

Commenters may also mail them to:  
Office of Management and Budget,  
Office of Information and Regulatory  
Affairs, New Executive Office Building,  
Room 10102, Washington, DC 20503.

**Rose Shannon,**

*Director, Division of Executive  
Correspondence.*

[FR Doc. 2011-25374 Filed 9-30-11; 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 and Instrumented Initial Testing Facilities 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 and  
Instrumented Initial Testing Facilities  
(IITF) currently certified to meet the  
standards 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);  
September 30, 1997 (62 FR 51118);  
April 13, 2004 (69 FR 19644); November  
25, 2008 (73 FR 71858); December 10,  
2008 (73 FR 75122); and on April 30,  
2010 (75 FR 22809).

A notice listing all currently certified  
Laboratories and Instrumented Initial  
Testing Facilities (IITF) is published in  
the **Federal Register** during the first  
week of each month. If any Laboratory/  
IITF's certification is suspended or  
revoked, the Laboratory/IITF will be  
omitted from subsequent lists until such  
time as it is restored to full certification  
under the Mandatory Guidelines.

If any Laboratory/IITF has withdrawn  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](http://www.workplace.samhsa.gov) and [http://  
www.drugfreeworkplace.gov](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 initially  
developed in accordance with Executive  
Order 12564 and section 503 of Public  
Law 100-71. The "Mandatory  
Guidelines for Federal Workplace Drug  
Testing Programs," as amended in the  
revisions listed above, requires {or set}  
strict standards that Laboratories and  
Instrumented Initial Testing Facilities  
(IITF) must meet in order to conduct  
drug and specimen validity tests on  
urine specimens for Federal agencies.

To become certified, an applicant  
Laboratory/IITF must undergo three  
rounds of performance testing plus an  
on-site inspection. To maintain that  
certification, a Laboratory/IITF must  
participate in a quarterly performance  
testing program plus undergo periodic,  
on-site inspections.

Laboratories and Instrumented Initial  
Testing Facilities (IITF) in the applicant  
stage of certification are not to be  
considered as meeting the minimum  
requirements described in the HHS  
Mandatory Guidelines. A Laboratory/  
IITF must have its letter of certification  
from HHS/SAMHSA (formerly: HHS/  
NIDA) which attests that it has met  
minimum standards.

In accordance with the Mandatory  
Guidelines dated November 25, 2008  
(73 FR 71858), the following  
Laboratories and Instrumented Initial  
Testing Facilities (IITF) meet the  
minimum standards to conduct drug  
and specimen validity tests on urine  
specimens:

#### Instrumented Initial Testing Facilities (IITF)

None.

#### Laboratories

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

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

Alere Toxicology Services, 450  
Southlake Blvd., Richmond, VA  
23236, 804-378-9130. (Formerly:  
Kroll Laboratory Specialists, Inc.,  
Scientific Testing Laboratories, Inc.;  
Kroll Scientific Testing Laboratories,  
Inc.)

Baptist Medical Center-Toxicology  
Laboratory, 11401 I-30, Little Rock,  
AR 72209-7056, 501-202-2783.  
(Formerly: Forensic Toxicology  
Laboratory Baptist Medical Center.)  
Clinical Reference Lab, 8433 Quivira  
Road, Lenexa, KS 66215-2802, 800-  
445-6917.

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

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

ElSohly Laboratories, Inc., 5 Industrial  
Park Drive, Oxford, MS 38655, 662-  
236-2609.

Gamma-Dynacare Medical  
Laboratories,\* A Division of the  
Gamma-Dynacare Laboratory  
Partnership, 245 Pall Mall Street,  
London, ONT, Canada N6A 1P4, 519-  
679-1630.

Laboratory Corporation of America  
Holdings, 7207 N. Gessner Road,  
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 Drive,  
Research Triangle Park, NC 27709,  
919-572-6900/800-833-3984.  
(Formerly: LabCorp Occupational  
Testing Service 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, 1120 Main Street,  
Southaven, MS 38671, 866-827-8042/  
800-233-6339. (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.)

MedTox Laboratories, Inc., 402 W. County Road D, St. Paul, MN 55112, 651-636-7466/800-832-3244.

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

Minneapolis Veterans Affairs Medical Center, Forensic Toxicology Laboratory, 1 Veterans Drive, Minneapolis, MN 55417, 612-725-2088.

National Toxicology Laboratories, Inc., 1100 California Ave., Bakersfield, CA 93304, 661-322-4250/800-350-3515.

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.)

Pathology Associates Medical Laboratories, 110 West Cliff Dr., Spokane, WA 99204, 509-755-8991/800-541-7891 x7.

Phamatech, Inc., 10151 Barnes Canyon Road, San Diego, CA 92121, 858-643-5555.

Quest Diagnostics Incorporated, 1777 Montreal Circle, Tucker, GA 30084, 800-729-6432. (Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories.)

Quest Diagnostics Incorporated, 400 Egypt Road, Norristown, PA 19403, 610-631-4600/877-642-2216. (Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories.)

Quest Diagnostics Incorporated, 8401 Fallbrook Ave., West Hills, CA 91304, 800-877-2520. (Formerly: SmithKline Beecham Clinical Laboratories.)

S.E.D. Medical Laboratories, 5601 Office Blvd., Albuquerque, NM 87109, 505-727-6300/800-999-5227.

South Bend Medical Foundation, Inc., 530 N. Lafayette Blvd., South Bend, IN 46601, 574-234-4176 x1276.

Southwest Laboratories, 4625 E. Cotton Center Boulevard, Suite 177, Phoenix, AZ 85040, 602-438-8507/800-279-0027.

St. Anthony Hospital Toxicology Laboratory, 1000 N. Lee St., Oklahoma City, OK 73101, 405-272-7052.

STERLING Reference Laboratories, 2617 East L Street, Tacoma, Washington 98421, 800-442-0438.

Toxicology & Drug Monitoring Laboratory, University of Missouri Hospital & Clinics, 301 Business Loop 70 West, Suite 208, Columbia, MO 65203, 573-882-1273.

Toxicology Testing Service, Inc., 5426 N.W. 79th Ave., Miami, FL 33166, 305-593-2260.

U.S. Army Forensic Toxicology Drug Testing Laboratory, 2490 Wilson St., Fort George G. Meade, MD 20755-5235, 301-677-7085.

\* The Standards Council of Canada (SCC) voted to end its Laboratory Accreditation Program for Substance Abuse (LAPSA) effective May 12, 1998. Laboratories certified through that program were accredited to conduct forensic urine drug testing as required by U.S. Department of Transportation (DOT) regulations. As of that date, the certification of those accredited Canadian laboratories will continue under DOT authority. The responsibility for conducting quarterly performance testing plus periodic on-site inspections of those LAPSA-accredited laboratories was transferred to the U.S. HHS, with the HHS' NLCP contractor continuing to have an active role in the performance testing and laboratory inspection processes. Other Canadian laboratories wishing to be considered for the NLCP may apply directly to the NLCP contractor just as U.S. laboratories do.

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 30, 2010 (75 FR 22809). 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.

Dated: September 21, 2011.

**Elaine Parry,**

*Director, Office of Management, Technology, and Operations, SAMHSA.*

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**BILLING CODE 4160-20-P**

## DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR-5481-N-13]

### Notice of Proposed Information Collection: Comment Request Self-Help Homeownership Opportunity Program (SHOP)

**AGENCY:** Office of the Assistant Secretary for Community Planning and Development, HUD.

**ACTION:** Notice of proposed information collection.

**SUMMARY:** The proposed information collection requirement described below will be submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

**DATES:** *Comment Due Date:* December 2, 2011.

**ADDRESSES:** Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB Control Number and should be sent to: Rudene Thomas, Reports Liaison Officer, U.S. Department of Housing and Urban Development, 451 Seventh Street, SW., Room 7233, Washington, DC 20410-4500.

**FOR FURTHER INFORMATION CONTACT:** Ginger Macomber, SHOP Program Manager, Office of Affordable Housing Programs, U.S. Department of Housing and Urban Development, 451 Seventh Street, SW., Room 7162, Washington, DC 20410-4500; telephone 202-402-4605 (this is not a toll-free number) or by e-mail at [ginger.macomber@hud.gov](mailto:ginger.macomber@hud.gov).

**SUPPLEMENTARY INFORMATION:** The Department is submitting the proposed information collection to OMB for review, as required by the Paperwork Reduction Act of 1995 (44 U.S.C. chapter 35, as amended).

This Notice is soliciting comments from members of the public and affected agencies concerning the proposed collection of information to: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond; including through the use of appropriate automated collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

The Self-Help Homeownership Opportunity Program (SHOP) is authorized by the Housing Opportunity Program Extension Act of 1996, Section 11. The purpose of SHOP is to provide grant funds to facilitate and encourage innovative homeownership opportunities on a national, geographically diverse basis through the provision of self-help homeownership housing programs. SHOP funds are appropriated by Congress, generally annually. HUD publishes a SHOP Notice of Funding Availability (NOFA) that announces the amount of SHOP grant funds and the application criteria, including the rating and ranking system HUD will use to select grantees.

SHOP grant funds may be used for land acquisition, the installation or