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

<table>
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<th>Type of respondent</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average time per response (in hours)</th>
<th>Total annual burden hour</th>
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<td></td>
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</table>

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 (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
Guidelines using Urine dated January
Urine Drug Testing
Testing Facilities Certified To Conduct
HHS-Certified Instrumented Initial
validity tests on oral fluid specimens.

Guidelines using Oral Fluid dated
Conduct Oral Fluid 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:

Dynamac, 6628 50th Street NW,
Edmonton AB Canada T6B 2N7, 780–
784–1190, (Formerly: Gamma-
Dynamac 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 Services, 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

Dynamac *, 245 Pall Mall Street,
London, ONT, Canada N6A 1P4, 519–
679–1630, (Formerly: Gamma-
Dynamac 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., Armitage, 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., CompChem
Laboratories, Inc.; CompChem
Laboratories, Inc., A Subsidiary of
Roche Biomedical Laboratory; Roche
CompChem 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, 913–838–3927/800–873–8845,
(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
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.


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.

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