

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.*

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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If any Laboratory/IITF has withdrawn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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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.

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