

**SUPPLEMENTARY INFORMATION:** Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: written comments and/or suggestions from the public and affected agencies are invited to address 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.

*Proposed Collection Title:* Early Career Reviewer Program Online Application and Vetting System, 0925–0695, REVISION, exp., date 05/31/2020, Center for Scientific Review (CSR), National Institutes of Health (NIH).

*Need and Use of Information Collection:* The Center for Scientific Review (CSR) is the portal for NIH grant applications and their review for scientific merit. Our mission is to see that all NIH grant applications receive fair, independent, expert, and timely reviews—free from inappropriate influences—so NIH can fund the most promising research. To accomplish this goal, Scientific Review Officers (SRO) form study sections consisting of scientists who have the technical and scientific expertise to evaluate the merit of grant applications. Study section members are generally scientists who have established independent programs of research as demonstrated by their publications and their grant award experiences.

The CSR Early Career Reviewer program was developed to identify and train qualified scientists who are early in their scientific careers and who have not had prior CSR review experience. The goals of the program are to expose these early career scientists to the peer review experience so that they become more competitive as applicants as well as to enrich the existing pool of NIH reviewers. Currently, the online application software, the Early Career

Reviewer Application and Vetting System, is accessed online by applicants to the Early Career Reviewer Program who provide information such as their name, contact information, a description of their areas of expertise, their study section preferences, and their professional Curriculum Vitae. This Information Collection Request (ICR) is to revise the Early Career Reviewer Application and Vetting System to include additional questions and be more user friendly. Additional questions are in line with NIH's renewed Interest in Diversity (NOT–OD–20–031) and include questions such as applicants' race, ethnicity, gender, disability, and disadvantage backgrounds. Applicants can choose if they would like to answer these additional questions (*i.e.* optional). Applicants are also now able to check their eligibility before applying to the program.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 505.

#### ESTIMATED ANNUALIZED BURDEN HOURS

| Type of respondent        | Number of respondents | Number of responses per respondent | Average time per response (in hours) | Total annual burden hour |
|---------------------------|-----------------------|------------------------------------|--------------------------------------|--------------------------|
| Research scientists ..... | 1212                  | 1                                  | 25/60                                | 505                      |
| Total .....               | .....                 | 1212                               | .....                                | 505                      |

Dated: January 22, 2020.

**Hope M. Cummings,**

*Project Clearance Liaison, Center for Scientific Review (CSR), National Institutes of Health.*

[FR Doc. 2020–01798 Filed 1–30–20; 8:45 am]

**BILLING CODE 4140–01–P**

#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

##### Substance Abuse and Mental Health Services Administration

##### Current List of HHS-Certified Laboratories and Instrumented Initial Testing Facilities Which Meet Minimum Standards To Engage in Urine and Oral Fluid 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 and Instrumented Initial Testing Facilities (IITFs) currently certified to meet the standards of the Mandatory Guidelines for Federal Workplace Drug Testing Programs using Urine or Oral Fluid (Mandatory Guidelines).

A notice listing all currently HHS-certified laboratories and IITFs is published in the **Federal Register** during the first week of each month. If any laboratory or IITF certification is suspended or revoked, the laboratory or IITF will be omitted from subsequent lists until such time as it is restored to full certification under the Mandatory Guidelines.

If any laboratory or IITF 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 <https://www.samhsa.gov/workplace/resources/drug-testing/certified-lab-list>.

##### FOR FURTHER INFORMATION CONTACT:

Anastasia Donovan, Division of Workplace Programs, SAMHSA/CSAP, 5600 Fishers Lane, Room 16N06B, Rockville, Maryland 20857; 240–276–2600 (voice); [Anastasia.Donovan@samhsa.hhs.gov](mailto:Anastasia.Donovan@samhsa.hhs.gov) (email).

**SUPPLEMENTARY INFORMATION:** The Department of Health and Human Services (HHS) notifies federal agencies of the laboratories and Instrumented Initial Testing Facilities (IITFs) currently certified to meet the standards of the Mandatory Guidelines for Federal Workplace Drug Testing Programs (Mandatory Guidelines) using Urine and of the laboratories currently certified to meet the standards of the Mandatory Guidelines using Oral Fluid.

The Mandatory Guidelines using Urine 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); September 30, 1997 (62 FR 51118); April 13, 2004 (69 FR 19644); November 25, 2008 (73 FR 71858); December 10, 2008 (73 FR 75122); April 30, 2010 (75 FR 22809); and on January 23, 2017 (82 FR 7920).

The Mandatory Guidelines using Oral Fluid were first published in the **Federal Register** on October 25, 2019 (84 FR 57554) with an effective date of January 1, 2020.

The Mandatory Guidelines were initially developed in accordance with Executive Order 12564 and section 503 of Public Law 100–71 and allowed urine drug testing only. The Mandatory Guidelines using Urine have since been revised, and new Mandatory Guidelines allowing for oral fluid drug testing have been published. The Mandatory Guidelines require strict standards that laboratories and IITFs must meet in order to conduct drug and specimen validity tests on specimens for federal agencies. HHS does not allow IITFs for oral fluid testing.

To become certified, an applicant laboratory or IITF must undergo three rounds of performance testing plus an on-site inspection. To maintain that certification, a laboratory or IITF must participate in a quarterly performance testing program plus undergo periodic, on-site inspections.

Laboratories and IITFs in the applicant stage of certification are not to be considered as meeting the minimum requirements described in the HHS Mandatory Guidelines using Urine and/or Oral Fluid. An HHS-certified laboratory or IITF must have its letter of certification from HHS/SAMHSA (formerly: HHS/NIDA), which attests that the test facility has met minimum standards. HHS does not allow IITFs for oral fluid testing.

#### **HHS-Certified Laboratories Certified To Conduct Oral Fluid Drug Testing**

In accordance with the Mandatory Guidelines using Oral Fluid dated October 25, 2019 (84 FR 57554), the following HHS-certified laboratories meet the minimum standards to conduct drug and specimen validity tests on oral fluid specimens:

At this time, there are no laboratories certified to conduct drug and specimen validity tests on oral fluid specimens.

#### **HHS-Certified Instrumented Initial Testing Facilities Certified To Conduct Urine Drug Testing**

In accordance with the Mandatory Guidelines using Urine dated January 23, 2017 (82 FR 7920), the following

HHS-certified IITFs meet the minimum standards to conduct drug and specimen validity tests on urine specimens:

Dynacare, 6628 50th Street NW, Edmonton, AB Canada T6B 2N7, 780–784–1190, (Formerly: Gamma-Dynacare Medical Laboratories)

#### **HHS-Certified Laboratories Certified To Conduct Urine Drug Testing**

In accordance with the Mandatory Guidelines using Urine dated January 23, 2017 (82 FR 7920), the following HHS-certified laboratories meet the minimum standards to conduct drug and specimen validity tests on urine specimens:

Alere Toxicology Services, 1111 Newton St., Gretna, LA 70053, 504–361–8989/800–433–3823, (Formerly: Kroll Laboratory Specialists, Inc., Laboratory Specialists, Inc.)

Alere Toxicology Services, 450 Southlake Blvd., Richmond, VA 23236, 804–378–9130, (Formerly: Kroll Laboratory Specialists, Inc., Scientific Testing Laboratories, Inc.; Kroll Scientific Testing Laboratories, Inc.)

Clinical Reference Laboratory, Inc., 8433 Quivira Road, Lenexa, KS 66215–2802, 800–445–6917

Cordant Health Solutions, 2617 East L Street, Tacoma, WA 98421, 800–442–0438, (Formerly: STERLING Reference Laboratories)

Desert Tox, LLC, 10221 North 32nd Street Suite J, Phoenix, AZ 85028, 602–457–5411

DrugScan, Inc., 200 Precision Road, Suite 200, Horsham, PA 19044, 800–235–4890

Dynacare \*, 245 Pall Mall Street, London, ONT, Canada N6A 1P4, 519–

\* 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 January 23, 2017 (82 FR 7920). 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.

679–1630, (Formerly: Gamma-Dynacare Medical Laboratories)  
ElSohly Laboratories, Inc., 5 Industrial Park Drive, Oxford, MS 38655, 662–236–2609

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 TW 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.)

Legacy Laboratory Services Toxicology, 1225 NE 2nd Ave., Portland, OR 97232, 503–413–5295/800–950–5295, (Formerly: Legacy Laboratory Services—MetroLab)

MedTox Laboratories, Inc., 402 W County Road D, St. Paul, MN 55112, 651–636–7466/800–832–3244

Minneapolis Veterans Affairs Medical Center, Forensic Toxicology Laboratory, 1 Veterans Drive, Minneapolis, MN 55417, 612–725–2088, Testing for Veterans Affairs (VA) Employees Only

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., 15175 Innovation Drive, San Diego, CA 92128, 888–635–5840

Quest Diagnostics Incorporated, 1777 Montreal Circle, Tucker, GA 30084, 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)  
Redwood Toxicology Laboratory, 3700  
Westwind Blvd., Santa Rosa, CA  
95403, 800-255-2159  
U.S. Army Forensic Toxicology Drug  
Testing Laboratory, 2490 Wilson St.,  
Fort George G. Meade, MD 20755-  
5235, 301-677-7085, Testing for  
Department of Defense (DoD)  
Employees Only

Anastasia Marie Donovan,  
*Policy Analyst.*

[FR Doc. 2020-01776 Filed 1-30-20; 8:45 am]

BILLING CODE 4160-20-P

## DEPARTMENT OF HOMELAND SECURITY

### Federal Emergency Management Agency

[Internal Agency Docket No. FEMA-4427-  
DR; Docket ID FEMA-2020-0001]

#### Tennessee; Amendment No. 2 to Notice of a Major Disaster Declaration

**AGENCY:** Federal Emergency  
Management Agency, DHS.

**ACTION:** Notice.

**SUMMARY:** This notice amends the notice  
of a major disaster declaration for State  
of Tennessee (FEMA-4427-DR), dated  
April 17, 2019, and related  
determinations.

**DATES:** This change occurred on January  
7, 2020.

**FOR FURTHER INFORMATION CONTACT:**  
Dean Webster, Office of Response and  
Recovery, Federal Emergency  
Management Agency, 500 C Street SW,  
Washington, DC 20472, (202) 646-2833.

**SUPPLEMENTARY INFORMATION:** The  
Federal Emergency Management Agency  
(FEMA) hereby gives notice that  
pursuant to the authority vested in the  
Administrator, under Executive Order  
12148, as amended, Myra M. Shird, of  
FEMA is appointed to act as the Federal  
Coordinating Officer for this disaster.

This action terminates the  
appointment of Manny J. Toro as  
Federal Coordinating Officer for this  
disaster.

The following Catalog of Federal Domestic  
Assistance Numbers (CFDA) are to be used  
for reporting and drawing funds: 97.030,  
Community Disaster Loans; 97.031, Cora  
Brown Fund; 97.032, Crisis Counseling;

97.033, Disaster Legal Services; 97.034,  
Disaster Unemployment Assistance (DUA);  
97.046, Fire Management Assistance Grant;  
97.048, Disaster Housing Assistance to  
Individuals and Households in Presidentially  
Declared Disaster Areas; 97.049,  
Presidentially Declared Disaster Assistance—  
Disaster Housing Operations for Individuals  
and Households; 97.050, Presidentially  
Declared Disaster Assistance to Individuals  
and Households—Other Needs; 97.036,  
Disaster Grants—Public Assistance  
(Presidentially Declared Disasters); 97.039,  
Hazard Mitigation Grant.

**Pete Gaynor,**

*Acting Administrator, Federal Emergency  
Management Agency.*

[FR Doc. 2020-01897 Filed 1-30-20; 8:45 am]

BILLING CODE 9111-23-P

## DEPARTMENT OF HOMELAND SECURITY

### Federal Emergency Management Agency

[Internal Agency Docket No. FEMA-4469-  
DR; Docket ID FEMA-2019-0001]

#### South Dakota; Amendment No. 1 to Notice of a Major Disaster Declaration

**AGENCY:** Federal Emergency  
Management Agency, DHS.

**ACTION:** Notice.

**SUMMARY:** This notice amends the notice  
of a major disaster declaration for the  
State of South Dakota (FEMA-4469-  
DR), dated November 18, 2019, and  
related determinations.

**DATES:** This amendment was issued  
January 8, 2020.

**FOR FURTHER INFORMATION CONTACT:**  
Dean Webster, Office of Response and  
Recovery, Federal Emergency  
Management Agency, 500 C Street SW,  
Washington, DC 20472, (202) 646-2833.

**SUPPLEMENTARY INFORMATION:** The notice  
of a major disaster declaration for the  
State of South Dakota is hereby  
amended to include the following area  
among those areas determined to have  
been adversely affected by the event  
declared a major disaster by the  
President in his declaration of  
November 18, 2019.

Aurora County for Individual Assistance  
(already designated for Public Assistance).

The following Catalog of Federal Domestic  
Assistance Numbers (CFDA) are to be used  
for reporting and drawing funds: 97.030,  
Community Disaster Loans; 97.031, Cora  
Brown Fund; 97.032, Crisis Counseling;  
97.033, Disaster Legal Services; 97.034,  
Disaster Unemployment Assistance (DUA);  
97.046, Fire Management Assistance Grant;  
97.048, Disaster Housing Assistance to  
Individuals and Households in Presidentially  
Declared Disaster Areas; 97.049,

Presidentially Declared Disaster Assistance—  
Disaster Housing Operations for Individuals  
and Households; 97.050 Presidentially  
Declared Disaster Assistance to Individuals  
and Households—Other Needs; 97.036,  
Disaster Grants—Public Assistance  
(Presidentially Declared Disasters); 97.039,  
Hazard Mitigation Grant.

**Pete Gaynor,**

*Acting Administrator, Federal Emergency  
Management Agency.*

[FR Doc. 2020-01887 Filed 1-30-20; 8:45 am]

BILLING CODE 9111-23-P

## DEPARTMENT OF HOMELAND SECURITY

### Federal Emergency Management Agency

[Internal Agency Docket No. FEMA-4469-  
DR; Docket ID FEMA-2020-0001]

#### South Dakota; Amendment No. 2 to Notice of a Major Disaster Declaration

**AGENCY:** Federal Emergency  
Management Agency, DHS.

**ACTION:** Notice.

**SUMMARY:** This notice amends the notice  
of a major disaster declaration for the  
State of South Dakota (FEMA-4469-  
DR), dated November 18, 2019, and  
related determinations.

**DATES:** This amendment was issued  
January 8, 2020.

**FOR FURTHER INFORMATION CONTACT:**  
Dean Webster, Office of Response and  
Recovery, Federal Emergency  
Management Agency, 500 C Street SW,  
Washington, DC 20472, (202) 646-2833.

**SUPPLEMENTARY INFORMATION:** The notice  
of a major disaster declaration for the  
State of South Dakota is hereby  
amended to include the following areas  
among those areas determined to have  
been adversely affected by the event  
declared a major disaster by the  
President in his declaration of  
November 18, 2019.

Clark, Codington, and Day Counties for  
Public Assistance.

Lincoln County for Public Assistance  
(already designated for Individual  
Assistance).

The following Catalog of Federal Domestic  
Assistance Numbers (CFDA) are to be used  
for reporting and drawing funds: 97.030,  
Community Disaster Loans; 97.031, Cora  
Brown Fund; 97.032, Crisis Counseling;  
97.033, Disaster Legal Services; 97.034,  
Disaster Unemployment Assistance (DUA);  
97.046, Fire Management Assistance Grant;  
97.048, Disaster Housing Assistance to  
Individuals and Households in Presidentially  
Declared Disaster Areas; 97.049,  
Presidentially Declared Disaster Assistance—  
Disaster Housing Operations for Individuals  
and Households; 97.050 Presidentially