DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, National Institutes of Health; Notice of Meetings

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended, notice is hereby given of meetings of the Advisory Committee on Research on Women’s Health.

The meetings will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meetings.

The meetings will also be videocast and can be accessed from the NIH Videocasting and Podcasting website (http://videocast.nih.gov/).

Name of Committee: Advisory Committee on Research on Women’s Health.
Date: April 21, 2020.
Time: 9:00 a.m. to 5:00 p.m.
Agenda: Director’s Report and Scientific Presentations.
Place: National Institutes of Health, Porter Neuroscience Center, Building 35A, Conference Room 620/630, 35 Center Drive, Bethesda, MD 20892.
Contact Person: Elizabeth Spencer, R.N., Deputy Director, Office of Research on Women’s Health, Executive Secretary, ACRWH, National Institutes of Health, 6707 Democracy Blvd., Room 7W444, Bethesda, MD 20817. (301) 402–1770. elizabeth.spencer@nih.gov.

Any member of the public interested in presenting oral comments to the committee may notify the Contact Person listed on this notice at least 10 days in advance of the meetings. Interested individuals and representatives of organizations may submit a letter of intent, a brief description of the organization represented, and a short description of the oral presentation. Only one representative of an organization may be allowed to present oral comments and if accepted by the committee, presentations may be limited to five minutes. Both printed and electronic copies are requested for the record. In addition, any interested person may file written comments with the committee by forwarding their 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.

In the interest of security, NIH has instituted stringent procedures for entrance onto the NIH campus. All visitor vehicles, including taxicabs, hotel, and airport shuttles will be inspected before being allowed on campus. Visitors will be asked to show one form of identification (for example, a government-issued photo ID, driver’s license, or passport) and to state the purpose of their visit.

Information is also available on the Institute’s/Center’s home page: https://orwh.od.nih.gov/, where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.14, Intramural Research Training Award; 93.22, Clinical Research Loan Repayment Program for Individuals from Disadvantaged Backgrounds; 93.232, Loan Repayment Program for Research Generally; 93.39, Academic Research Enhancement Award; 93.936, NIH Acquired Immunodeficiency Syndrome Research Loan Repayment Program; 93.187, Undergraduate Scholarship Program for Individuals from Disadvantaged Backgrounds, National Institutes of Health, HHS)

Miguelina Perez, Program Analyst, Office of Federal Advisory Committee Policy.

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Center for Substance Abuse Prevention; Notice of Meeting

Pursuant to Public Law 92–463, notice is hereby given for the meeting of the Substance Abuse and Mental Health Services Administration’s (SAMHSA) Center for Substance Abuse Prevention National Advisory Council (CSAP NAC) on March 17, 2020.

The Council was established to advise the Secretary, Department of Health and Human Services (HHS); the Assistant Secretary for Mental Health and Substance Use, SAMHSA; and Director, CSAP concerning matters relating to the activities carried out by and through the Center and the policies respecting such activities.

The meeting will be open to the public and will include the discussion of the Evidence-Based Practices Resource Center; new SAMHSA publications; adolescent prevention programs/activities; and Fostering Healthy Mental, Emotional, and Behavioral Development. The meeting will also include updates on CSAP program developments. The meeting will be held in Rockville, Maryland. Attendance by the public on-site will be limited to the space available. Interested persons may present data, information, or views, orally or in writing, on issues pending before the Council. Written submissions should be forwarded to the contact person on or before one week prior to the meeting. Oral presentations from the public will be scheduled at the conclusion of the meeting. Individuals interested in making oral presentations should notify the contact on or before one week prior to the meeting. Five minutes maximum will be allotted for each presentation.

To attend onsite, submit written or brief oral comments, or request special accommodations for persons with disabilities, please register at the SAMHSA Committees’ website, https://snaregister.samhsa.gov/MeetingList.aspx, or communicate with the CSAP Council’s Designated Federal Officer (see contact information below).

Substantive program information may be obtained after the meeting by contacting the SAMHSA Committee website, https://www.samhsa.gov/about-us/advisory-councils, or by contacting the Designated Federal Officer.

Committee Name: Substance Abuse and Mental Health Services Administration, Center for Substance Abuse Prevention National Advisory Council.

Date/Time/Type: March 17, 2020, from 9:30 a.m. to 5:00 p.m. EDT (OPEN).
Place: SAMHSA, 5600 Fishers Lane, Room SN54, Rockville, MD 20852, Adobe Connect webcast: https://samhsa-csap.adobeconnect.com/nac/.
Contact: Matthew J. Aumen, Designated Federal Officer, SAMHSA CSAP NAC, 5600 Fishers Lane, Rockville, MD 20852, Telephone: 240–276–2440, Fax: 301–460–8480, Email: matthew.aumen@samhsa.hhs.gov.
Carlos Castillo,
Committee Management Officer, SAMHSA.

BILLING CODE 4162–20–P

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).

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 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 51110); 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 Oral Fluid 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:

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

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, 504–361–8989/800–433–4276 (Formerly: Kroll Laboratory Specialists, Inc., Laboratory Specialists, Inc.).

Written comments and recommendations concerning the proposed information collection should be sent by April 1, 2020 to the SAMHSA Desk Officer at the Office of Management and Budget (OMB). To ensure timely receipt of comments, and to avoid potential delays in OMB’s receipt and processing of mail sent through the U.S. Postal Service, commenters are encouraged to submit their comments to OMB via email to: OIRA_Submission@omb.eop.gov.

Although commenters are encouraged to submit their comments in writing, electronic submission of comments is preferred. Comments may be submitted to OIRA during business hours (Monday through Friday, 8:45 am to 5:00 pm Eastern Time) via email to: OIRA_Submission@omb.eop.gov. They may also be faxed to (202) 395-7380 or mailed to: OIRA Submission Desk Officer, Office of Information and Regulatory Affairs, U.S. Office of Management and Budget, Room 2101, 700 7th Street NW., Washington, DC 20503.

The estimated response burden is as follows:

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<th>Form name</th>
<th>Number of respondents</th>
<th>Responses per respondent</th>
<th>Total responses</th>
<th>Hours per response</th>
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<td>SOAR Web-based Data Form (Part I)</td>
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<tr>
<td>Annual Report Questions (Part II)</td>
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There are two parts to the SOAR Web-based Data Form. Part I of the SOAR Web-based Data Form is intended for SOAR-trained case workers to enter the outcomes of SOAR-assisted SSI/SSDI applications. Part II of the SOAR Web-based Data Form includes two sections reserved for SOAR State Team Leads to report annually. The first section of Part II collects quantitative summary data from states that do not track SOAR-assisted SSI/SSDI applications using the SOAR Web-based Data Form Part I. The second section of Part II collects qualitative (open-ended) questions on annual SOAR accomplishments, identified challenges, and collaborations.

For more information, including a description of the data collected, visit the SOAR Technical Assistance Center site: https://soartrack.prainc.com.