these diseases. Hispanics, now the largest minority population in the U.S., are influenced by factors associated with immigration from different cultural settings and environments, including changes in diet, activity, community support, working conditions, and health care access. This project is a multicenter, six-and-a-half year epidemiologic study and will recruit 16,000 Hispanic men and women aged 18–74 in four community-based cohorts in Chicago, Miami, San Diego, and the Bronx. The study will also examine measures of obesity, physical activity, nutritional habits, diabetes, lung and sleep function, cognitive function, hearing, and dental conditions. Closely integrated with the research component will be a community and professional education component, with the goals of bringing the research results back to the community, improving recognition and control of risk factors, and attracting and training Hispanic researchers in epidemiology and population-based research. Frequency of Response: The participants will be contacted annually. Affected Public: Individuals or households; Businesses or other for profit; Small businesses or organizations. Type of Respondents: Individuals or households; physicians. The annual reporting burden is as follows: Estimated Number of Respondents: 10,801; Estimated Number of Responses per Respondent: 1.0; Average Burden Hours Per Response: 3.6; and Estimated Total Annual Burden Hours Requested: 38,401. The annualized cost to respondents is estimated at $506,613, assuming respondents time at the rate of $13 per hour and physician time at the rate of $50 per hour. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

ESTIMATE OF ANNUAL HOUR BURDEN

<table>
<thead>
<tr>
<th>Type of response</th>
<th>Number of respondents</th>
<th>Frequency of response</th>
<th>Average hours per response</th>
<th>Annual hour burden</th>
</tr>
</thead>
<tbody>
<tr>
<td>Participant Examinations and Questionnaires</td>
<td>5,334</td>
<td>1.0</td>
<td>6.5</td>
<td>34,671</td>
</tr>
<tr>
<td>Physician, Medical Examiner, and Next-of-kin Follow-up 1</td>
<td>200</td>
<td>1.0</td>
<td>1.0</td>
<td>200</td>
</tr>
<tr>
<td>Total</td>
<td>10,801</td>
<td></td>
<td></td>
<td>38,401</td>
</tr>
</tbody>
</table>

1 Annual burden is placed on doctors and respondent relatives/informants through requests for information which will help in the compilation of the number and nature of new fatal and nonfatal events.

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) 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) 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) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to 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.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Dr. Larissa Aviles-Santa, Deputy Project Officer, NIH, NHLBI, 6701 Rockledge Drive, MSC 7936, Bethesda, MD 20892–7934, or call non-toll-free number 301–435–1284 or E-mail your request, including your address to: AvilesantaL@NHLBI.NIH.GOV.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.


Peter Savage, Acting Director, DPPS.

Suzanne Freeman, NHLBI Project Clearance Liaison, National Institutes of Health.

[FR Doc. E7–13384 Filed 7–10–07; 8:45 am]

BILLING CODE 4140–01–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 (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 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 dated April 13, 2004 which attests that it has met minimum standards. Mandatory Guidelines dated April 13, 2004 (Formerly: HHS/NIDA) which attests that it has met minimum standards.

In accordance with Subpart C of the HHS Mandatory Guidelines dated April 13, 2004 (Formerly: HHS/NIDA) which attests that it has met minimum standards.

- 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).
- Doctors Laboratory, Inc., 2906 Julia Ave., Nashville, TN 37210, 615–526–2400/800–437–4986 (Formerly: Roche Biomedical 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, 13112 Evening Creek Drive, Suite 100, San Diego, CA 92128, 858–668–3710/800–882–7272 (Formerly: Poisonlab, Inc.).
- Laboratory Corporation of America Holdings, 550 17th Ave., Suite 300, Seattle, WA 98122, 206–923–7020/800–896–0180 (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).
- 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 Inc., 6740 Campbell Road, Mississauga, ON, Canada L5N 2L8, 905–817–5700 (Formerly: NOVAMANN (Ontario), Inc.).
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.


U.S. 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 June 19, 2007:


* 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 24924). 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.

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

DEPARTMENT OF THE INTERIOR
Office of the Secretary

DEPARTMENT OF THE INTERIOR
Fish and Wildlife Service

Proposed Information Collection; Export of Fertilized Live Eggs, Caviar, or Meat from Aquacultured Paddlefish or Sturgeon (CITES)

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice; request for comments.

SUMMARY: We (Fish and Wildlife Service) will ask the Office of Management and Budget (OMB) to approve the information collection (IC) described below. As required by the Paperwork Reduction Act of 1995 and as part of our continuing efforts to reduce paperwork and respondent burden, we invite the general public and other Federal agencies to take this opportunity to comment on this IC.

DATES: You must submit comments on or before September 10, 2007.

ADDRESSES: Send your comments on the IC to Hope Grey, Information Collection Clearance Officer, Fish and Wildlife Service, MS 222–ARLSQ, 4401 North Fairfax Drive, Arlington, VA 22203 (mail); hope.greyserv@fws.gov (e-mail); or (703) 358–2269 (fax).

FOR FURTHER INFORMATION CONTACT: To request additional information about this IC, contact Hope Grey by mail, fax, or e-mail (see ADDRESSES) or by telephone at (703) 358–2482.

SUPPLEMENTARY INFORMATION:

I. Abstract

This information collection is associated with regulations implementing the Convention on International Trade in Endangered Species of Wild Fauna and Flora (CITES). CITES regulates international trade in listed species through a system of permits and certificates. We assess permit requests according to criteria in CITES and Federal regulations (50 CFR parts 13 and 23) for the issuance, suspension, revocation, or denial of permits.

We have developed a new permit application form (FWS Form 3–200–80) specific to permit requests for the export of fertilized live eggs, caviar, or meat from aquacultured paddlefish or sturgeon. In the past, we have used FWS Form 3–200–24 (Export of Captive Born Wildlife) to collect the information necessary for us to evaluate these permit requests. When using that general form, applicants have had considerable difficulty understanding what information is necessary and how to supply it. The proposed form clarifies these issues. The information we plan to collect includes, but is not limited to: