within 10 calendar days of completed data submission. If the submission is returned for incomplete data, the 10-day clock for rebate payment will restart when all necessary data is submitted.

5. Plan must ensure that 340B rebates are not denied based on eligibility or compliance concerns with diversion or Medicaid duplicate discounts, pursuant to section 340B(a)(5)(A) and (B) of the Public Health Service Act and should provide for rationale and specific documentation for reasons claims are denied (*e.g.,* nonduplication of discounts for a selected drug for which the MFP is required under the MDPNP or 340B rebate provided to another covered entity on the same claim). Rebates may not be denied for perceived lack of WAC purchases. If a manufacturer has concerns regarding Medicaid duplicate discounts, diversion, eligibility, or insufficient WAC purchases to support rebate requests, the manufacturer must raise those concerns directly with HRSA/OPA or utilize the 340B statutory mechanisms, such as audits and administrative dispute resolution, for addressing such issues. Covered entities are also afforded opportunities to raise concerns with HRSA/OPA if there are issues with rebate denials through reporting tools sent to *340Bpricing@hrsa.gov.*

6. Plan must ensure that its implementation of the Pilot is limited to using the 340B rebates model only on sales of active selected drugs for the initial price applicability years 2026 or 2027, as included on the CMS Medicare Drug Price Negotiation Selected Drug List (''List''),[41] regardless of payer, or indication, and only during the selected drug's effective dates of negotiated prices. The NDC–11s of the selected drug are included in the Pilot only to the extent they are on the List, and the selected drug is in its price applicability period in the MDPNP.

D. Data

1. All data requested as part of the Plan should be limited to only the following claim fields:

Pharmacy claims data fields	Medical claims data fields
Date of Service	Date of Service.
Date Prescribed	Claim Line Number.
Rx number	Claim Number.
Fill number	Unit of Measure.
NDC–11	NDC–11.
Quantity Dispensed	Quantity.
Prescriber ID	Rendering Physician ID.
Service Provider ID	Service Provider ID.
340B ID	340B ID.
RX BIN	Health Plan Name.
RX PCN	Health Plan ID.
	Health Plan ID Qualifier (if available).

- Data definitions for each field must be submitted with the plan for HRSA's approval to ensure consistency and make it available for covered entities. Purchasing data and encounter data requests should not be requested as part of the pilot at this time.
- For BIN, PCN, and Health Plan fields for uninsured or cash paying patients, please allow the submission in the fields to be marked ''CASH''.
- Covered entities must be permitted to resubmit data if a rebate request is deemed incomplete or missing data.
- Instructions for providing data regarding wasted or undispensed units must be provided as part of the manufacturer's plan and communicated to covered entities.

Covered entity data that is handled by technology platforms and received by manufacturers as a part of this Pilot should not be used for any purpose other than those explicitly identified in this Pilot. This limitation extends to any collecting, aggregating, sharing, or licensing of Pilot data by manufacturers or technology platforms.

Thomas J. Engels,

Administrator.

[FR Doc. 2026–15633 Filed 7–31–26; 8:45 am]

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[41] *https://www.cms.gov/files/zip/medicare-drug-price-negotiation-selected-drug-list.zip.*

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request; National Health Service Corps Scholar/Students to Service Travel Worksheet, OMB No. 0906–0087—Revision

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

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, HRSA submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period. OMB may act on HRSA's ICR only after the 30-day comment period for this notice has closed.

DATES: Comments on this ICR should be received no later than September 2, 2026.

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 Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email Samantha Miller, the HRSA Information Collection Clearance Officer, at *paperwork@hrsa.gov* or call (301) 443–3983.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: National Health Service Corps Scholar/Students to Service Travel Worksheet, OMB No. 0906–0087—Revision

Abstract: Clinicians participating in the HRSA National Health Service Corps (NHSC) Scholarship Program and the Students to Service (S2S) Loan Repayment Program use the online Travel Request Worksheet to request and receive travel funds from the federal government to, in accordance with the Public Health Service Act, section 331(c)(1), visit eligible NHSC sites to which they may be assigned.

The travel approval process is initiated when an NHSC scholar or S2S participant notifies the NHSC of an impending interview at one or more NHSC-approved practice sites. The Travel Request Worksheet is also used to initiate the relocation reimbursement process, in accordance with the Public Health Service Act, section 331(c)(3), after an NHSC scholar or S2S participant has successfully been matched to an approved practice site. Upon receipt of a completed Travel Request Worksheet, the NHSC will review and approve or disapprove the request and promptly notify the NHSC scholar or S2S participant and the NHSC logistics contractor regarding travel arrangements and authorization of the funding for the site visit or relocation.

A 60-day notice published in the **Federal Register** on May 20, 2026, vol. 91, No. 97; pp. 29498–99. There were no public comments.

Need and Proposed Use of the Information: This information will facilitate NHSC scholar and S2S participants' receipt of federal travel funds that are used to visit high-need NHSC-approved practice sites. The Travel Request Worksheet is also used to initiate the relocation process after an NHSC scholar or S2S participant has successfully matched to an approved practice site.

Likely Respondents: Clinicians participating in the NHSC Scholarship Program and the S2S Loan Repayment Program.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install, and utilize technology and systems for the purpose of collecting, validating, and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

Total Estimated Annualized Burden Hours:

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Travel Request Worksheet	400	2	800	0.0667	53.36
Total	400		800		53.36

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2026–15596 Filed 7–31–26; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Mental Health; Notice of Closed Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the Board of Scientific Counselors, National Institute of Mental Health.

The meeting will be closed to the public as indicated below in accordance with the provisions set forth in section 552b(c)(6), Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual intramural programs and projects conducted by the National Institute of Mental Health, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Board of Scientific Counselors, National Institute of Mental Health.

Date: September 16–18, 2026.

Time: September 16, 2026, 12:00 p.m. to 4:10 p.m.

Agenda: To review and evaluate personnel qualifications and performance, and competence of individual investigators.

Address: Porter Neuroscience Research Center, Building 35A, Room GE620/630, 35 Convent Drive, Bethesda, MD 20892.

Meeting Format: In Person and Virtual Meeting.

Time: September 17, 2026, 9:30 a.m. to 6:20 p.m.

Agenda: To review and evaluate personnel qualifications and performance, and competence of individual investigators.

Address: Porter Neuroscience Research Center, Building 35A, Room GE620/630, 35 Convent Drive, Bethesda, MD 20892.

Meeting Format: In Person and Virtual Meeting.

Time: September 18, 2026, 10:00 a.m. to 2:20 p.m.

Agenda: To review and evaluate personnel qualifications and performance, and competence of individual investigators.

Address: Porter Neuroscience Research Center, Building 35A, Room GE620/630, 35 Convent Drive, Bethesda, MD 20892.

Meeting Format: In Person and Virtual Meeting.

Contact Person: Jennifer E. Mehren, Ph.D., Scientific Advisor, Division of Intramural Research Programs, National Institute of Mental Health, NIH, 35A Convent Drive, Room GE 412, Bethesda, MD 20892, 301–496–3501, *mehrenj@mail.nih.gov*.

Dated: July 30, 2026.

Rosalind M. Niamke,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2026–15682 Filed 7–31–26; 8:45 am]

BILLING CODE 4167–05–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the AIDS Research Advisory Committee, NIAID.

The meeting will be open to the public, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting. The meeting can be accessed from the NIH Videocast at the following link: *https://videocast.nih.gov/.*

Name of Committee: AIDS Research Advisory Committee, NIAID.

Date: September 23, 2026.

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

Agenda: Report of Division Director and Division Staff.

Address: National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Grand Hall, Rockville, MD 20892 (Virtual Meeting).

Contact Person: Martin Gutierrez, Program Coordinator, Scientific Planning and Operations, Division of AIDS, National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 8D50, Rockville, MD 20892, 240–292–4844, *mgutierrez@mail.nih.gov*.

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

Dated: July 30, 2026.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2026–15696 Filed 7–31–26; 8:45 am]

BILLING CODE 4167–05–P