

COMMUNITY LEVEL SURVEY BURDEN ESTIMATE—Continued

Survey section	Content description	Number of respondents	Number of responses	Hourly burden/response	Total hourly burden
172–178 .....	Contextual Factors and Closing Questions.	390	2	1.0	780
Part II, 1–52 .....	Intervention Specific Information and Adaptations.	390	3	2.0	2,340
53–60 .....	Intervention Outcomes .....	390	6	1.0	2,340
Sub-forms .....	Intervention Component Information	390	6	1.0	2,340
Review of past responses .....	.....	390	2	1.0	780
Total Community Level Year 3 Burden.	.....	.....	24	7.9	10,062
Average Annual Community Burden.	.....	.....	21	8.1	8,827

Send comments to Summer King, SAMHSA Reports Clearance Officer, Room 71–1044, One Choke Cherry Road, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: December 30, 2005.

**Anna Marsh,**

*Director, Office of Program Services.*

[FR Doc. E6–95 Filed 1–9–06; 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 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If any laboratory has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end, and will be omitted from the monthly listing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 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:

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 Road, Lenexa, KS 66215–2802. 800–445–6917.

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

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.

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 Drive, Oxford, MS 38655. 662–236–2609.

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

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

- General Medical Laboratories, 36 South Brooks St., Madison, WI 53715. 608-267-6225.
- Kroll Scientific Testing Laboratories, Inc., 450 Southlake Blvd., Richmond, VA 23236. 804-378-9130. (Formerly: Scientific Testing Laboratories, Inc.)
- 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 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, 550 17th Ave., Suite 300, Seattle, WA 98122. 206-923-7020/800-898-0180. (Formerly: DrugProof, Division of Dynacare/Laboratory of Pathology, LLC; Laboratory of Pathology of Seattle, Inc.; DrugProof, Division of Laboratory of Pathology of Seattle, 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, ONT, Canada L5N 2L8. 905-817-5700. (Formerly: 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.)
- 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. 509-755-8991/800-541-7897x7.
- 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, 10101 Renner Blvd., Lenexa, KS 66219. 913-888-3927/800-873-8845. (Formerly: LabOne, Inc.; Center for Laboratory Services, a Division of LabOne, Inc.)
- 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, 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).
- Quest Diagnostics Incorporated, 2282 South Presidents Drive, Suite C, West Valley City, UT 84120. 801-606-6301/800-322-3361. (Formerly:
- Northwest Toxicology, a LabOne Company; LabOne, Inc., dba Northwest Toxicology; NWT Drug Testing, Northwest Toxicology, Inc.; Northwest Drug Testing, a division of NWT Inc.)
- 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.
- US 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 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.

**Anna Marsh,**

*Director, Office Program Services, SAMHSA.*

[FR Doc. E6-96 Filed 1-9-06; 8:45 am]

**BILLING CODE 4162-20-P**

## DEPARTMENT OF THE INTERIOR

### Fish and Wildlife Service

#### Endangered Species Recovery Permit Application

**AGENCY:** Fish and Wildlife Service, Interior.

**ACTION:** Notice of receipt of a permit application.

**SUMMARY:** The following applicant has applied for an enhancement of propagation or survival permit to conduct certain activities with endangered species pursuant to section 10(a)(1)(A) of the Endangered Species Act. The U.S. Fish and Wildlife Service ("we") solicits review and comment from local, State and Federal agencies, and the public on the following permit request.

**DATES:** Comments on this permit application must be received on or before February 9, 2006.

**ADDRESSES:** Written data or comments should be submitted to the U.S. Fish and Wildlife Service, Chief, Endangered Species, Ecological Services, 911 NE. 11th Avenue, Portland, Oregon 97232-4181 (fax: 503-231-6243). Please refer to the permit number when submitting comments. All comments received, including names and addresses, will become part of the official administrative record and may be made available to the public.

**FOR FURTHER INFORMATION CONTACT:** Documents and other information submitted with this application are available for review, subject to the requirements of the Privacy Act and Freedom of Information Act, by any party who submits a written request for a copy of such documents within 30 days of the date of publication of this notice to the address above (telephone: 503-231-2063). Please refer to the permit number when requesting copies of documents.

**SUPPLEMENTARY INFORMATION:**

**Permit No. TE-114934**

*Applicant:* Idaho Department of Fish and Game, Boise, Idaho.

The applicant requests a permit to take (harass by survey, capture, handle, collar, take blood samples, relocate, sacrifice, and release) the gray wolf (*Canis lupus*) in conjunction with wolf management activities in the State of Idaho north of Interstate 90, for the purpose of enhancing its survival.

The applicant proposes to: (a) Conduct monitoring of wolf populations; and (b) coordinate non-lethal and lethal control actions to reduce wolf conflicts with livestock and dogs. These actions are currently coordinated by the U.S. Fish and Wildlife Service (Service). If the permit is issued, the applicant would take on responsibility for managing wolves in northern Idaho. Wolf management activities would be in accordance with the requirements of the State of Idaho Wolf Conservation and Management Plan (March 2002) and the Service's Interim Wolf Control Plan for Northwestern Montana and the Panhandle of Northern Idaho (Control Plan) (September 1999).

If issued, the permit would not affect ongoing wolf management in the remainder of the State of Idaho conducted in accordance with the non-essential experimental population regulations found at 50 CFR 17.84(n).

We have determined that a practical, responsive management program is essential to enhancing survival of the wolf in the wild (Service 1987; Service 1994; Service 1999). The program must respond to wolf-livestock conflicts, while promoting wolf recovery objectives. The Control Plan provides guidelines for: (a) Determining problem wolf status (including investigative procedures and criteria), (b) conducting wolf control actions, and (c) disposition of problem wolves.

We have made a preliminary determination that issuance of this permit would be categorically excluded from further consideration under the National Environmental Policy Act of 1969 (NEPA). If issued, the permit would authorize Idaho Department of Fish and Game to manage wolves in the same manner that the Service has in the 6 years since the Control Plan was adopted for northern Idaho. No additional environmental impacts would be expected beyond those analyzed in the Service's 1988 Environmental Assessment, the Service's 1999 Environmental Action Memorandum, and the Control Plan. The effect of the permit would be to allow continuation of previously analyzed and authorized activities; therefore, its issuance would be an administrative action.

Our preliminary NEPA categorical exclusion determination, the two wolf plans noted above, and the Idaho Department of Fish and Game permit application, can be found at: <http://www.fws.gov/pacific/ecoservices/Endangered/Recovery/default.htm>.

Additional information about wolf recovery and conservation in the northwestern United States, including control of problem wolves, can be found in various reports at: <http://westerngraywolf.fws.gov/>.

All comments received from individuals become part of the official public record. Requests for such comments will be handled in accordance with the Freedom of Information Act and the Council on Environmental Quality's National Environmental Protection Act regulations [40 CFR 1506.6(f)]. Our practice is to make comments, including names and addresses of respondents, available for public review during regular business hours. Individual respondents may request that we withhold their home address from the record, but this must be stated prominently at the beginning of their comments. We will honor these requests to the extent allowable by law.

We solicit public review and comment on this recovery permit application.

**Authority:** This document is published under the authority of the Endangered Species Act of 1973, as amended (16 U.S.C. 1531 *et seq.*).

Dated: December 13, 2005.

**David J. Wesley,**

*Acting Regional Director, Region 1, U.S. Fish and Wildlife Service.*

[FR Doc. E6-93 Filed 1-9-06; 8:45 am]

**BILLING CODE 4310-55-P**

## DEPARTMENT OF THE INTERIOR

### Bureau of Land Management

[NV-030-1610-DU]

#### Notice of Intent To Prepare an Amendment to the Carson City Field Office Consolidated Resource Management Plan

**AGENCY:** Bureau of Land Management, Interior.

**ACTION:** Notice of intent.

**SUMMARY:** This document provides notice that the Bureau of Land Management (BLM) intends to prepare a Resource Management Plan (RMP) amendment to address Lands and Land Tenure Issues, and Recreation and Travel Management. The appropriate state, tribal and local governments will