DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration


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).

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

SUPPLEMENTARY INFORMATION: In accordance with Section 9.19 of the 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If any laboratory or IITF has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end and will be omitted from the monthly listing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.

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 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 does not allow IITFs to conduct oral fluid testing.

HHS-Certified Laboratories Approved 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 Approved 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 T6E 2N7, 780–784–1190 (Formerly: Gamma-Dynacare Medical Laboratories)

HHS-Certified Laboratories Approved 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.)


- 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)


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

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

- ElSholly 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, 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
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, 800–326–6942 (Formerly: Centinel Hospital Airport Toxicology Laboratory)
Phamatech, Inc., 15175 Innovation Drive, San Diego, CA 92128, 888–635–5840
Quest Diagnostics Incorporated, 400 Egypt Road, Norristown, PA 19403, 610–631–4600/877–642–2216 (Formerly: SmithKline Beecham Clinical Laboratories; SmithKline BioScience Laboratories)
Redwood Toxicology Laboratory, 3700 Westwind Blvd., Santa Rosa, CA 95403, 800–255–2159
US Army Forensic Toxicology Drug Testing Laboratory, 2490 Wilson St., Fort George G. Meade, MD 20755–5255, 301–677–7085, Testing for Department of Defense (DoD) Employees Only
* 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.

Carlos Castillo, Committee Management Officer.

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DEPARTMENT OF HOMELAND SECURITY

Coast Guard

[Docket No. USCG–2021–0407]

Guidance: Change 2 to NVIC 02–18 Guidelines on Qualification for STCW Endorsements as Officer in Charge of a Navigational Watch of Vessels of Less Than 500 GT

AGENCY: Coast Guard, Homeland Security (DHS).

ACTION: Notice of availability.

SUMMARY: The Coast Guard announces the availability of change 2 to Navigation and Vessel Inspection Circular (NVIC) 02–18, Guidelines on Qualification for STCW Endorsements as Officer in Charge of a Navigational Watch of Vessels of Less Than 500 GT. This NVIC provides guidance to mariners concerning regulations governing endorsements to Merchant Mariner Credentials for service on vessels of less than 500 Gross Tons (GT) (i.e., not limited to near-coastal waters). This change notice revises NVIC 02–18 to indicate that the Coast Guard will not enforce the 3 month maximum allowable substitution for credible sea service in a rating capacity.

DATES: The policies announced in Change–2 to NVIC 02–18 are effective as of August 26, 2021.


FOR FURTHER INFORMATION CONTACT: For information about this document, contact the James Cavo, Mariner Credentialing Program Policy Division (CG–MMC–2), Coast Guard; telephone 202–372–1205; email MMCPolicy@uscg.mil.

SUPPLEMENTARY INFORMATION: NVIC 02–18 describes the Coast Guard’s policy for merchant mariners to qualify for and renew Standards of Training, Certification and Watchkeeping for Seafarers (STCW) endorsements to Merchant Mariner Credentials under 46 CFR 11.319 for service as officer in charge of a navigational watch (OICNW) on vessels of less than 500 GT upon all waters (i.e., not limited to near-coastal waters). The Coast Guard has become aware that mariners seeking an endorsement as OICNW of vessels of less than 500 GT are experiencing difficulties meeting the bridge watchkeeping requirements for the endorsement in a rating capacity.

As specified in § 11.319(a)(2), mariners must obtain at least six months of service performing bridge watchkeeping duties. This section also limits the amount of service in a rating capacity that can be used to meet this requirement to not more than 6 months of experience, which is accepted as a maximum of three months of creditable service. The mariner would have to obtain three additional months of service performing bridge watchkeeping duties in a non-rating capacity, such as an officer capacity. The OICNW of vessels less than 500 GT is considered a first certificate of competence, in that it may be the first STCW officer endorsement a mariner obtains, mariners typically qualify by serving as a rating such as able seaman. It is often impossible for these mariners to meet the 6 month bridge watchkeeping requirement with the 3 month limit on credible rating service because they cannot serve as an officer to accrue the remaining 3 months of bridge watchkeeping without this OICNW first certificate of competence.

This CH–2 will remedy that barrier to qualifying for the OICNW of vessels less than 500 GT endorsement by not enforcing the 3 month maximum allowable substitution for service as a rating to meet the required service performing bridge watchkeeping duties. This change notice revises NVIC 02–18 to indicate that the Coast Guard will accept, on a day for day basis and without limitation, service in any capacity performing bridge watchkeeping duties under the supervision of an officer holding the STCW endorsement as Master, Chief Mate, or OICNW. Allowing mariners to accrue bridge watchkeeping service in a