

Jeffrey M. Zirger,

*Lead, Information Collection Review Office,
Office of Scientific Integrity, Office of Science,
Centers for Disease Control and Prevention.*

[FR Doc. 2022-04191 Filed 2-28-22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

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

The meeting 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, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92-463. 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: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—SIP22-005, Building Resilience Against Climate Effects (BRACE): Enhancing Practical Guidance to Support Climate and Health Adaptation Planning.

Date: May 4, 2022.

Time: 11:00 a.m.–6:00 p.m., EDT.

Place: Teleconference.

Agenda: To review and evaluate grant applications.

FOR FURTHER INFORMATION CONTACT: Jaya Raman, Ph.D., Scientific Review Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 4770 Buford Highway, Mailstop S107-B, Atlanta, Georgia 30341, Telephone: (770) 488-6511, email: JRaman@cdc.gov.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and

Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

*Director, Strategic Business Initiatives Unit,
Office of the Chief Operating Officer, Centers
for Disease Control and Prevention.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

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

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Name of Committee: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)-RFA-CK-22-001, Investigation of Monkeypox and Other Zoonotic Diseases in the Democratic Republic of the Congo (DRC); RFA-CK-22-002, Technological Advancement of Global Rabies Surveillance and Control; and RFA-CK-22-004, Optimization and Standardization of Methods to Suppress Ixodes scapularis and Disrupt Zoonotic Pathogen Transmission in Settings Posing an Elevated Risk to Humans.

Date: April 28, 2022.

Time: 10:00 a.m.–5:00 p.m. (EDT).

Place: Teleconference, Centers for Disease Control and Prevention, Room 1080, 8 Corporate Square Blvd., Atlanta, GA 30329.

Agenda: To review and evaluate grant applications.

For Further Information Contact: Gregory Anderson, M.S., M.P.H., Scientific Review Officer, National Center for HIV, Viral Hepatitis, STD, and TB Prevention, CDC, 1600 Clifton Road NE, Mailstop US8-1, Atlanta,

Georgia 30329, (404) 718-8833, ganderson@cdc.gov.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

*Director, Strategic Business Initiatives Unit,
Office of the Chief Operating Officer, Centers
for Disease Control and Prevention.*

[FR Doc. 2022-04261 Filed 2-28-22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

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The meeting 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, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92-463. 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: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—PAR 20-297, NIOSH Centers of Excellence for Total Worker Health (TWH).

Date: April 21, 2022.

Time: 1:00 p.m.–4:00 p.m., EDT.

Place: Video-Assisted Meeting.

Agenda: To review and evaluate grant applications.

For Further Information Contact: Dan Hartley, Ed.D., Scientific Review Officer, Office of Extramural Programs, National Institute for Occupational Safety and Health, CDC, 1095 Willowdale Road, Morgantown, West

Virginia 26505, Telephone: (304) 285–5812; Email: DHartley@cdc.gov.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–22–22CX; Docket No. CDC–2022–0031]

Proposed Data Collections Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), in the Department of Health and Human Services (HHS), as part of its continuing effort to reduce public burden, invites the general public and other federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Preferences for Longer-Acting Preexposure Prophylaxis (LA-PrEP) Methods Among Persons in U.S. Populations at Highest Need: A Discrete Choice Experiment. The proposed project is designed to understand preferences for LA-PrEP products for HIV prevention among potential users and providers.

DATES: CDC must receive written comments on or before May 2, 2022.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2022–0031, by either of the following methods:

- *Federal eRulemaking Portal:* [Regulations.gov](https://www.regulations.gov). Follow the instructions for submitting comments.

- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to [regulations.gov](https://www.regulations.gov).

Please note: Submit all comments through the Federal eRulemaking portal ([regulations.gov](https://www.regulations.gov)) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329; phone: 404–639–7118; Email: omb@cdc.gov.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
2. Evaluate 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. Enhance the quality, utility, and clarity of the information to be collected;
4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submissions of responses; and

5. Assess information collection costs.

Proposed Project

Preferences for Longer-Acting Preexposure Prophylaxis (LA-PrEP) Methods Among Persons in U.S. Populations at Highest Need: A Discrete Choice Experiment—New—National Center for HIV, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The 2022–2025 National HIV/AIDS Strategy includes a goal of increasing pre-exposure prophylaxis (PrEP) coverage to 50 percent among persons with indications, from a 2017 baseline of 13.2 percent. Despite successes in development and scale up of daily oral PrEP as a biomedical HIV prevention product, studies consistently show obstacles to its uptake and continuation. The Centers for Disease Control and Prevention (CDC) and its partners must engage in early planning for the implementation of longer-acting PrEP (LA-PrEP) agents to help achieve the U.S. Ending the HIV Epidemic (EHE) goal of reducing incident HIV infections by 90 percent by 2030. Understanding providers' and priority populations' preferences for different LA-PrEP agents, and perceived advantages and disadvantages of each product, will be critical to estimating future uptake and use of the various products that are recently or soon likely to become available for prescription.

The goal of this study is to understand preferences for LA-PrEP products for HIV prevention among potential users and providers, including product characteristics and other service delivery factors that may facilitate or hinder future uptake of these products. In cooperation with partners, CDC will conduct a discrete choice experiment (DCE) among providers and potential users of LA-PrEP products to elicit their preferences for characteristics of LA-PrEP and delivery programs to maximize uptake of LA-PrEP among people in need of HIV prevention methods. Results from this experiment will be used to identify factors key to adoption and implementation of each product and increase implementation efficiency by identifying strategies to support decision making and address potential challenges.

The study design is a cross-sectional, online survey comprised of a DCE and additional questions to directly elicit participant preferences and gather data on socioeconomic, behavioral, and attitudinal factors. DCE methods are based on the principle that products or