vaccines with the VICP even before such vaccines are added as a separate and distinct category to the Table through rulemaking.

DATES: This Notice is effective on December 1, 2004. As described below, Hepatitis A vaccines will be covered under the VICP on December 1, 2004.

FOR FURTHER INFORMATION CONTACT: Geoffrey Evans, M.D., Medical Director, Division of Vaccine Injury Compensation, Healthcare Systems Bureau, Health Resources and Services Administration, Parklawn Building, Room 11C–26, 5600 Fishers Lane, Rockville, Maryland 20857; telephone number (301) 443–4198.

SUPPLEMENTARY INFORMATION: The statute authorizing the VICP provides for the inclusion of additional vaccines in the VICP when they are recommended by the Centers for Disease Control and Prevention (CDC) to the Secretary for routine administration to children. See section 2114(e)(2) of the Public Health Service (PHS) Act, 42 U.S.C. 300aa–14(e)(2). Consistent with section 13632(a)(3) of Public Law 103–66, the regulations governing the VICP provide that such vaccines will be included in the Table as covered vaccines as of the effective date of an excise tax to provide funds for the payment of compensation with respect to such vaccines (42 CFR 100.3(c)(5)).

The two prerequisites for adding Hepatitis A vaccines to the VICP as covered vaccines as well as to the Table have been satisfied. First, the CDC published its recommendation that Hepatitis A vaccines be routinely administered to certain children in the October 1, 1999, issue of the Morbidity and Mortality Weekly Report (MMWR). Specifically, the CDC recommended that all children in States, counties, and communities with rates of Hepatitis A that are twice the 1987–1997 national average or greater (*i.e.*, greater than or equal to 20 cases per 100,000 population) receive the Hepatitis A vaccine.

Second, on October 22, 2004, the excise tax for Hepatitis A vaccines was enacted by Public Law 108–357, the ''American Jobs Creation Act of 2004.'' Section 889 of this Act adds all vaccines against Hepatitis A to section 4132(a)(1) of the Internal Revenue Code of 1986, which defines all taxable vaccines. Unlike the CDC's recommendation, the American Jobs Creation Act of 2004 does not distinguish between Hepatitis A vaccines administered in areas in which rates of Hepatitis A are at least twice the national average and Hepatitis A vaccines administered in other areas of the country. For this reason, all Hepatitis A vaccines manufactured or produced in the United States, or entered into the United States for consumption, use, or warehousing, will be subject to this excise tax (26 U.S.C. 4132(a)(1)).

Under the regulations governing the VICP, Item XIV of the Table specifies that ''[a]ny new vaccine recommended by the Centers for Disease Control and Prevention for routine administration to children, after publication by the Secretary of a notice of coverage'' is a covered vaccine under the Table. (42 CFR 100.3(a), Item XIV.) As explained above, the CDC's recommendation was accepted. This notice serves to satisfy the regulation's publication requirement. Through this notice, Hepatitis A vaccines are included as covered vaccines under Category XIV of the Table. As explained above, because the American Jobs Creation Act of 2004 enacted an excise tax for Hepatitis A vaccines administered throughout the United States, all Hepatitis A vaccines will be covered under the VICP and under the Table.

Under section 2114(e) of the PHS Act, as amended by section 13632(a) of the Omnibus Budget Reconciliation Act of 1993, coverage for a vaccine recommended by the CDC for routine administration to children shall take effect upon the effective date of the tax enacted to provide funds for compensation with respect to the vaccine included as a covered vaccine in the Table. The American Jobs Creation Act of 2004 provides that the addition of Hepatitis A vaccines to the list of taxable vaccines applies to sales and uses on or after the first day of the first month which begins more than 4 weeks after the date of the enactment of the Act. It further provides that if the vaccines were sold before or on the effective date of the excise tax, but delivered after this date, the delivery date of such vaccines shall be considered the sale date. Because the American Jobs Creation Act of 2004 was enacted on October 22, 2004, the effective date of the excise tax adding Hepatitis A vaccines as taxable vaccines is December 1, 2004. Thus, Hepatitis A vaccines are included as covered vaccines under Category XIV of the Table as of December 1, 2004. Petitioners may file petitions related to Hepatitis A vaccines as of December 1, 2004.

Petitions filed concerning vaccine-related injuries or deaths associated with Hepatitis A vaccines must, of course, be filed within the applicable statute of limitations. The statutes of limitations applicable to petitions filed with the VICP are set out in section 2116(a) of the PHS Act (42 U.S.C. 300aa–16(a)). In addition, section 2116(b) of the PHS Act lays out specific exceptions to these statutes of limitations that apply when the effect of a revision to the Table makes a previously ineligible person eligible to receive compensation or when an eligible person's likelihood of obtaining compensation significantly increases. Under this provision, a person who may be eligible to file a petition based on the addition of a new vaccine under Category XIV of the Table may file a petition for compensation not later than 2 years after the effective date of the revision if the injury or death occurred not more than 8 years before the effective date of the revision of the Table (42 U.S.C. 300aa–16(b)). Thus, persons whose petitions may not satisfy the limitations periods described in section 2116(a) of the PHS Act may still file petitions concerning vaccine-related injuries or deaths associated with Hepatitis A vaccines until December 1, 2006, as long as the vaccine-related injury or death occurred on or after December 1, 1996 (8 years prior to the effective date of the addition that included Hepatitis A as a covered vaccine).

The Secretary plans to amend the Table through the rulemaking process by including Hepatitis A vaccines as a separate category of vaccines in the Table. December 1, 2004, will remain the applicable effective date when the Secretary makes a corresponding amendment to add Hepatitis A vaccines as a separate category on the Table through rulemaking.

Dated: November 22, 2004.

Elizabeth M. Duke,

Administrator, HRSA.

[FR Doc. 04–26273 Filed 11–30–04; 8:45 am]

BILLING CODE 4165–15–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 (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://workplace.samhsa.gov* and *http://www.drugfreeworkplace.gov*.

FOR FURTHER INFORMATION CONTACT: Mrs. Giselle Hersh or Dr. Walter Vogl, Division of Workplace Programs, SAMHSA/CSAP, Room 2–1035, 1 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 Pub. L. 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:

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.

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.

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

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

DrugProof, Division of Dynacare/ Laboratory of Pathology, LLC, 1229 Madison St., Suite 500, Nordstrom Medical Tower, Seattle, WA 98104, 206–386–2661/800–898–0180, (Formerly: Laboratory of Pathology of Seattle, Inc., DrugProof, Division of Laboratory of Pathology of Seattle, Inc.).

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

Dynacare Kasper Medical Laboratories*, 10150–102 St., Suite 200, Edmonton, Alberta, Canada T5J 5E2, 780–451–3702/800–661–9876.

ElSohly Laboratories, Inc., 5 Industrial Park Dr., Oxford, MS 38655, 662–236–2609.

Express Analytical Labs, 3405 7th Ave., Suite 106, Marion, IA 52302, 319–377–0500.

General Medical Laboratories, 36 South Brooks St., Madison, WI 53715, 608–267–6225.

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

LabOne, Inc., 10101 Renner Blvd., Lenexa, KS 66219, 913–888–3927/800–873–8845, (Formerly: Center for Laboratory Services, a Division of LabOne, Inc.).

LabOne, Inc., d/b/a Northwest Toxicology, 1141 E. 3900 S., Salt Lake City, UT 84124, 801–293–2300/800–322–3361, (Formerly: NWT Drug Testing, NorthWest Toxicology, Inc.; Northwest Drug Testing, a division of NWT Inc.).

Laboratory Corporation of America Holdings, 7207 N. Gessner Rd., 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 Dr., Research Triangle Park, NC 27709, 919–572–6900/800–833–3984, (Formerly: LabCorp Occupational Testing Services, 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, 10788 Roselle St., San Diego, CA 92121, 800–882–7272 (Formerly: Poisonlab, Inc.).

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).

Marshfield Laboratories, Forensic Toxicology Laboratory, 1000 North Oak Ave., Marshfield, WI 54449, 715–389–3734/800–331–3734.

MAXXAM Analytics Inc.,* 6740 Campobello Road, Mississauga, ON, Canada L5N 2L8, 905–817–5700, (Formerly: NOVAMANN (Ontario) Inc.).

MedTox Laboratories, Inc., 402 W. County Rd. 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 Dr., 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).

Oregon Medical Laboratories, P.O. Box 972, 722 East 11th Ave., Eugene, OR 97440–0972, 541–687–2134.

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, 509–755–8991/800–541–7897 x7.
Physicians Reference Laboratory, 7800 West 110th St., Overland Park, KS 66210, 913–339–0372/800–821–3627.
Quest Diagnostics Incorporated, 3175 Presidential Dr., Atlanta, GA 30340, 770–452–1590/800–729–6432, (Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories).
Quest Diagnostics Incorporated, 4770 Regent Blvd., Irving, TX 75063, 800–824–6152, (Moved from the Dallas location on 03/31/01; Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories).
Quest Diagnostics Incorporated, 4230 South Burnham Ave., Suite 250, Las Vegas, NV 89119–5412, 702–733–7866/800–433–2750, (Formerly: Associated Pathologists Laboratories, Inc.).
Quest Diagnostics Incorporated, 400 Egypt Rd., Norristown, PA 19403, 610–631–4600/877–642–2216, (Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories).
Quest Diagnostics Incorporated, 506 E. State Pkwy., Schaumburg, IL 60173, 800–669–6995/847–885–2010, (Formerly: SmithKline Beecham Clinical Laboratories; International Toxicology Laboratories).
Quest Diagnostics Incorporated, 7600 Tyrone Ave., Van Nuys, CA 91405, 818–989–2520/800–877–2520, (Formerly: SmithKline Beecham Clinical Laboratories).
Scientific Testing Laboratories, Inc., 450 Southlake Blvd., Richmond, VA 23236, 804–378–9130.
Sciteck Clinical Laboratories, Inc., 317 Rutledge Rd., Fletcher, NC 28732, 828–650–0409.
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 x276.
Southwest Laboratories, 4645 E. Cotton Center Boulevard, Suite 177, Phoenix, AZ 85040, 602–438–8507/800–279–0027.
Sparrow Health System, Toxicology Testing Center, St. Lawrence Campus, 1210 W. Saginaw, Lansing, MI 48915, 517–364–7400, (Formerly: St. Lawrence Hospital & Healthcare System).
St. Anthony Hospital Toxicology Laboratory, 1000 N. Lee St., Oklahoma City, OK 73101, 405–272–7052.
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.

Anna Marsh,

Executive Officer, SAMHSA.

[FR Doc. 04–26426 Filed 11–30–04; 8:45 am]

BILLING CODE 4160–20–P

DEPARTMENT OF THE INTERIOR

Bureau of Land Management

[NV–912–05–1990–PO–241A–006F]

Sierra Front-Northwestern Great Basin Resource Advisory Council; Notice of Meeting Location and Time

AGENCY: Bureau of Land Management, Interior.

ACTION: Notice of meeting location and time for the Sierra Front-Northwestern Great Basin Resource Advisory Council (Nevada).

SUMMARY: In accordance with the Federal Land Policy and Management Act and the Federal Advisory Committee Act of 1972 (FACA), a meeting of the U.S. Department of the Interior, Bureau of Land Management (BLM) Sierra Front-Northwestern Great Basin Resource Advisory Council (RAC), Nevada, will be held as indicated below. Topics for discussion at the meeting will include, but are not limited to: Manager's reports of current field office activities; review of the BLM's Resource Management Planning Process; review of the Pine Nut Mountain RMP Amendment DEIS; a review of Native American tribe consultation procedures; review of proposed 2005 Wild Horse Herd Management Area gathers in the Northwest Great Basin; a panel discussion on water resources transportation issues in Nevada; and additional topics the council may raise during the meeting.

Date & Time: The RAC will meet on Thursday, January 27, 2005, from 9 a.m. to 5 p.m., at the BLM-Carson City Field Office, 5665 Morgan Mill Road, Carson City, Nevada; and on Friday, January 28, 2005, from 8 a.m. to 12 p.m., at the BLM-Nevada State Office, Great Basin A&B Conference Room, 1340 Financial Blvd., Reno, Nevada. All meetings are open to the public. A general public comment period, where the public may submit oral or written comments to the RAC, will be held at 4 p.m. on January 27, 2005.

A final detailed agenda, with any additions/corrections to agenda topics, will be available on the Internet no later than January 13, 2005, at *http://www.nv.blm.gov/rac;* hard copies can also be mailed or sent via FAX. Individuals who need special assistance such as sign language interpretation or other reasonable accommodations, or who wish a hard copy of the agenda, should contact Mark Struble, Carson City Field Office, 5665 Morgan Mill Road, Carson City, NV 89701, telephone (775) 885–6107, no later than January 13, 2005.

FOR FURTHER INFORMATION CONTACT: Mark Struble, Public Affairs Officer, BLM Carson City Field Office, 5665 Morgan Mill Road, Carson City, NV 89701. Telephone: (775) 885–6107. E-mail: *mstruble@nv.blm.gov.*

Dated: November 23, 2004.

Don Hicks,

Field Office Manager, BLM-Carson City Field Office.

[FR Doc. 04–26491 Filed 11–30–04; 8:45 am]

BILLING CODE 4310–HC–P

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* 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 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.