on Birth Defects and Developmental Disabilities as well as the results of the BSC’s review of the National Center for Chronic Disease Prevention and Health Promotion as an organizational unit at CDC.

**Providing Oral or Written Comments:** It is the policy of the BSC, CCHP to provide a brief period for oral public comments. In general, each individual or group requesting to make an oral presentation will be limited to a total time of five minutes, if time permits.

**Contact Person for Additional Information:** Karen Steinberg, PhD, Senior Science Officer, Coordinating Center for Health Promotion, CDC, 4770 Buford Highway, NE., Mailstop E–70, Atlanta, Georgia 30341; telephone (404) 498–6700; fax (404) 498–6880; or e-mail at Karen.Steinberg@cchp.dhhs.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register Notices pertaining to announcements of meetings and other committee management activities, for CDC and the Agency for Toxic Substances and Disease Registry.