

www.fda.gov/NewsEvents/MeetingsConferencesWorkshops/ucm592778.htm. Please provide complete contact information for each attendee, including name, title, affiliation, address, email, and telephone.

Registration is free and based on space availability, with priority given to early registrants. Persons interested in attending this public workshop must register by April 25, 2018, 5 p.m. Eastern Time. Early registration is recommended because seating is limited; therefore, FDA may limit the number of participants from each organization. Registrants will receive confirmation when their registration has been received. If time and space permit, onsite registration on the day of the public workshop will be provided beginning an hour prior to the start of the meeting.

If you need special accommodations due to a disability, please contact Nicole Wolanski, at 301-796-6570, or OOPDOrphanEvents@fda.hhs.gov no later than April 25, 2018.

An agenda for the workshop and any other background materials will be made available 5 days before the workshop at <https://www.fda.gov/NewsEvents/MeetingsConferencesWorkshops/ucm592778.htm>.

Streaming Webcast of the Public Workshop: For those unable to attend in person, FDA will provide a live webcast of the workshop. To register for the streaming webcast of the public workshop, please visit the following website by May 8, 2018: <https://www.fda.gov/NewsEvents/MeetingsConferencesWorkshops/ucm592778.htm>.

If you have never attended a Connect Pro event before, test your connection at https://collaboration.fda.gov/common/help/en/support/meeting_test.htm. To get a quick overview of the Connect Pro program, visit https://www.adobe.com/go/connectpro_overview. FDA has verified the website addresses in this document, as of the date this document publishes in the **Federal Register**, but websites are subject to change over time.

Transcripts: Please be advised that as soon as a transcript of the public workshop is available, it will be accessible at <https://www.regulations.gov>. It may be viewed at the Dockets Management Staff (see **ADDRESSES**). A link to the transcript will also be available on the internet at <https://www.fda.gov/NewsEvents/MeetingsConferencesWorkshops/ucm592778.htm>.

Dated: February 22, 2018.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2018-03961 Filed 2-26-18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Service Administration

Women's Preventive Services Guidelines

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: Applicable as of December 29, 2017, HRSA updated the HRSA-supported Women's Preventive Services Guidelines for purposes of health insurance coverage for preventive services that address health needs specific to women based on clinical recommendations from the Women's Preventive Services Initiative. This 2017 update adds two additional services—Screening for Diabetes Mellitus after Pregnancy and Screening for Urinary Incontinence—to the nine preventive services included in the 2016 update to the HRSA-supported Women's Preventive Services Guidelines. The nine services included in the 2016 update are as follows: Breast Cancer Screening for Average Risk Women, Breastfeeding Services and Supplies, Screening for Cervical Cancer, Contraception, Screening for Gestational Diabetes Mellitus, Screening for Human Immunodeficiency Virus Infection, Screening for Interpersonal and Domestic Violence, Counseling for Sexually Transmitted Infections, and Well-Woman Preventive Visits. This notice serves as an announcement of the decision to update the guidelines as listed below. Please see <https://www.hrsa.gov/womens-guidelines/index.html> for additional information.

FOR FURTHER INFORMATION CONTACT: Kimberly C. Sherman, Maternal and Child Health Bureau, HRSA at phone: (301) 443-0543; email: wellwomancare@hrsa.gov.

SUPPLEMENTARY INFORMATION: The complete set of updated 2017 HRSA-supported Women's Preventive Services Guidelines includes those that were accepted by the Acting HRSA Administrator on December 20, 2016, as well as two new services, Screening for Diabetes Mellitus After Pregnancy and Screening for Urinary Incontinence. For a complete listing and detailed

information about the December 20, 2016, updates, please see <https://www.federalregister.gov/documents/2016/12/27/2016-31129/updating-the-hrsa-supported-womens-preventive-services-guidelines>. In addition, the December 20, 2016, updates, including information related to coverage of contraceptive services and exemption for objecting organizations from requirements related to the provision of contraceptive services, can be found at <https://www.hrsa.gov/womens-guidelines-2016/index.html>. Information regarding the two new services that were accepted by the HRSA Administrator on December 29, 2017, is set out below:

1. Screening for Diabetes Mellitus After Pregnancy

The Women's Preventive Services Initiative recommends women with a history of gestational diabetes mellitus (GDM) who are not currently pregnant and who have not previously been diagnosed with type 2 diabetes mellitus should be screened for diabetes mellitus. Initial testing should ideally occur within the first year postpartum and can be conducted as early as 4–6 weeks postpartum.

Women with a negative initial postpartum screening test result should be rescreened at least every 3 years for a minimum of 10 years after pregnancy. For women with a positive postpartum screening test result, testing to confirm the diagnosis of diabetes is indicated regardless of the initial test (*e.g.*, oral glucose tolerance test, fasting plasma glucose, or hemoglobin A1c). Repeat testing is indicated in women who were screened with hemoglobin A1c in the first six months postpartum regardless of the result (see Implementation Considerations below).

2. Screening for Urinary Incontinence

The Women's Preventive Services Initiative recommends screening women for urinary incontinence annually. Screening should ideally assess whether women experience urinary incontinence and whether it impacts their activities and quality of life. The Women's Preventive Services Initiative recommends referring women for further evaluation and treatment if indicated.

HRSA-Supported Women's Preventive Services Guidelines

The HRSA-supported Women's Preventive Services Guidelines were originally established in 2011 based on recommendations from an HHS commissioned study by the Institute of Medicine, now known as the National

Academy of Medicine (NAM). Since then, there have been advancements in science and gaps identified in the existing guidelines, including a greater emphasis on practice-based clinical considerations. To address these, HRSA awarded a 5-year cooperative agreement in March 2016 to convene a coalition of clinician, academic and consumer-focused health professional organizations and conduct a scientifically rigorous review to develop recommendations for updated Women's Preventive Services Guidelines in accordance with the model created by the NAM *Clinical Practice Guidelines We Can Trust*. The American College of Obstetricians and Gynecologists was awarded the cooperative agreement and formed an expert panel called the Women's Preventive Services Initiative.

Under section 2713 of the Public Health Service Act, non-grandfathered group health plans and issuers of non-grandfathered group and individual health insurance coverage are required to cover specified preventive services without a copayment, coinsurance, deductible, or other cost sharing, including preventive care and screenings for women as provided for in comprehensive guidelines supported by HRSA for this purpose. Non-grandfathered group health plans and health insurance issuers offering non-grandfathered group or individual coverage (generally, plans or policies created or sold after March 23, 2010, or older plans or policies that have been changed in certain ways since that date) are required to provide coverage without cost sharing for preventive services listed in the updated HRSA-supported guidelines (which include the nine preventive services set out in the 2016 update, as well as the two services added in this update) beginning with the first plan year (in the individual market, policy year) that begins on or after December 29, 2018.

Dated: February 20, 2018.

George Sigounas,
Administrator.

[FR Doc. 2018-03840 Filed 2-26-18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR-16-366 Dual Purpose with Dual Benefit: Research in Biomedicine and Agriculture.

Date: March 21-22, 2018.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Tera Bounds, DVM, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3214, MSC 7808, Bethesda, MD 20892, 301-435-2306, boundst@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict.

Date: March 21, 2018.

Time: 11:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Kate Fothergill, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3142, Bethesda, MD 20892, 301-435-2309, fothergillke@mail.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: The Biostatistical Methods and Research Design.

Date: March 21, 2018.

Time: 1:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Ping Wu, Ph.D., Scientific Review Officer, HDM IRG, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3166, Bethesda, MD 20892, 301-451-8428, wup4@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR Panel: AIDS and Related Research.

Date: March 22, 2018.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: The Fairmont Washington, DC, 2401 M Street NW, Washington, DC 20037.

Contact Person: Robert Freund, Ph.D., Scientific Review Officer, Center for

Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5216, MSC 7852, Bethesda, MD 20892, 301-435-1050, freundr@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowship: Infectious Diseases and Microbiology.

Date: March 22-23, 2018.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Embassy Suites DC Convention Center, 900 10th Street NW, Washington, DC 20001.

Contact Person: Tamara Lyn McNealy, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3188, Bethesda, MD 20747, 301-827-2372, tamara.mcnealy@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR Panel: Cancer Health Disparities.

Date: March 22, 2018.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Janet M. Larkin, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5142, MSC 7840, Bethesda, MD 20892, 301-806-2765, larkinja@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Innovative Immunology.

Date: March 22, 2018.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: The William F. Bolger Center, 9600 Newbridge Drive, Potomac, MD 20854.

Contact Person: Andrea Keane-Myers, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4218, Bethesda, MD 20892, 301-435-1221, andrea.keane-myers@nih.gov.

Name of Committee: AIDS and Related Research Integrated Review Group; AIDS Discovery and Development of Therapeutics Study Section.

Date: March 22, 2018.

Time: 8:00 a.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.

Place: The Fairmont Washington, DC, 2401 M Street NW, Washington, DC 20037.

Contact Person: Shiv A. Prasad, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5220, MSC 7852, Bethesda, MD 20892, 301-443-5779, prasads@csr.nih.gov.

Name of Committee: AIDS and Related Research Integrated Review Group; NeuroAIDS and other End-Organ Diseases Study Section.

Date: March 22, 2018.

Time: 8:00 a.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.