practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

**Proposed Project**

**HRSA Competing Training Grant Application, Instructions and Related Regulations—(0915–0060)—Extension and Revision**

The Health Resources and Services Administration uses the information in the application to determine the eligibility of applicants for awards, to calculate the amount of each award, and to judge the relative merit of applications. This is a request for renewed clearance with several changes in the application form. The form will be distributed electronically via the Internet; the budget will be negotiated for all years of the project period based on this application, and program-specific instructions will include greater standardization of content for the project summary and the detailed description of the project. Regulations which authorize the application form and other reporting requirements for various programs are cleared in this package. No changes were made to the regulations.

The estimated annual application burden is as follows:

<table>
<thead>
<tr>
<th>Type of collection</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden response (hours)</th>
<th>Total burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Basic Application</td>
<td>1769</td>
<td>1</td>
<td>61.25</td>
<td>108,351</td>
</tr>
<tr>
<td>Statutory Requirements*</td>
<td>1121</td>
<td>1</td>
<td>105</td>
<td>117,705</td>
</tr>
</tbody>
</table>

* In 1992, a law was passed which required applicants for selected grant programs to provide specified data in the grant application.

The burden for the regulatory requirements included in this package are as follows:

<table>
<thead>
<tr>
<th>Type of requirement</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden response (hours)</th>
<th>Total burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Reporting Requirements</td>
<td>28</td>
<td>1.4</td>
<td>1</td>
<td>39</td>
</tr>
<tr>
<td>Disclosure Requirements</td>
<td>148</td>
<td>1.4</td>
<td>3.3</td>
<td>669</td>
</tr>
</tbody>
</table>

The total burden for these activities is 226,934 hours.

Send comments to Patricia Royston, HRSA Reports Clearance Officer, Room 14–36, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: August 29, 1996.

J. Henry Montes,
Associate Administrator for Policy Coordination.

[FR Doc. 96–22605 Filed 9–4–96; 8:45 am]

**BILLING CODE 4160–15–P**

**Substance Abuse and Mental Health Services Administration**

**Current List of Laboratories Which Meet Minimum Standards To Engage in Urine Drug Testing for Federal Agencies, and Laboratories That Have Withdrawn From the Program**

**AGENCY:** Substance Abuse and Mental Health Services Administration, HHS (Formerly: National Institute on Drug Abuse, ADAMHA, 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 similar notice listing all currently certified laboratories will be published during the first week of each month, and updated to include laboratories which subsequently apply for and complete the certification process. If any listed laboratory’s certification is totally suspended or revoked, the laboratory will be omitted from updated 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 identified as such at the end of the current list of certified laboratories, and will be omitted from the monthly listing thereafter.

**FOR FURTHER INFORMATION CONTACT:** Mrs. Giselle Hersh, Division of Workplace Programs, Room 13A–54, 5600 Fishers Lane, Rockville, Maryland 20857; Tel.: (301) 443–6014.

**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:

American Medical Laboratories, Inc., 14225 Newbrook Dr., Chantilly, VA 22021, 703–802–6200
Associated Regional and University Pathologists, Inc. (ARUP), 500 Chipeta Way, Salt Lake City, UT 84108, 801–583–2787
Baptist Medical Center—Toxicology Laboratory, 9601 S. Sepulveda Blvd., Los Angeles, CA 90045, 310–215–6020
Centinelia Hospital Airport Toxicology Laboratory, 9601 S. Sepulveda Blvd., Los Angeles, CA 90045, 310–215–6020
Clinical Reference Laboratory, One Malcolm Ave., Teterboro, NJ 07608, 201–393–5000 (formerly: MetPath, Inc., CORNING MetPath Clinical Laboratories)
CORNING Clinical Laboratory, 4022 Willow Lake Blvd., Memphis, TN 38175, 901–795–1515
Notice of Availability of a Draft Recovery Plan for the Alabama Cave Shrimp (Palaemonias alabamae) for Review and Comment

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

**ACTION:** Notice of document availability and public comment period.

**SUMMARY:** The U.S. Fish and Wildlife Service (Service) announces the availability for public review of a draft recovery plan for the Alabama cave shrimp (Palaemonias alabamae). The albino Alabama cave shrimp has been found in five caves (three cave systems) near the city of Huntsville, Madison County, Alabama. One cave is found on the Redstone Arsenal, an army installation, while the other four caves are privately owned. The Service solicits review and comment from the public on this draft plan.

**DATES:** Comments on the draft recovery plan must be received on or before November 15, 1996, to receive consideration by the Service.

**ADDRESSES:** Persons wishing to review the draft recovery plan may obtain a copy by contacting the Jackson Field Office, U.S. Fish and Wildlife Service, 6578 Dogwood View Parkway, Suite A, Jackson, Mississippi 39213. Written comments and materials regarding the plan should be addressed to the Field Supervisor at the above address. Comments and materials received are available on request for public inspection, by appointment, during normal business hours at the above address.

**FOR FURTHER INFORMATION CONTACT:** Ms. Theresa Jacobson at the above address (601–965–4900, ext. 30).

**SUPPLEMENTARY INFORMATION:**

**Background**

Restoring endangered or threatened animals and plants to the point where they are again secure, self-sustaining members of their ecosystems is a primary goal of the U.S. Fish and Wildlife Service’s endangered species program. To help guide the recovery effort, the Service is working to prepare recovery plans for most of the listed species native to the United States. Recovery plans describe actions considered necessary for conservation of