

increased mental health capacity at Hch sites; less redundancy in data collection;

and enhanced screening for multi-dimensional issues.

The estimated response burden for this project is as follows:

Instrument	Number of responses	Responses/respondent	Burden/response (Hrs.)	Total burden hours
Administrative Interviews .....	24	3	1.5	108
Evaluator Interviews .....	12	3	1.0	36
Line Staff Interviews .....	48	3	1.0	144
Consumer Focus Groups .....	84	3	1.0	252
Other Key Informants .....	48	3	1.0	144
Activity Logs .....	12	10	2.0	240
Total .....	228	.....	.....	924
3-yr. Annual Average .....	228	.....	.....	308

A total of approximately 6,500 program participants are expected to be recruited from the 12 sites. Each site will collect GPRA data on these participants using the CMHS GPRA Core Client Outcome measures approved by the Office of Management and Budget under control number 0930-0208, which cover such domains as drug and alcohol use, family and living conditions, education, employment, and income, crime and criminal justice status, and mental and physical health problems and treatment. To obtain information on client-level outcomes the central Evaluation Center will work with each site to develop methods for obtaining relevant material from the GPRA data. It is expected that client-level data will be submitted to the Evaluation Center via electronic means. The Evaluation Center will provide training and technical assistance to all sites on data submission procedures.

Send comments to Nancy Pearce, SAMHSA Reports Clearance Officer, Room 16-105, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: January 27, 2003.

**Richard Kopanda,**

*Executive Officer, Substance Abuse and Mental Health Services Administration.*

[FR Doc. 03-2392 Filed 1-31-03; 8:45 am]

BILLING CODE 4162-20-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 notifies Federal agencies of the laboratories currently certified to meet standards of Subpart C of Mandatory Guidelines for Federal Workplace Drug Testing Programs (59 FR 29916, 29925). 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 Guidelines.

If any laboratory has withdrawn from the National Laboratory Certification Program 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 the following websites: <http://workplace.samhsa.gov> and <http://www.drugfreeworkplace.gov>.

**FOR FURTHER INFORMATION CONTACT:** Mrs. Giselle Hersch or Dr. Walter Vogl, Division of Workplace Programs, 5600 Fishers Lane, Rockwall 2 Building, Room 815, Rockville, Maryland 20857; Tel.: (301) 443-6014, Fax: (301) 443-3031.

#### **SUPPLEMENTARY INFORMATION:**

Mandatory Guidelines for Federal Workplace Drug Testing were developed in accordance with Executive Order 12564 and section 503 of Public Law 100-71. Subpart C of the Guidelines, "Certification of Laboratories Engaged in Urine Drug Testing for Federal Agencies," sets strict standards which laboratories must meet in order to conduct urine drug testing 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 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 expressed in the HHS Guidelines. A laboratory must have its letter of certification from SAMHSA, HHS (formerly: HHS/NIDA) which attests that it has met minimum standards.

In accordance with Subpart C of the Guidelines, the following laboratories meet the minimum standards set forth in the Guidelines:

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.

Alliance Laboratory Services, 3200 Burnet Ave., Cincinnati, OH 45229, 513-585-6870, (Formerly: Jewish Hospital of Cincinnati, Inc.).

Associated Pathologists Laboratories, Inc., 4230 South Burnham Ave., Suite 250, Las Vegas, NV 89119-5412, 702-733-7866 / 800-433-2750.

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.

Cox Health Systems, Department of Toxicology, 1423 North Jefferson Ave., Springfield, MO 65802, 800-876-3652 / 417-269-3093, (Formerly: Cox Medical Centers).

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