increase efficiency in our funding process.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 2295.

### ESTIMATED ANNUALIZED BURDEN HOURS

<table>
<thead>
<tr>
<th>Form</th>
<th>Type of respondent</th>
<th>Number of respondents</th>
<th>Frequency of response</th>
<th>Average time per response (in hours)</th>
<th>Annual hour burden</th>
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</thead>
<tbody>
<tr>
<td>NIMH Recruitment Milestone Reporting</td>
<td>Principal assistant, Principal assistant</td>
<td>900</td>
<td>3</td>
<td>45/60</td>
<td>2025</td>
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<tr>
<td>NIMH Recruitment Milestone Monthly Report</td>
<td>Investigators/Research Assistant</td>
<td>40</td>
<td>9</td>
<td>45/60</td>
<td>270</td>
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</tbody>
</table>

Dated: May 21, 2015.
Keisha L. Shropshire,
Project Clearance Liaison, NIMH, NIH.
[FR Doc. 2015–13112 Filed 5–29–15; 8:45 am]
BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Current List of HHS-Certified 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 29900); 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 HHS-certified laboratories and IITFs is published in the Federal Register during the first week of each month. If any laboratory or IITF certification is suspended or revoked, the laboratory or IITF will be omitted from subsequent lists until such time as it is restored to full certification under the Mandatory Guidelines.

If any laboratory or IITF has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end and will be omitted from the monthly listing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://beta.samhsa.gov/workplace.

FOR FURTHER INFORMATION CONTACT:
Giselle Hersh, Division of Workplace Programs, SAMHSA/CSAP, Room 7–1051, 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 strict standards that laboratories and IITFs must meet in order to conduct drug and specimen validity tests on urine specimens for federal agencies.

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

Laboratories and IITFs in the applicant stage of certification are not to be considered as meeting the minimum requirements described in the HHS Mandatory Guidelines. A HHS-certified laboratory or 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 HHS-certified laboratories and IITFs meet the minimum standards to conduct drug and specimen validity tests on urine specimens:

Alere Toxicology Services, 1111 Newton St., Gretna, LA 70053, 504–361–8989/800–433–3823, (Formerly: Kroll Laboratory Specialists, Inc., Laboratory Specialists, 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.
DrugScan, Inc., 200 Precision Road, Suite 200, Horsham, PA 19044, 800–235–4890.
Fortes Laboratories, Inc., 25740 SW Canyon Creek Road, Suite 600, Wilsonville, OR 97070, 503–486–1023.
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,
(Formerly: LabCorp Occupational
Testing Services, Inc., CompuChem
Laboratories, Inc.; CompuChem
Laboratories, Inc., A Subsidiary of
Roche Biomedical Laboratory;
Roche LabOne, 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: Lab-
Corp 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.).

MedTox Laboratories, Inc., 402 W.
County Road D, St. Paul, MN 55112,

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: Cantinela
Hospital Airport
Toxicology Laboratory).

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

Phamatech, Inc., 15175 Innovation
Drive, San Diego, CA 92128, 888–
635–5840, 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 Rd., Norristown, PA 19403,
(Formerly: SmithKline Beecham
Clinical Laboratories; SmithKline
Beecham Clinical Laboratories; SmithKline
Beecham Clinical Laboratories; SmithKline
Beecham Clinical Laboratories).

Quest Diagnostics Incorporated, 8401
Fallbrook Ave., West Hills, CA
91304, 818–737–6370, (Formerly:
SmithKline Beecham Clinical
Laboratories).

Redwood Toxicology Laboratory,
3700650 Westwind Blvd., Santa
Rosa, CA 95403, 800–235–2159.

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

STERLING Reference Laboratories, 2617
East I Street, Tacoma, Washington
98421, 800–442–0438.

U.S. Army Forensic Toxicology Drug
Testing Laboratory, 2490 Wilson
St., Fort George G. Meade, MD

* 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
certification, the laboratory will be
given certification
pursuant to CBP regulations, that
Intertek USA, Inc., as approved to
gauge and accredited to test petroleum
and petroleum products for customs
purposes, in accordance with the
provisions of 19 CFR 151.12 and 19 CFR
151.13.

Intertek USA, Inc., is approved for the
following gauging procedures for
petroleum and certain petroleum
products set forth by the American
Petroleum Institute (API):

<table>
<thead>
<tr>
<th>API Chapters</th>
<th>Title</th>
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<tbody>
<tr>
<td>3</td>
<td>Tank gauging.</td>
</tr>
<tr>
<td>7</td>
<td>Temperature Determination.</td>
</tr>
<tr>
<td>8</td>
<td>Sampling.</td>
</tr>
<tr>
<td>12</td>
<td>Calculations.</td>
</tr>
<tr>
<td>17</td>
<td>Maritime Measurements.</td>
</tr>
</tbody>
</table>

Intertek USA, Inc., is accredited for the
following laboratory analysis
procedures and methods for petroleum
and certain petroleum products set forth
by the U.S. Customs and Border
Protection Laboratory Methods (CBP)
and American Society for Testing and
Materials (ASTM):