CVM, or the date U.S. Bank notifies FDA that your payment in the full amount has been received, or when the U.S. Department of the Treasury notifies FDA of receipt of an electronic or wire transfer payment. U.S. Bank and the U.S. Department of the Treasury are required to notify FDA within 1 working day, using the PIN described previously.

The tax identification number of FDA is 53–0196965.

B. Application Cover Sheet Procedures

Step One: Create a user account and password. Log on to the ADUFA website at *https://www.fda.gov/industry/animal-drug-user-fee-act-adufa/animal-drug-user-fee-cover-sheet* and, under Application Submission Information, click on ''Create ADUFA User Fee Cover Sheet.'' For security reasons, each firm submitting an application will be assigned an organization identification number, and each user will also be required to set up a user account and password the first time you use this site. Online instructions will walk you through this process.

Step Two: Create an Animal Drug User Fee Cover Sheet, transmit it to FDA, and print a copy. After logging into your account with your username and password, complete the steps required to create an Animal Drug User Fee Cover Sheet. One cover sheet is needed for each animal drug application or supplement. Once you are satisfied that the data on the cover sheet are accurate and you have finalized the cover sheet, you will be able to transmit it electronically to FDA and you will be able to print a copy of your cover sheet showing your unique PIN.

Step Three: Send the payment for your application as described in section IX.A.

Step Four: Submit your application.

C. Product, Establishment, and Sponsor Fees

By December 31, 2022, FDA will issue invoices and payment instructions for product, establishment, and sponsor fees for FY 2023 using this fee schedule. Payment will be due by January 31, 2023. FDA will issue invoices in November 2023 for any products, establishments, and sponsors subject to fees for FY 2023 that qualify for fees after the December 2022 billing.

Dated: July 22, 2022.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2022–16176 Filed 7–27–22; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Request for Information (RFI): Inviting Comments and Suggestions on an ODS Strategic Plan 2022–2026

AGENCY: National Institutes of Health, HHS.

ACTION: Request for information.

SUMMARY: Since its inception in 1994, the National Institutes of Health (NIH), Office of Dietary Supplements (ODS) has used a structured planning process to develop five-year strategic plans. ODS is committed to engaging its stakeholders including representatives of the scientific community, industry, other federal agencies, and the public in the strategic planning process by soliciting their comments on the draft ODS Strategic Plan for Fiscal Years (FY) 2022–2026. Considering comments from representative stakeholder groups, and the general public will help ODS assess the outcomes of its investments and prioritize plans for the next five years.

DATES: The RFI is open for public comments and must be received by 11:59:59 p.m. (ET) on August 31, 2022, to ensure consideration.

ADDRESSES: All comments must be submitted electronically to *odsplan@od.nih.gov.*

FOR FURTHER INFORMATION CONTACT: Please direct all inquiries to: Barbara Cohen at *ODSplan@od.nih.gov* or (301) 435–2920.

SUPPLEMENTARY INFORMATION: This notice is in accordance with the 21st Century Cures Act, wherein NIH institutes are required to regularly update their strategic plans. The purpose of the FY 2022–2026 ODS Strategic Plan (*https://ods.od.nih.gov/About/StrategicPlan.aspx*) is to communicate how ODS will advance its mission to support, coordinate, and disseminate the results of scientific research, and provide leadership to help expand the knowledge, scientific evidence, and understanding of dietary supplements, thus fostering an enhanced quality of life and health for the U.S. population. The plan articulates ODS' priorities in five key areas (goals):

(1) Expand the scientific knowledge base on dietary supplements by stimulating and supporting a full range of biomedical research and by developing and contributing to relevant initiatives, workshops, meetings, and conferences;

(2) Enhance the dietary supplement research workforce through training and career development and simultaneously support the development of programs for diverse researchers who are underrepresented in science;

(3) Foster development and dissemination of research resources and tools to enhance the quality of dietary supplement research;

(4) Translate dietary supplement research findings into useful information for consumers, health professionals, researchers, and policymakers; and

(5) Coordinate and support the development of collaborative initiatives to address gaps in dietary supplement research.

ODS has completed a draft of its Five-Year Strategic Plan for FY 2022–2026 (*https://ods.od.nih.gov/About/StrategicPlan.aspx*) and is interested in receiving feedback from all interested parties on the following:

- Are there additional emerging public health issues that ODS can help address?
- Are there existing knowledge gaps that ODS can help address (not included in the current plan)?
- What can ODS do better to meet the needs of its stakeholders?

ODS encourages organizations to submit a single response reflective of the views of the organization as a whole.

Responses to this RFI are voluntary and may be submitted anonymously. Please do not include any personally identifiable information or any information that you do not wish to make public. Proprietary, classified, confidential, or sensitive information should not be included in your response. The NIH will use the information submitted in response to this RFI at its discretion. The NIH reserves the right to use any submitted information on public websites, in reports, in summaries of the state of the science, in any possible resultant solicitation(s), grant(s), or cooperative agreement(s), or in the development of future funding opportunity announcements. This RFI is for informational and planning purposes only and is not a solicitation for applications or an obligation on the part of the Government to provide support for any ideas identified in response to it. Please note that the Government will not pay for the preparation of any information submitted or for use of that information.

We look forward to your input and hope that you will share this RFI opportunity with your colleagues.

Dated: July 22, 2022.

Tara A. Schwetz,

Acting Principal Deputy Director, National Institutes of Health.

[FR Doc. 2022–16152 Filed 7–27–22; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Environmental Health Sciences; 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. 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: National Institute of Environmental Health Sciences Special Emphasis Panel: Mechanism for Time-Sensitive Research Opportunities in Environmental Health Sciences (R21).

Date: August 9, 2022.

Time: 12:00 p.m. to 1:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute of Environmental Health Sciences, Keystone Building, 530 Davis Drive, Durham, NC 27709 (Virtual Meeting).

Contact Person: Qingdi Quentin Li, Ph.D., MD, Scientific Review Officer, Scientific Review Branch, Division of Extramural Research and Training, Nat'l Institute of Environmental Health Sciences, Research Triangle Park, NC 27709, (240) 858–3914, *liquenti@mail.nih.gov.*

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(Catalogue of Federal Domestic Assistance Program Nos. 93.115, Biometry and Risk Estimation—Health Risks from Environmental Exposures; 93.142, NIEHS Hazardous Waste Worker Health and Safety Training; 93.143, NIEHS Superfund Hazardous Substances—Basic Research and Education; 93.894, Resources and Manpower Development in the Environmental Health Sciences; 93.113, Biological Response to Environmental Health Hazards; 93.114, Applied Toxicological Research and Testing, National Institutes of Health, HHS)

Dated: July 21, 2022.

David W. Freeman,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2022–16157 Filed 7–27–22; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; Conference, Meeting, Workshop, Registration and Challenges Generic Clearance (Office of the Director)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Ms. Mikia P. Currie, Chief, Project Clearance Branch (PCB), Office of Policy for Extramural Research Administration, 6705 Rockledge Drive, Suite 803–B, Bethesda, Maryland, 20892 or call non-toll-free number (301) 435–0941 or email your request, including your address to: *curriem@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: This proposed information collection was previously published in the **Federal Register** on May 5, 2022, pages 26768 & 26769 (87 FR 26768) and allowed 60 days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment.

The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below.

Proposed Collection: Conference, Meeting, Workshop, Registration and Challenges Generic Clearance, –0925–0740—REVISION, expiration date 07/31/2022, National Institutes of Health (NIH).

Need and Use of Information Collection: This is a revision of a currently approved generic clearance to include Challenges and Competitions as a means of promoting innovative solutions. As a result of including Challenges and Competitions, the title of this generic has been revised. This generic will continue to provide a quick and efficient process to create registration forms for NIH sponsored conference, meetings, workshops, poster sessions, presentations, and panels. NIH directly sponsors, organizes, and conducts research-related activities such as conferences, workshops, meetings, poster sessions, and training courses. These activities are designed to be relevant to the current state of research in a given field or to the current state of participant's research projects or careers, and other resource limitations and determine the number of possible participants. For such activities to be timely and to optimally use available resources to address needs and opportunities within the research community, it is necessary for NIH to have a means to register and select the most appropriate participants, according to the type or purpose of a given activity.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 10,375.