along with copies of requested SF 278 reports. Those who choose to respond can complete and return the survey to OGE via the self-contained postage-paid postcards (the reverse side of the survey form, when folded, becomes a pre-addressed postcard). The purpose of this anonymous survey is to determine through customer responses how well OGE is responding to such requests and how OGE can improve its customer service in this important area. The current paperwork approval for the survey form is scheduled to expire at the end of January 2002.

On June 18, 2001, OGE issued its first round **Federal Register** notice to announce its forthcoming request to OMB for paperwork renewal of the customer service survey form. See 66 FR 32823–32824 with comments due by September 4, 2001. (OGE did not receive any comments or requests for copies of the customer service survey form). In that notice, and this one, OGE has proposed minor changes to survey question 4 to achieve greater clarity. That question currently asks whether OGE's requirement to fax or mail requests that involve more than six filers creates a problem for the requester. Based on an analysis of customer responses to question 4, OGE believes that the following statement should be added: ''SKIP this question if your request involved six or fewer filers.'' Additionally, one of the three requested responses to question 4, ''Not Applicable,'' is being changed to ''My request did not have to be faxed or mailed.''

Pursuant to the Paperwork Reduction Act, OGE has not included in its public burden estimate for the survey form the limited number of access requests filed by other Federal agencies or Federal employees. Nor has OGE included in that estimate the limited number of requests for copies of other records covered under the special Ethics Act public access provision (such as certificates of divestiture) since the survey is only sent to persons who request copies of SF 278 reports. As so defined, the total number of access survey forms for copies of SF 278s estimated to be filed annually at OGE over the next three years by members of the public (primarily by news media representatives, public interest group members and private citizens) is 50. This estimate is based on a calculation of the number of survey forms received at OGE between April 1999 and June 2001 (70 surveys). This number also takes into account the increase in the number of public requests experienced as a result of the transition and the new Presidential administration. The estimated average amount of time to read the instructions on the proposed revised customer service survey form, and to complete the form remains at three minutes. Thus, the overall estimated annual public burden for the OGE Public Financial Disclosure Access Customer Service Survey as proposed for revision will be three hours (rounded up from two and a half hours(= 50 forms × 3 minutes per form)).

In this second round notice, public comment is again invited on all aspects of OGE's customer service survey form as proposed for renewal with minor revision, including specifically views on: the accuracy of OGE's public burden estimate; the potential for enhancement of quality, utility, and clarity of the information to be collected; and the minimization of burden (including the possibility of use of information technology). The Office of Government Ethics, in consultation with OMB, will consider all comments received, which will become a matter of public record.

Approved: December 31, 2001.

Amy L. Comstock,

Director, Office of Government Ethics.

[FR Doc. 02–327 Filed 1–4–02; 8:45 am]

BILLING CODE 6345–01–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 notifies Federal agencies of the laboratories currently certified to meet standards of Subpart C of Mandatory Guidelines for Federal Workplace Drug Testing Programs (59 FR 29916, 29925). 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 Guidelines.

If any laboratory has withdrawn from the National Laboratory Certification Program 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 the following Web sites: *http://workplace.samhsa.gov; http://www.drugfreeworkplace.gov;* and *http://www.health.org/workplace.*

FOR FURTHER INFORMATION CONTACT: Mrs. Giselle Hersh or Dr. Walter Vogl, Division of Workplace Programs, 5600 Fishers Lane, Rockwall 2 Building, Room 815, Rockville, Maryland 20857; Tel.: (301) 443–6014, Fax: (301) 443–3031.

SUPPLEMENTARY INFORMATION:

Mandatory Guidelines for Federal Workplace Drug Testing were developed in accordance with Executive Order 12564 and section 503 of Pub. L. 100–71. Subpart C of the Guidelines, ''Certification of Laboratories Engaged in Urine Drug Testing for Federal Agencies,'' sets strict standards which laboratories must meet in order to conduct urine drug testing 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 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 expressed in the HHS Guidelines. A laboratory must have its letter of certification from SAMHSA, HHS (formerly: HHS/NIDA) which attests that it has met minimum standards.

In accordance with Subpart C of the Guidelines, the following laboratories meet the minimum standards set forth in the Guidelines:

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, 716–429–2264

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

Alliance Laboratory Services, 3200 Burnet Ave., Cincinnati, OH 45229, 513–585–9000 (Formerly: Jewish Hospital of Cincinnati, Inc.)

American Medical Laboratories, Inc., 14225 Newbrook Dr., Chantilly, VA 20151, 703–802–6900

Associated Pathologists Laboratories, Inc., 4230 South Burnham Ave., Suite 250, Las Vegas, NV 89119–5412, 702–733–7866/800–433–2750

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 Laboratory Partners, LLC, 129 East Cedar St., Newington, CT 06111, 860–696–8115 (Formerly: Hartford Hospital Toxicology Laboratory)
Clinical Reference Lab, 8433 Quivira Rd., Lenexa, KS 66215–2802, 800–445–6917
Cox Health Systems, Department of Toxicology, 1423 North Jefferson Ave., Springfield, MO 65802, 800–876–3652/417–269–3093 (Formerly: Cox Medical Centers)
Diagnostic Services Inc., dba DSI, 12700 Westlinks Drive, Fort Myers, FL 33913, 941–561–8200/800–735–5416
Doctors Laboratory, Inc., P.O. Box 2658, 2906 Julia Dr., Valdosta, GA 31602, 912–244–4468
DrugProof, Divison of Dynacare, 543 South Hull St., Montgomery, AL 36103, 888–777–9497/334–241–0522 (Formerly: Alabama Reference Laboratories, Inc.)
DrugProof, Division of Dynacare/Laboratory of Pathology, LLC, 1229 Madison St., Suite 500, Nordstrom Medical Tower, Seattle, WA 98104, 206–386–2672/800–898–0180 (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,* 14940–123 Ave., Edmonton, Alberta, Canada T5V 1B4, 780–451–3702/800–661–9876
ElSohly Laboratories, Inc., 5 Industrial Park Dr., Oxford, MS 38655, 662–236–2609
Express Analytical Labs, 3405 7th Avenue, Suite 106, Marion, IA 52302, 319–377–0500
Gamma-Dynacare Medical Laboratories,* A Division of the Gamma-Dynacare Laboratory Partnership, 245 Pall Mall St., London, ONT, Canada N6A 1P4, 519–679–1630
General Medical Laboratories, 36 South Brooks St., Madison, WI 53715, 608–267–6267
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, 913–888–3927/800–728–4064 (Formerly: Center for Laboratory Services, a Division of LabOne, 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 Street, San Diego, CA 92121, 800–882–7272 (Formerly: Poisonlab, Inc.)
Laboratory Corporation of America Holdings, 1120 Stateline Road West, 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.*, 5540 McAdam Rd., Mississauga, ON, Canada L4Z 1P1, 905–890–2555, (Formerly: NOVAMANN (Ontario) Inc.)
Medical College Hospitals Toxicology Laboratory, Department of Pathology, 3000 Arlington Ave., Toledo, OH 43699, 419–383–5213
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 Drive, Minneapolis, Minnesota 55417, 612–725–2088
National Toxicology Laboratories, Inc., 1100 California Ave., Bakersfield, CA 93304, 661–322–4250/800–350–3515
Northwest Drug Testing, a division of NWT Inc., 1141 E. 3900 South, Salt Lake City, UT 84124, 801–293–2300/800–322–3361, (Formerly: NWT Drug Testing, NorthWest Toxicology, Inc.)
One Source Toxicology Laboratory, Inc., 1705 Center Street, Deer Park, TX 77536, 713–920–2559, (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, 6160 Variel Ave., Woodland Hills, CA 91367, 818–598–3110/800–328–6942, (Formerly: Centinela Hospital Airport Toxicology Laboratory
Pathology Associates Medical Laboratories, 110 West Cliff Drive, Spokane, WA 99204, 509–755–8991/800–541–7891x8991
PharmChem Laboratories, Inc., 4600 N. Beach, Haltom City, TX 76137, 817–605–5300, (Formerly: PharmChem Laboratories, Inc., Texas Division; Harris Medical Laboratory)
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, (Formerly: SmithKline Beecham Clinical Laboratories, SmithKline Bio-Science Laboratories)
Quest Diagnostics Incorporated, 4770 Regent Blvd., Irving, TX 75063, 800–842–6152, (Moved from the Dallas location on 03/31/01; Formerly: SmithKline Beecham Clinical Laboratories, SmithKline Bio-Science Laboratories)
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, 7470 Mission Valley Rd., San Diego, CA 92108–4406, 619–686–3200/800–446–4728 (Formerly: Nichols Institute, Nichols Institute Substance Abuse Testing (NISAT), CORNING Nichols Institute, CORNING Clinical 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., 463 Southlake Blvd., Richmond, VA 23236, 804–378–9130
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, 219–234–4176
Southwest Laboratories, 2727 W. Baseline Rd., Tempe, AZ 85283, 602–438–8507/800–279–0027
Sparrow Health System, Toxicology Testing Center, St. Lawrence Campus, 1210 W. Saginaw, Lansing, MI 48915, 517–377–0520 (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, 2703 Clark Lane, Suite B, Lower Level, Columbia, MO 65202, 573–882–1273
Toxicology Testing Service, Inc., 5426 N.W. 79th Ave., Miami, FL 33166, 305–593–2260
Universal Toxicology Laboratories (Florida), LLC, 5361 NW 33rd Avenue, Fort Lauderdale, FL 33309, 954–717–0300, 800–419–7187x419 (Formerly: Integrated Regional Laboratories, Cedars Medical Center, Department of Pathology)
Universal Toxicology Laboratories, LLC, 9930 W. Highway 80, Midland, TX 79706, 915–561–8851/888–953–8851
US Army Forensic Toxicology Drug Testing Laboratory, Fort Meade, Building 2490, Wilson Street, 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. DHHS, with the DHHS' National Laboratory Certification Program (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, the DHHS will recommend that DOT certify the laboratory (**Federal Register**, 16 July 1996) as meeting the minimum standards of the ''Mandatory Guidelines for Workplace Drug Testing'' (59 FR, 9 June 1994, Pages 29908–29931). After receiving the DOT certification, the laboratory will be included in the monthly list of DHHS certified laboratories and participate in the NLCP certification maintenance program.

Richard Kopanda,

Executive Officer, Substance Abuse and Mental Health Services Administration.

[FR Doc. 02–277 Filed 1–4–02; 8:45 am]

BILLING CODE 4160–20–P

DEPARTMENT OF THE INTERIOR

Office of the Secretary

Establishment of the Battle of Midway National Memorial Planning Committee

AGENCY: Office of the Secretary, Interior.

ACTION: Notice of establishment.

SUMMARY: We are publishing this notice in accordance with section 9a of the Federal Advisory Committee Act (Public Law 92–463). Following consultation with the General Services Administration, the Secretary of the Interior hereby establishes the Battle of Midway National Memorial Advisory Committee. The Committee will develop a strategy for a public dedication of the memorial, identify and plan for appropriate exhibits to commemorate this important event, and offer recommendations on improving visitor services on Midway Atoll National Wildlife Refuge.

DATES: On January 22, 2002, we will file a copy of the charter with the Committee on Environment and Public Works, United States Senate; Committee on Resources, House of Representatives; General Services Administration; and Library of Congress.

ADDRESSES: You may submit comments to Barbara Maxfield, Fish and Wildlife Service, P.O. Box 50617, Honolulu, Hawaii, 96850–5167, phone number (808) 541–1201.

FOR FURTHER INFORMATION CONTACT: Barbara Maxfield, U.S. Fish and Wildlife Service, (808) 541–1201.

SUPPLEMENTARY INFORMATION: The Committee will provide advice to the Secretary of the Interior through the Director, Fish and Wildlife Service on the management of the Battle of Midway National Memorial. The FY 2000 Interior Appropriations bill directed us to designate the Battle of Midway National Memorial on the Midway Atoll National Wildlife Refuge to commemorate the pivotal World War II Battle of Midway. The appropriations language also directed that we consult on a regular basis with other agencies and organizations on the management of the national memorial.

The Committee will be comprised of representatives from the Fish and Wildlife Service, National Park Service, Naval Historical Center, International Midway Memorial Foundation, Inc., Midway-Phoenix Corporation, Sixth Defense Battalion, the National Wildlife Refuge Association, Friends of Midway Atoll National Wildlife Refuge, National Trust for Historic Preservation, and a member of the Battle of Midway veterans' community. These agencies, organizations, and the veteran have demonstrated an interest and expertise in commemorating and preserving historical features associated with the Battle of Midway and reflect a balanced, cross-sectional representation of public and private sector organizations.

The Committee will function solely as an advisory body and in compliance with provisions of the Federal Advisory Committee Act.

The Certification for establishment of the committee is published below.

Certification

I hereby certify that the Battle of Midway National Memorial Planning Committee is necessary and in the public interest in connection with the performance of duties imposed on the Department of the Interior by the Consolidated Appropriations Act for FY 2000, the National Historic Preservation Act of 1966, as amended, and the National Wildlife Refuge System Improvement Act of 1997. The Committee will assist the Fish and Wildlife Service by providing advice and developing recommendations for the long-term management and interpretation of the Battle of Midway National Memorial.

Dated: October 11, 2001.

Gale A. Norton,

Secretary of the Interior.

[FR Doc. 02–293 Filed 1–4–02; 8:45 am]

BILLING CODE 4310–55–P

DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

Information Collections for Approval Under the Paperwork Reduction Act for Neotropical Migratory Bird Conservation Act Program

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice of information collection; request for comments.

SUMMARY: The collection of information described below has been submitted to the Office of Management and Budget (OMB) for emergency approval under the provisions of the Paperwork Reduction Act of 1995, and received OMB approval number 1018–0113 with an expiration date of 6/30/2002. Copies of the specific information collection requirements, related forms and explanatory material may be obtained by contacting the Service Information Collection Clearance Officer at the address provided below.

DATES: Consideration will be given to all comments received on or before March 8, 2002. OMB has up to 60 days to approve or disapprove information collection but may respond after 30 days. Therefore, to ensure maximum consideration, OMB should receive public comments by the above referenced date.

ADDRESSES: Comments and suggestions on the requirement should be sent to Rebecca Mullin, Service Information Collection Clearance Officer, U.S. Fish and Wildlife Service, ms 860—ARLSQ, 1849 C Street, NW, Washington, DC 20240.

FOR FURTHER INFORMATION CONTACT: To request a copy of the information collection request, explanatory information and related forms, contact Rebecca A. Mullin at 703/358–2287, or electronically to *rmullin@fws.gov.*

SUPPLEMENTARY INFORMATION:

The OMB regulations at 5 CFR part 1320, which implement provisions of the Paperwork Reduction Act of 1995 (Pub. L. 104–13), require that interested members of the public and affected agencies have an opportunity to comment on information collection and recordkeeping activities (*see* 5 CFR 1320.8(d)). On December 19, 2001, the U.S. Fish and Wildlife Service (Service) provided information to OMB for collection of information in order to begin a grants program conducted under the Neotropical Migratory Bird Conservation Act (Public Law 106–247). The assigned OMB information collection control number is 1018–[to be assigned] , and temporary approval expires on [unknown]. The Service is requesting a three year term of approval for this information collection activity. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

Comments are invited on : (1) Whether the collection of information is