information to national organizations included in The Union on all questions concerning scientific and sociological study on TB. The Union is uniquely qualified to combat TB and lung disease globally and to offer training and other capacity-building activities leading to health solutions for the poor in resource limited countries, the key activities under this NOFO, due to its WHO-recognized accomplishments and leadership role in the global TB fight since its founding in 1920.

Summary of the Award

Recipient: International Union Against Tuberculosis and Lung Disease (The Union).

Purpose of the Award: The purpose of this award is to continue developing and updating TB scientific and programmatic resources, disseminating TB best practices, and building TB capacity.

Amount of Award: For The Union, the approximate year 1 funding amount will be $500,000 in Federal Fiscal Year (FYY) 2024 funds, subject to the availability of funds. Funding amounts for years 2–5 will be set at continuation.

Non-PEPFAR Authority: This program is authorized under section 307 of the Public Health Service Act (42 U.S.C. 242l), as amended and section 301(a) of the Public Health Service Act (42 U.S.C. 241(a)), as amended.

Period of Performance: The period for this award will be September 30, 2024, through September 29, 2029.

Dated: March 13, 2024.

Jamie Legier,

Acting Director, Office of Grants Services, Centers for Disease Control and Prevention.

[FR Doc. 2024–05771 Filed 3–18–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2024–N–1181]

Air Products and Chemicals, Inc.; Withdrawal of Approval of a New Drug Application and New Animal Drug Application for Helium, USP

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is withdrawing approval of the new drug application (NDA) 205864 and the new animal drug application (NADA) 141–395 for the designated medical gas Helium, USP held by Air Products and Chemicals, Inc., 1940 Air Products Blvd., Allentown, PA 18106–5500 (Air Products). Air Products notified the Agency in writing that the drug product was no longer marketed and requested that the approval of the application be withdrawn.

DATES: Approval is withdrawn as of April 18, 2024.

FOR FURTHER INFORMATION CONTACT: Kimberly Lehrfeld, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6226, Silver Spring, MD 20993–0002, 301–796–3137, *Kimberly.Lehrfeld@fda.hhs.gov;* or Scott Fontana (HFV–180), Center for Veterinary Medicine, Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–402–0656, *Scott.Fontana@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Air Products has informed FDA that it is no longer marketing the designated medical gas Helium, USP and has requested that FDA withdraw approval of NDA 205864 and NADA 141–395 under the processes in § 314.150© (21 CFR 314.150(c)) and § 514.115(d) (21 CFR 514.115(d)). Air Products has also, by its request, waived its opportunity for a hearing. Withdrawal of approval of an application or abbreviated application under § 314.150(c) or an NADA or abbreviated new animal drug application under § 514.115(d) is without prejudice to refiling.

Therefore, approval of NDA 205864 and NADA 141–395, and all amendments and supplements thereto, are hereby withdrawn as of April 18, 2024. Introduction or delivery for introduction into interstate commerce of Helium, USP, without an approved new drug application or an approved new animal drug application violates sections 505(a), 512(a), 301(a), and 301(d) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(a), 360b(a)(1), 331(a), and 331(d)). Any Helium, USP manufactured by Air Products pursuant to these applications that is in inventory on April 18, 2024 may continue to be dispensed until the inventories have been depleted or the drug product has reached its expiration date or otherwise become violative, whichever occurs first.

Dated: March 13, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024–05742 Filed 3–18–24; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Committee on Vital and Health Statistics

AGENCY: Centers for Disease Control and Prevention, Department of Health and Human Services.

ACTION: Notice of meeting.

SUMMARY: Pursuant to the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) announces the following advisory committee meeting. This meeting is open to the public. The public is welcome to attend in person or to obtain the link to attend this meeting by following the instructions posted on the Committee website: *https://ncvhs.hhs.gov/meetings/full-committee-meeting-16/.*

Name: National Committee on Vital and Health Statistics (NCVHS) Meeting.

DATES: Thursday, April 11, 2024: 9:15 a.m.–5:30 p.m. EDT and Friday, April 12, 2024: 8:30 a.m.–3:00 p.m. EDT.

ADDRESSES: In-person/hybrid (includes virtual attendance option).

FOR MORE INFORMATION CONTACT: Substantive program information may be obtained from Rebecca Hines, MHS, Executive Secretary, NCVHS, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Hyattsville, Maryland 20782, or via electronic mail to *vgh4@cdc.gov;* or by telephone (301) 458–4715. Summaries of meetings and a roster of Committee members are available on the NCVHS website *https://ncvhs.hhs.gov/,* where further information including an agenda and instructions to access the broadcast of the meeting will be posted.

Should you require reasonable accommodation, please telephone the CDC Office of Equal Employment Opportunity at (770) 488–3210 as soon as possible.

SUPPLEMENTARY INFORMATION:

Purpose: As outlined in its Charter, the National Committee on Vital and Health Statistics assists and advises the Secretary of HHS on health data, data standards, statistics, privacy, national health information policy, and the Department's strategy to best address those issues. Under the Health Insurance Portability and Accountability Act of 1996 (HIPAA),[1] NCVHS advises the Secretary on administrative simplification standards, including those for privacy, security,

[1] Public Law 104–191, 110 Stat. 1936 (Aug 21, 1996), available at *https://www.congress.gov/104/plaws/publ191/PLAW-104publ191.pdf.*

adoption and implementation of transaction standards, unique identifiers, code sets, and operating rules adopted under the Patient Protection and Affordable Care Act (ACA).[2]

The meeting agenda will include discussion of the 2024 workplan including the NCVHS Report to Congress, and briefings and discussions with invited experts on several health data policy topics, including: standards for SDOH data elements; possible implications of Value Based Care Models vs Fee-For-Service on HIPAA standards; an overview of the key elements of the Trusted Exchange Framework and Common Agreement (TEFCA) published by the Office of the National Coordinator for Health IT (ONC) in November 2023; and exploration of privacy and security in AI in technology and healthcare.

The NCVHS Workgroup on Timely and Strategic Action to Inform ICD–11 Policy for morbidity will report to the full Committee on Phase II of its work focusing on analysis of the recent Request for Information (RFI), published in October 2023.[3]

The Committee will reserve time on the agenda for public comment. Meeting times and topics are subject to change. Please refer to the agenda posted on the NCVHS website for updates: *https://ncvhs.hhs.gov/meetings/full-committee-meeting-16/.*

Sharon Arnold,

Associate Deputy Assistant Secretary, Office of Science and Data Policy, Office of the Assistant Secretary for Planning and Evaluation.

[FR Doc. 2024–05779 Filed 3–18–24; 8:45 am]

BILLING CODE 4150–05–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Neurological Disorders and Stroke; Notice of Closed Meeting

Pursuant to section 1009 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 Neurological Disorders and Stroke Special Emphasis Panel; P01 and R03 Review.

Date: April 3, 2024.

Time: 10:30 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852.

Contact Person: M. Catherine Bennett, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, NINDS/NIH/HHS, NSC, 6001 Executive Blvd., Rockville, MD 20852, 301–435–1766, *bennettc3@csr.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.853, Clinical Research Related to Neurological Disorders; 93.854, Biological Basis Research in the Neurosciences, National Institutes of Health, HHS).

Dated: March 13, 2024.

Lauren A. Fleck,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024–05727 Filed 3–18–24; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer at (240) 276–1243.

Comments are invited on: (a) whether the proposed collections of information are necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Project: Minority AIDS Initiative—Management Reporting Tools (MAI–MRTs)—(OMB No. 0930–0357)—Renewal

The Substance Abuse and Mental Health Services Administration (SAMHSA), Center for Substance Abuse Prevention (CSAP) is requesting from the Office of Management and Budget (OMB) approval for the renewal of Minority AIDS Initiative (MAI) monitoring tools, which includes both youth and adult questionnaires, as well as the quarterly progress report. This revision includes the inclusion of new cohorts, substantial revisions to the youth and adult questionnaires, updates to the data used to estimate response rates and expected numbers of participants by service duration (see Table 1 below).

The cohorts of grantees funded by the MAI and included in this clearance request are:

- Prevention Navigators 2019
- Prevention Navigators 2020
- Prevention Navigators 2021
- Prevention Navigators 2022
- Prevention Navigators 2023

The target population for the grantees will be at-risk minority adolescents and young adults. All MAI grantees are expected to report their monitoring data using SAMHSA's Strategic Prevention Framework (SPF) and to target minority populations, as well as other high-risk groups residing in communities of color with high prevalence of Substance Abuse and HIV/AIDS. The primary objectives of the monitoring tools include:

- Assess the success of the MAI in reducing risk factors and increasing protective factors associated with the transmission of the Human Immunodeficiency Virus (HIV), Hepatitis C Virus (HCV) and other sexually transmitted diseases (STD).
- Measure the effectiveness of evidence-based programs and infrastructure development activities such as: outreach and training, mobilization of key stakeholders, substance abuse and HIV/AIDS counseling and education, testing, referrals to appropriate medical treatment and/or other intervention strategies (*i.e.,* cultural enrichment

[2] Public Law 111–148, 124 Stat.119, available at *https://www.congress.gov/111/plaws/publ148/PLAW-111publ148.pdf.*

[3] **Federal Register** Notice, October 16, 2023: *https://www.federalregister.gov/documents/2023/10/16/2023-22753/national-committee-on-vital-and-health-statistics.*