

ESTIMATES OF ANNUAL BURDEN HOURS

Type of respondents	Survey instrument	Number of respondents	Frequency of response	Average time per response (hours)	Annual burden hours
Clinical Trials	Initial Registration	5,500	1	2.0	11,000
	Amendment	5,500	4	1.0	22,000
Total					33,000

Request for Comments: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Attention: NIH Desk Officer, Office of Management and Budget, at OIRA_submission@omb.eop.gov or by fax to 202-395-6974. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact John Speakman, Associate Director for Clinical Trials Products and Programs, Center for Biomedical Informatics and Information Technology, National Cancer Institute, NIH, DHHS, 2115 E. Jefferson Street, Suite 6000, Rockville, MD 20892 or call non-toll-free number 301-451-8786 or e-mail your request, including your address to: john.speakman@nih.gov.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 15 days of the date of this publication.

Dated: June 9, 2009.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. E9-14089 Filed 6-15-09; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Office of Refugee Resettlement; Single-Source Program Expansion Supplement to the Lutheran Social Services of South Dakota (LSS-SD) Under the South Dakota Wilson-Fish Program, Award

AGENCY: Office of Refugee Resettlement, ACF, HHS.

ACTION: Notice to award a single-Source program expansion supplement to the Lutheran Social Services of South Dakota (LSS-SD) under the South Dakota Wilson-Fish Program.

CFDA Number: 93.583.

Legislative Authority: The Refugee Act of 1980 as amended, Wilson-Fish Amendment, 8 U.S.C. 1522(e)(7); section 412(e)(7)(A) of the Immigration and Nationality Act.

Amount of Award: \$125,000.

Project Period: 09/30/2007-09/29/2010

Justification for the Exception to Competition: The Wilson-Fish program is an alternative to the traditional State-administered refugee assistance program for providing integrated assistance and services to refugees, asylees, Amerasian Immigrants, Cuban and Haitian Entrants, Trafficking Victims and Iraqi/Afghani SIV's. South Dakota is one of 12 sites that has chosen this alternative approach.

The supplemental funds will allow the grantee, LSS-SD, located in Sioux Falls, SD, to provide refugee cash assistance through the end of this fiscal year to eligible refugees (and others eligible for refugee benefits) under the South Dakota Wilson-Fish Program.

The primary reason for the grantee's supplemental request is a higher number of arrivals than anticipated when the grantee's budget was submitted and approved last year. The Refugee Act of 1980 mandates that the Office of Refugee Resettlement (ORR) reimburse States and Wilson-Fish projects for the costs of cash and medical assistance for newly arriving

refugees. Since 1991, ORR has reimbursed States and Wilson-Fish agencies for providing cash and medical assistance to eligible individuals during their first eight months in the United States. Hence, the supplement is consistent with the purposes of the Wilson-Fish Program, the Refugee Act of 1980, and ORR policy.

CONTACT FOR FURTHER INFORMATION: Carl Rubenstein, Wilson-Fish Program Manager, Office of Refugee Resettlement, Aerospace Building, 8th Floor West, 901 D Street, SW., Washington, DC 20447. Telephone: 202-205-5933, *E-mail:* crubenstein@acf.hhs.gov.

Dated: 06/04/2009.

David H. Siegel,

Acting Director, Office of Refugee Resettlement.

[FR Doc. E9-14140 Filed 6-15-09; 8:45 am]

BILLING CODE 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://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:

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, 345 Hill Ave., Nashville, TN 37210, 615-255-

2400 (Formerly: Aegis Sciences Corporation, Aegis Analytical Laboratories, Inc.).
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).
Clendo Reference Laboratory, Avenue Santa Cruz #58, Bayamon, Puerto Rico 00959, 787-620-9095.
Clinical Reference Lab, 8433 Quivira Road, Lenexa, KS 66215-2802, 800-445-6917.
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.
DynaLIFE Dx*, 10150-102 St., Suite 200, Edmonton, Alberta, Canada T5J 5E2, 780-451-3702/800-661-9876 (Formerly: Dynacare Kasper Medical Laboratories).
ElSohly Laboratories, Inc., 5 Industrial Park Drive, Oxford, MS 38655, 662-236-2609.
Gamma-Dynacare Medical Laboratories*, A Division of the Gamma-Dynacare Laboratory Partnership, 245 Pall Mall Street, London, ONT, Canada N6A 1P4, 519-679-1630.
Kroll Laboratory Specialists, Inc., 1111 Newton St., Gretna, LA 70053, 504-361-9898/800-433-3823 (Formerly: Laboratory Specialists, Inc.).
Kroll Laboratory Specialists, Inc., 450 Southlake Blvd., Richmond, VA 23236, 804-378-9130 (Formerly: Scientific Testing Laboratories, Inc.; Kroll 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, 1120 Main Street, Southaven, MS 38671, 866-827-8042/800-233-6339 (Formerly: LabCorp 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.).
Maxxam Analytics*, 6740 Campobello Road, Mississauga, ON, Canada L5N 2L8, 905-817-5700, (Formerly: Maxxam Analytics Inc., 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).
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, 509-755-8991/800-541-7891x7.
Phamatech, Inc., 10151 Barnes Canyon Road, San Diego, CA 92121, 858-643-5555.
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, 400 Egypt Road, Norristown, PA 19403, 610-631-4600/877-642-2216 (Formerly: SmithKline Beecham Clinical Laboratories; SmithKline Bio-Science Laboratories).
Quest Diagnostics Incorporated, 7600 Tyrone Ave., Van Nuys, CA 91405, 866-370-6699/818-989-2521 (Formerly: SmithKline Beecham Clinical Laboratories).
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, 4625 E. Cotton Center Boulevard, Suite 177, Phoenix, AZ 85040, 602-438-8507/800-279-0027.

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.

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 following laboratory voluntarily withdrew from the NLCP on May 30, 2009:

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

*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.

Dated: June 8, 2009.

Elaine Parry,

Director, Office of Program Services, SAMHSA.

[FR Doc. E9-14084 Filed 6-15-09; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket Numbers NIOSH-083 Supplied Air Respirators, NIOSH 148 Air Fed Ensembles, NIOSH-168 Total Inward Leakage (for respirators other than filtering facepieces and halfmasks)]

Notice of Public Meeting To Discuss NIOSH's Respirator Standards Development Efforts

Authority: 29 U.S.C. 651 et seq.

AGENCY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of a public meeting.

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and prevention (CDC), will conduct a public meeting to discuss current respirator standards development projects for Supplied Air Respirators (SAR); Air Fed Ensembles; and Total Inward Leakage (TIL) for respirators other than filtering facepieces and halfmasks. There will be an opportunity for discussion following NIOSH's presentations and an accompanying poster session.

Public Meeting Time and Date: 8:30 a.m. to 5 p.m., September 17, 2009. On-site registration will be held beginning at 7:45 a.m.

Place: Hyatt Regency Pittsburgh International Airport, 1111 Airport Boulevard, Pittsburgh, PA 15231. Interested parties should make hotel reservations directly with the Hyatt Regency Pittsburgh International Airport by calling (800) 233-1234, before the cut-off date of September 2, 2009. You must reference the NIOSH room block to receive the special group rate of \$114.00 per night that has been negotiated for meeting guests.

Status: The meeting will be open to the public, limited only by the space available. The meeting room accommodates approximately 200 people.

Instructions: Requests to make presentations at the public meeting

should be mailed to the NIOSH Docket Office, Robert A. Taft Laboratories, MS-C34, 4676 Columbia Parkway, Cincinnati, OH 45226. Requests may also be submitted by telephone (513) 533-8611, facsimile (513) 533-8285, or e-mailed to niocindocket@cdc.gov. All requests to present should contain the name, address, telephone number, and relevant business affiliations of the presenter, topic of the presentation, and the approximate time requested for the presentation. Oral presentations should be limited to 15 minutes.

After reviewing the requests for presentations, NIOSH will notify the presenter that his/her presentation is scheduled. If a participant is not present when their presentation is scheduled to begin, the remaining participants will be heard in order. At the conclusion of the meeting, an attempt will be made to allow presentations by any scheduled participants who missed their assigned times. Attendees who wish to speak but did not submit a request for the opportunity to make a presentation may be given this opportunity at the conclusion of the meeting, at the discretion of the presiding officer.

This meeting will also be using Audio/LiveMeeting Conferencing, remote access capabilities where interested parties may listen in and review the presentations over the internet simultaneously. Parties remotely accessing the meeting will have the opportunity to ask questions during the open comment period. To register to use this capability, please contact the National Personal Protective Technology Laboratory (NPPTL), Policy and Standards Development Branch, Post Office Box 18070, 626 Cochran's Mill Road, Pittsburgh, PA 15236, telephone (412) 386-5200, facsimile (412) 386-4089. This option will be available to participants on a first come, first serve basis and is limited to the first 50 participants.

Background: NIOSH, National Personal Protective Technology Laboratory (NPPTL), will present information to attendees concerning the development of the concepts being considered regarding updated performance criteria for the various classes of respirators in 42 Code of Federal Regulations, Part 84. Participants will be given an opportunity to ask questions and to present individual comments that they may wish to have considered.

FOR FURTHER INFORMATION CONTACT: Jonathan Szalajda, NPPTL, Policy and Standards Development Branch, Post Office Box 18070, 626 Cochran's Mill Road, Pittsburgh, PA 15236, telephone