Send comments to Summer King, SAMHSA Reports Clearance Officer, Room 7–1044, 1 Choke Cherry Road, Rockville, MD 20857 AND e-mail her a copy at summer.king@samhsa.hhs.gov. Written comments should be received within 60 days of this notice.

Elaine Parry,  
Director, Office of Program Services.  
[FR Doc. E9–11377 Filed 5–14–09; 8:45 am]

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If any laboratory has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end, and will be omitted from the monthly listing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://www.workplace.samhsa.gov and 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 developed in accordance with Executive Order 12564 and section 503 of Public Law 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:

<table>
<thead>
<tr>
<th>Instrument/form</th>
<th>Number of respondents</th>
<th>Responses per respondent</th>
<th>Total responses</th>
<th>Hours per response</th>
<th>Total annualized hour burden per respondent</th>
</tr>
</thead>
<tbody>
<tr>
<td>Family Program Discharge Form</td>
<td>4</td>
<td>50</td>
<td>200</td>
<td>.16</td>
<td>8</td>
</tr>
<tr>
<td>Volunteer/Staff Survey</td>
<td>4</td>
<td>1</td>
<td>4</td>
<td>.25</td>
<td>.25</td>
</tr>
<tr>
<td>Support Services Supervisor:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Adolescent Telephone Support QA Checklist</td>
<td>1</td>
<td>12</td>
<td>12</td>
<td>1</td>
<td>12</td>
</tr>
<tr>
<td>Social Networking QA Checklist</td>
<td>1</td>
<td>12</td>
<td>12</td>
<td>.5</td>
<td>6</td>
</tr>
<tr>
<td>Family Program QA Checklist</td>
<td>1</td>
<td>12</td>
<td>12</td>
<td>1</td>
<td>12</td>
</tr>
<tr>
<td>Volunteer/Staff Survey</td>
<td>18</td>
<td>1</td>
<td>18</td>
<td>.25</td>
<td>.25</td>
</tr>
<tr>
<td>Column Total</td>
<td>418</td>
<td>2580</td>
<td>713.67</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Total Annualized Hour Burden Per Respondent = Responses Per Respondent × Hours Per Response.

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


South Bend Medical Foundation, Inc., 530 N. Lafayette Blvd., South Bend, IN 46601, 574–234–4176 x276.


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.


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

The following laboratory voluntarily withdrew from the NLCP on March 31, 2009:

Sparrow Health System, Toxicology Testing Center, St. Lawrence Campus, 1210 W. Saginaw, Lansing, MI 48915, 517–364–7400, (Formerly: St. Lawrence Hospital & Healthcare System).

The Standards Council of Canada (SCC) voted to end its Laboratory Accreditation Program for Substance Abuse (LAPSA) effective May 12, 1998. Laboratories accredited through that program were accredited to conduct forensic urine drug testing as required by U.S. Department of Transportation


Elaine Parry,
Director, Office of Program Services,
SAMHSA.

[FR Doc. E9–11374 Filed 5–14–09; 8:45 am]
BILLING CODE 4160–20–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Advisory Committee on Rural Health and Human Services

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), notice is hereby given that the following committee will convene its sixty-second meeting.

Name: National Advisory Committee on Rural Health and Human Services

Dates and Times: June 9, 2009, 9 a.m.—4:45 p.m.

June 10, 2009, 8:45 a.m.—3 p.m.

June 11, 2009, 8:45 a.m.—11 a.m.

Place: Hampton Inn, 1720 Rapp Street, Rapid City, South Dakota 57701.

Phone: 605–348–1911.

Status: The meeting will be open to the public.

Purpose: The National Advisory Committee on Rural Health and Human Services provides advice and recommendations to the Secretary with respect to the delivery, research, development and administration of health and human services in rural areas.

Agenda: Tuesday morning, at 9 a.m., the meeting will be called to order by the Chairperson of the Committee, the Honorable David Beasley. The first presentation will be an overview of rural South Dakota by Dr. Sidney Goss, Professor of Demography, South Dakota School of Mines and Technology. The Committee will be formally welcomed by the South Dakota Office of Rural Health, Sandra Durick, Director. The Committee will hear presentations on the three chosen

(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 13, 2004 (69 FR 19644). 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.