

funds. Funds generated by grant-supported efforts are considered "project income," and may not be used for matching purposes.

It is the policy of this program to reject all requests for indirect or overhead costs as well as "in-kind" match contributions. In addition, grant funds must not be used to supplant private or local/state government funds currently spent for committee purposes. Funding requests from existing committees should focus entirely on the costs associated with the expansion efforts. Also, under no circumstances may business or labor officials participating on a labor-management committee be compensated out of grant funds for *time* spent at committee meetings or *time* spent in committee training sessions. Applicants generally will not be allowed to claim all or a portion of *existing* full-time staff as an expense or match contribution. For a more complete discussion of cost allowability, applicants are encouraged to consult the FY2005 FMCS Financial and Administrative Grants Manual, which will be included in the application kit.

G. Application Submission and Review Process

The Application for Federal Assistance (SF-424) form must be signed by *both* a labor and management representative. In lieu of signing the SF-424 form, representatives may be type their name, title, and organization on plain bond paper with a signature line signed and dated, in accordance with block 18 of the SF-424 form. We will accept applications beginning May 15, 2005 and continue to do so until all FY 2005 grant funds have been obligated, with awards being made by September 30, 2006. While proposals may be accepted at any time between May 15, 2005 and September 30, 2006, proposals received late in the cycle have a greater risk of not being funded due to unavailability of funds. Offerors are highly advised to contact the grants director prior to committing any resources to the preparation of a proposal. An original application containing numbered pages, plus *three* copies, should be addressed to the Federal Mediation and Conciliation Service, Labor-Management Grants Program, 2100 K Street, NW., Washington, DC 20427. FMCS will not consider videotaped submissions or video attachments to submissions. FMCS will confirm receipt of all applications within 10 days thereof.

All eligible applications will be reviewed and scored preliminarily by one or more Grant Review Boards. The

Board(s) will recommend selected applications for rejection or further funding consideration. The Director, Labor-Management Grants Program, will finalize the scoring and selection process. The individual listed as contact person in Item 6 on the application form will generally be the only person with whom FMCS will communicate during the application review process. Please be sure that person is available once the application has been submitted.

All FY2005 grant applicants will be notified of results and all grant awards will be made September 30, 2006. Applications that fail to adhere to eligibility or other major requirements will be administratively rejected by the Director, Labor-Management Grants Program.

H. Contact

Individuals wishing to apply for funding under this program should contact the Federal Mediation and Conciliation Service as soon as possible to obtain an application kit. Please consult the FMCS Web site (<http://www.fmcs.gov>) to download forms and information.

These kits and additional information or clarification can be obtained free of charge by contacting the Federal Mediation and Conciliation Service, Labor-Management Grants Program, 2100 K Street, NW., Washington, DC 20427 at (202) 606-8181, (202) 271-8868 or jlrorber@fmcs.gov.

Fran Leonard,

Director, Budget and Finance, Federal Mediation and Conciliation Service.

[FR Doc. 05-4292 Filed 3-4-05; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Committee on Vital and Health Statistics: Meeting

Pursuant to the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) announces the following advisory committee meeting.

Name: National Committee on Vital and Health Statistics (NCVHS).

Time and Date: March 3, 2005, 9 a.m.-3 p.m., March 4, 2005, 10 a.m.-3:15 p.m.

Place: Hubert H. Humphrey Building, 200 Independence Avenue SW., Room 505A, Washington, DC 20201.

Status: Open.

Purpose: At this meeting the Committee will hear presentations and hold discussions on several health data policy topics. On the morning of the first day the Committee will hear updates and status reports from the

Department on topics including Clinical Data Standards, the Consolidated Health Informatics Initiative, and the HIPAA Privacy Rule compliance. In the afternoon the Committee will hear an update on the National Health Information Infrastructure and will discuss various materials prepared by NCVHS Subcommittees.

On the second day the Committee will hear an updated from the Board of Scientific Counselors at the National Center for Health Statistics (NCHS) and reports on two HHS initiatives in geocoding. The Committee will also discuss plans for its annual report to Congress and there will be reports from the Subcommittees and a discussion of agendas for future Committee meetings.

The times shown above are for the full committee meeting. Subcommittee breakout sessions are scheduled for late in the afternoon of the first day and in the morning prior to the full Committee meeting on the second day. Agendas for these breakout sessions will be posted on the NCVHS Web site (URL below) when available.

Contact Person for More Information: Substantive program information as well as summaries of meetings and a roster of committee members may be obtained from Marjorie S. Greenberg, Executive Secretary, NCVHS, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Room 2402, Hyattsville, Maryland 20782, telephone (301) 458-4245. Information also is available on the NCVHS home page of the HHS Web site: <http://www.ncvhs.hhs.gov/>, where further information including an agenda will be posted when available.

Should you require reasonable accommodation, please contact the CDC Office of Equal Employment Opportunity on (301) 458-4EEO (4336) as soon as possible.

Dated: March 1, 2005.

James Scanlon,

Acting Deputy Assistant Secretary for Science and Data Policy, Office of the Assistant Secretary for Planning and Evaluation.

[FR Doc. 05-4311 Filed 3-4-05; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Funding Opportunity Number: RFA 05045]

Academic Partners Public Health Training Grant; Notice of Availability of Funds—Amendment

A notice announcing the availability of fiscal year (FY) 2005 funds for a grant to: (a) provide trainees the opportunity to learn about broad, cross-cutting public health policy and program development at the Federal, state and local government level and (b) make progress toward achieving the prevention objectives of Healthy People 2010 was published in the **Federal**

Register on February 15, 2005, Vol. 70, No. 30, pages 7739–7744.

The notice is amended as follows: On page 7740, Column 1, First line: change to “Application Deadline: April 1, 2005.” On page 7743, Column 1, Section 1V.5, please include a third bullet that states: Indirect costs cannot exceed 8% of the total direct cost.

Dated: March 1, 2005.

William P. Nichols,

*Director, Procurement and Grants Office,
Centers for Disease Control and Prevention.*

[FR Doc. 05–4315 Filed 3–4–05; 8:45 am]

BILLING CODE 4163–18–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, 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 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, 414–328–7840/800–877–7016 (Formerly: Bayshore Clinical Laboratory)

ACM Medical Laboratory, Inc., 160 Elmgrove Park, Rochester, NY 14624, 585–429–2264

Advanced Toxicology Network, 3560 Air Center Cove, Suite 101, Memphis, TN 38118, 901–794–5770/888–290–1150

Aegis Analytical Laboratories, Inc., 345 Hill Ave., Nashville, TN 37210, 615–255–2400

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) Clinical Reference Lab, 8433 Quivira Rd., Lenexa, KS 66215–2802, 800–445–6917

Diagnostic Services Inc., dba DSI, 12700 Westlinks Dr., Fort Myers, FL 33913, 239–561–8200/800–735–5416

Doctors Laboratory, Inc., 2906 Julia Drive, Valdosta, GA 31602, 229–671–2281

DrugProof, Division of Dynacare/Laboratory of Pathology, LLC, 1229 Madison St., Suite 500, Nordstrom Medical Tower, Seattle, WA 98104,

206–386–2661/800–898–0180 (Formerly: Laboratory of Pathology of Seattle, Inc., DrugProof, Division of Laboratory of Pathology of Seattle, Inc.)

DrugScan, Inc., P.O. Box 2969, 1119 Mearns Rd., Warminster, PA 18974, 215–674–9310

Dynacare Kasper Medical Laboratories,* 10150–102 St., Suite 200, Edmonton, Alberta, Canada T5J 5E2, 780–451–3702/800–661–9876

ElSohly Laboratories, Inc., 5 Industrial Park Dr., Oxford, MS 38655, 662–236–2609

Express Analytical Labs, 3405 7th Ave., Suite 106, Marion, IA 52302, 319–377–0500

General Medical Laboratories, 36 South Brooks St., Madison, WI 53715, 608–267–6225

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, 7207 N. Gessner Rd., 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 Dr., 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, 10788 Roselle St., San Diego, CA 92121, 800–882–7272 (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,