

consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Board of Scientific Counselors, National Institute of Neurological Disorders and Stroke.

Date: November 8–9, 2027.

Time: November 8, 2027, 9:00 a.m. to 9:00 p.m.

Agenda: To review and evaluate personnel qualifications and performance, and competence of individual investigators.

Address: National Institute of Neurological Disorders and Stroke, National Institutes of Health, Porter Neuroscience Research Center, Building 35A, 35 Convent Drive, Room GF–149 Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Time: November 9, 2027, 10:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate personnel qualifications and performance, and competence of individual investigators.

Address: National Institute of Neurological Disorders and Stroke, National Institutes of Health, Porter Neuroscience Research Center, Building 35A, 35 Convent Drive, Room GF–149 Bethesda, MD 20892

Meeting Format: Virtual Meeting.

Contact Person: Jeffrey S. Diamond, Ph.D., Acting Scientific Director, c/o Caren Collins, Program Analyst, Division of Intramural Research, NINDS, Office of the Scientific Director, National Institute of Neurological Disorders and Stroke, National Institutes of Health, Porter Neuroscience Research Center, Building 35A, 35 Convent Drive, Room GF–149 Bethesda, MD 20892, 301–435–1896, diamondj@ninds.nih.gov; collinsca@ninds.nih.gov.

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

(Catalogue of Federal Domestic Assistance Program Nos. 93.853, Extramural Research Programs in the Neurosciences and Neurological Disorders; 93.854, Biological Basis Research in the Neurosciences, National Institutes of Health, HHS).

Dated: September 28, 2026.

Rosalind M. Niamke,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2026–20059 Filed 9–29–26; 8:45 am]

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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) provides notice 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 the laboratories currently certified to meet the standards of the Mandatory Guidelines using Oral Fluid.

FOR FURTHER INFORMATION CONTACT:

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

SUPPLEMENTARY INFORMATION: The Department of Health and Human Services (HHS) publishes a notice listing all HHS-certified laboratories and Instrumented Initial Testing Facilities (IITFs) in the **Federal Register** monthly, in accordance with Section 9.19 of the Mandatory Guidelines for Federal Workplace Drug Testing Programs (Mandatory Guidelines) using Urine and Section 9.17 of the Mandatory Guidelines using Oral Fluid. 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/drug-testing-resources/certified-lab-list>.

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); January 23, 2017 (82 FR 7920); and on October 12, 2023 (88 FR 70768).

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, and subsequently revised in the **Federal Register** on October 12, 2023 (88 FR 70814).

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 to conduct 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-Certified Laboratories Approved To Conduct Oral Fluid Drug Testing

In accordance with the Mandatory Guidelines using Oral Fluid effective October 10, 2023 (88 FR 70814), 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 Approved To Conduct Urine Drug Testing

In accordance with the Mandatory Guidelines using Urine effective February 1, 2024 (88 FR 70768), 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).

Note: DOT does not allow IITFs to test DOT-regulated specimens.

HHS-Certified Laboratories Approved To Conduct Urine Drug Testing

In accordance with the Mandatory Guidelines using Urine effective February 1, 2024 (88 FR 70768), 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.).

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

Desert Tox, LLC, 5425 E Bell Rd., Suite 125, Scottsdale, AZ, 85254, 602-457-5411/623-748-5045.

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

Dynacare, 245 Pall Mall Street, London, ON, Canada N6A 1P4, 519-679-1630, (Formerly: Gamma-Dynacare Medical Laboratories).

ElSohly Laboratories, Inc., 5 Industrial Park Drive, Oxford, MS 38655, 662-236-2609.

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.).

Laboratory Corporation of America Holdings, 7207 N Gessner Road, Houston, TX 77040, 713-856-8288/800-800-2387.

Laboratory Corporation of America Holdings, 1904 T.W. 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).

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).
Phamatech, Inc., 15175 Innovation Drive, San Diego, CA 92128, 888-635-5840.
US 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.

Carlos R. Castillo,

Public Health Analyst, OAS.

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DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR-7106-N-38]

Privacy Act of 1974; System of Records

AGENCY: Office of Public and Indian Housing, HUD.

ACTION: Notice of a modified system of records.

SUMMARY: Pursuant to the provisions of the Privacy Act of 1974, as amended, the Department of Housing and Urban Development (HUD), Office of Public and Indian Housing (PIH), is modifying the system of records notice for the Inventory Management System (IMS), also known as the Public and Indian Housing Information Center (IMS/PIC), and the Housing Information Portal (HIP). The modification incorporates two routine uses that were previously added by separate notices and adds three new routine uses and updates the system manager and agency contact information. The incorporated and new routine uses are listed as: (17), (18), (19), (20), and (21). The modification also clarifies the Categories of Records in the System, Record Source Categories, and methods of electronic collection and submission for IMS/PIC and HIP. These updates are explained in the **SUPPLEMENTARY INFORMATION** section of this notice. IMS/PIC and HIP serve as a national repository of information related to Public Housing Agencies (PHA)s, Tribally Designated Housing Entities (TDHE), HUD-assisted families, HUD assisted properties, and other HUD programs, for the purpose of monitoring and evaluating the effectiveness of PIH rental housing assistance programs. IMS/PIC and HIP allow PHAs, TDHEs,

and their hired management agents to electronically submit information to HUD that is related to the administration of HUD's PIH programs. They collect data for PIH operations, including data submitted via the internet from HUD's field offices, and accurately track activities and processes. IMS/PIC and HIP also help to increase sharing of information throughout PIH and HUD, which improves staff awareness of activities related to the administration of HUD subsidized housing programs.

DATES: Comments will be accepted on or before October 30, 2026. This proposed action will be effective on the date following the end of the comment period unless comments are received which result in a contrary determination.

ADDRESSES: You may submit comments, identified by docket number or by one of these methods:

Federal e-Rulemaking Portal: <http://www.regulations.gov>. Follow the instructions provided on that site to submit comments electronically.

Fax: 202-619-8365.

Email: privacy@hud.gov.

Mail: Attention: Privacy Office; Kimberly Morton, Acting Chief Privacy Officer, Office of the Executive Secretariat; 451 7th Street SW, Room 10139, Washington, DC 20410-0001.

Instructions: All submissions received must include the agency name and docket number for this notice.

Docket: For access to the docket to read background documents or comments received go to <http://www.regulations.gov>.

FOR FURTHER INFORMATION CONTACT:

Kimberly Morton, Acting Chief Privacy Officer, 451 7th Street SW, Room 10139, Washington, DC 20410-0001, telephone number (804) 822-4801 (this is not a toll-free number). HUD welcomes and is prepared to receive calls from individuals who are deaf or hard of hearing, as well as individuals with speech or communication disabilities. To learn more about how to make an accessible telephone call, please visit <https://www.fcc.gov/consumers/guides/telecommunications-relay-service-trs>.

SUPPLEMENTARY INFORMATION: HUD is publishing this notice to add three new routine uses to the Routine Uses section of this System of Records Notice. The changes also include updates to the system manager and agency contact information. Additional revisions clarify the Categories of Records in the System, Record Source Categories, and methods of electronic collection and submission associated with IMS/PIC and HIP based