

Dated: February 5, 2026.

Sheila R. Garrity,

Director, Office of Research Integrity, Office of the Assistant Secretary for Health.

[FR Doc. 2026-02505 Filed 2-6-26; 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 1009 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 Panel: Topics in Clinical Care, Disease Management and Health Outcomes.

Date: February 24, 2026.

Time: 10:00 a.m. to 1:00 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Erica Charlot Spears, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 480-3211, spearsec@csr.nih.gov.

This notice is being published less than 15 days from the meeting date due to exceptional circumstances. As a result of the 43-day government shutdown, due to lapsed appropriations, the above meeting was canceled. This meeting was to assess the scientific and technical merit of NIH grant applications, required by statute to disburse NIH funds. The meeting must take place urgently so that evaluations of biomedical research applications addressing multiple major public health priorities can be submitted to the national advisory councils for timely funding recommendations.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Innovations for Healthy Living and Technologies for Improving Health.

Date: February 24, 2026.

Time: 1:00 p.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Cheryl K. Nordstrom, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, 301-402-6711, cheryl.nordstrom@nih.gov.

This notice is being published less than 15 days from the meeting date due to exceptional circumstances. As a result of the 43-day government shutdown, due to lapsed appropriations, the above meeting was canceled. This meeting was to assess the scientific and technical merit of NIH grant applications, required by statute to disburse NIH funds. The meeting must take place urgently so that evaluations of biomedical research applications addressing multiple major public health priorities can be submitted to the national advisory councils for timely funding recommendations.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393-93.396, 93.837-93.844, 93.846-93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: February 4, 2026.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2026-02502 Filed 2-6-26; 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: Submission for OMB Review; Comment Request

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of information collection requests under Office of Management and Budget (OMB) review, in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these documents, email or call the SAMHSA Reports Clearance Officer at samhsapra@samhsa.hhs.gov or (240) 276-0166.

Proposed Project: Program Evaluation for Prevention Contract (PEPC) Evaluation (OMB No. 0930-0377)—Revision

The Substance Abuse and Mental Health Services Administration's (SAMHSA) Center for Behavioral Health Statistics and Quality (CBHSQ) aims to complete a cross-site evaluation of SAMHSA's Strategic Prevention Framework for Prescription Drugs (SPF

Rx). SPF Rx is designed to address nonmedical use of prescription drugs as well as opioid overdoses by raising awareness about the dangers of sharing medications and by working with pharmaceutical and medical communities on the risks of overprescribing. The SPF Rx program grantees also raise community awareness and bring activities related to prescription drug misuse prevention and education to schools, communities, parents, prescribers, and their patients. The SPF Rx program aims to promote collaboration between states, tribes, U.S. territories, and pharmaceutical and medical communities to understand the risks of overprescribing to youth ages 12-17 and adults 18 years of age and older. The program also aims to enhance capacity for, and access to, Prescription Drug Monitoring Program (PDMP) data for prevention purposes.

This request for data collection includes a revision to previously approved OMB instruments to allow for data collection through the end of the grant period with the FY 2021 and FY 2022 grantees. The FY 2021 cohort of grants focused on the dangers of sharing medications; the risks of overprescribing, especially to young adults; community awareness and education; and incorporation of PDMP data into grantees' needs assessments and strategic plans. The FY 2022 cohort of grants focused on raising awareness about the risks of sharing medications, taking fake or counterfeit pills, and overprescribing.

The SPF Rx program's indicators of success are reductions in opioid overdoses, reductions in prescription drug misuse, and improved use of PDMP data. Data collected through the tools described in this statement will be used for the national cross-site evaluation of SAMHSA's SPF Rx program. This request for revision covers continued data collection through FY 2028 (three years), through the end of the grant period for grants awarded in FY 2022 and covering the a year of data collection for an anticipated FY2027 cohort. The Program Evaluation for Prevention Control (PEPC) team will systematically collect and maintain an Annual Reporting Tool (ART) and Grantee and Community Level Outcomes data modules submitted by SPF Rx grantees through the online Data Management System (DMS), and conduct telephone interviews with SPF Rx grantees.

SAMHSA is requesting approval for data collection for the SPF Rx cross-site evaluation with the following instruments:

• *Annual Reporting Tool (ART)*—The ART is a survey instrument collected yearly to monitor state, territory, tribal entity, and community-level performance, and to evaluate the effectiveness of the SPF Rx program. This tool is completed by grantees and subrecipient community project directors and provides process data related to funding use and effectiveness, organizational capacity, collaboration

with community partners, data infrastructure, planned intervention targets, evaluation, contextual factors, and sustainability.

• *Grantee and Community-Level Outcomes Modules*—These modules collect data on key SPF Rx program outcomes, including opioid prescribing patterns and provider use of PDMP. Grantees will provide outcomes data at the grantee level for their state, tribal

area, or jurisdiction, as well as at the community level for each of their subrecipient communities.

• *Grantee-Level Interview*—This qualitative interview will be administered annually to obtain information from the grantee project directors on their programs, staffing, populations of focus, infrastructure, capacity, lessons learned, and collaboration.

AVERAGE ANNUALIZED DATA COLLECTION BURDEN FOR THE PEPC DATA COLLECTION ^a

Instrument	Average number of respondents	Average number of responses per respondent	Average number of responses	Hours per response	Average burden hours	Hourly wage ^{c,d}	Annualized data collection burden
ART	^b 107	1	107	1.5	161	\$30.56	\$4,920.16
	^c 21	1	21	1.5	31	50.85	1,576.35
Total Burden for ART	128	128	192	6,496.51
Grantee-Level PDMP Outcomes Module	^c 21	1	21	2.5	52	50.85	2,627.25
Community-Level PDMP Outcomes Module	^c 21	5.2	107	1.25	134.17	50.85	6,822.38
Grantee-Level Interview	^c 21	1	21	1.5	31	50.85	1,576.35
Total Annualized Burden	191	277	409.17	17,522.49

^a *Annualized Data Collection Burden* captures the average number of respondents and responses, burden hours, and respondent cost over the 3 years (FY 2026–FY 2028).

^b Community subrecipient respondent.

^c Grantee respondent.

^d *Grantee Project Director or Evaluator hourly wage* is based on the mean hourly wage for state government managers, as reported in the 2023 Occupational Employment (OES) by the Bureau of Labor Statistics (BLS) found at https://www.bls.gov/oes/current/naics4_999200.htm#11-0000.

^e *Subrecipient Staff hourly wage* is based on the mean hourly wage for local government counselors, social workers, and other community and social service specialists, as reported in the 2023 OES by the BLS found at https://www.bls.gov/oes/current/naics4_999300.htm.

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.

Carlos Graham,
Social Science Analyst.

[FR Doc. 2026–02494 Filed 2–6–26; 8:45 am]

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INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701–TA–458 and 731–TA–1154 (Third Review)]

Kitchen Appliance Shelving and Racks From China; Scheduling of Expedited Five-Year Reviews

AGENCY: United States International Trade Commission.

ACTION: Notice.

SUMMARY: The Commission hereby gives notice of the scheduling of expedited reviews pursuant to the Tariff Act of

1930 (“the Act”) to determine whether revocation of the antidumping and countervailing duty orders on kitchen appliance shelving and racks from China would be likely to lead to continuation or recurrence of material injury within a reasonably foreseeable time.

DATES: January 26, 2026.

FOR FURTHER INFORMATION CONTACT:

Juan-Carlos Pena-Flores (202–205–3169), Office of Investigations, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436.

Hearing-impaired persons can obtain information on this matter by contacting the Commission’s TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its internet server (<https://www.usitc.gov>). The public record for this proceeding may be viewed on the Commission’s electronic docket (EDIS) at <https://edis.usitc.gov>.

SUPPLEMENTARY INFORMATION:

Background.—On January 26, 2026, the Commission determined that the

domestic interested party group response to its notice of institution (90 FR 42443, September 2, 2025) of the subject five-year reviews was adequate and that the respondent interested party group response was inadequate. The Commission did not find any other circumstances that would warrant conducting full reviews.¹ Accordingly, the Commission determined that it would conduct expedited reviews pursuant to section 751(c)(3) of the Act (19 U.S.C. 1675(c)(3)).²

For further information concerning the conduct of these reviews and rules of general application, consult the Commission’s Rules of Practice and Procedure, part 201, subparts A and B (19 CFR part 201), and part 207, subparts A, D, E, and F (19 CFR part 207).

Staff report.—A staff report containing information concerning the subject matter of the reviews has been placed in the nonpublic record, and will be made available to persons on the

¹ A record of the Commissioners’ votes, the Commission’s statement on adequacy, and any individual Commissioner’s statements will be available from the Office of the Secretary and at the Commission’s website.

² Commissioner David S. Johanson voted to conduct full reviews.