ESTIMATED ANNUALIZED BURDEN HOUR TABLE—Continued

Type of respondent	Form name	Number of respondents	Number responses per respondent	Average burden per response (in hours)	Total burden hours
QHIN	QHIN Quarterly Report	15	4	2	120
QHIN	QHIN Attestation	15	1	1	15
QHIN	TEFCA Directory Submission (weekly).	15	1040	0.033	520
QHIN	Program Evaluation or Usability Testing.	15	4	4	240
QHIN	Other	15	4	2	120
Total					1,420

Authority: 44 U.S.C. 3501 *et seq.*

Catherine Howard,

Paperwork Reduction Act Reports Clearance Officer, Office of the Secretary.

[FR Doc. 2026–16766 Filed 8–14–26; 8:45 am]

BILLING CODE 4150–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Data Use Certification for the NIH National Institute on Drug Abuse Data Share Site

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The purpose of this notice is to inform the public of the information collection requirement for the NIDA Data Share, the data use terms and conditions, and to request a progress report statement at the time of renewal. In compliance with the requirement of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Institute on Drug Abuse (NIDA) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

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: Dr. Ming Zhan, Director, Office of Data Science and Coordination, National Institute on Drug Abuse, 3WFN Room 09D14, 11601 Landsdown St, Rockville MD 20852, or call non-toll-free number (301) 480–3506 or email your request, including your address to: *ming.zhan@nih.gov.*

Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) 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) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Proposed Collection Title: Data Use Certification for the NIDA Data Share, OMB#0925–NEW, National Institute on Drug Abuse (NIDA), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of this proposal is to inform data requestors about terms and conditions for using data stored in the NIDA Data Share System, and to obtain signed agreements from requestors and their institutional officials attesting to their commitment to abide by the data use terms and conditions. These include using data for research purposes; adhering to human subjects research requirements; not distributing the data to non-authorized users; minimizing risk of participant identifiability; using data ethically and responsibly; and keeping the data secure. Recipients must include a brief description of their research project and submit their signed data use agreements to the data repository to gain access to Data Share data.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 127.

Estimated Annualized Burden Hours

ANNUAL BURDEN HOURS ESTIMATE

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
Data Request	100	1	1	100
Data Request Renewal Annual Progress Report	40	1	30/60	20
Data Request Renewal	40	1	10/60	7
Total	180	180		127

Dated: August 12, 2026.

Thomas Clarke,

Project Clearance Liaison, National Institute on Drug Abuse, National Institutes of Health.

[FR Doc. 2026–16683 Filed 8–14–26; 8:45 am]

BILLING CODE 4167–05–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection: 30-Day Comment Request; Specimen Resource Locator (National Cancer Institute)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995 to provide an opportunity for public comment on proposed data collection projects, the National Institutes of Health, National Cancer Institute (NCI) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

DATES: Comments regarding this information collection are best assured of having their full effect if received by September 16, 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 30-day Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Melissa Park, PRA Liaison, Office of Management Policy and Compliance, National Cancer Institute, 9609 Medical Center Drive, Room 2E196, Bethesda, MD 20892 or call non-toll-free number (240) 276–5717 or email your request, including your address to: *vivian.horovitchkelley@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 June 12, 2026 (Vol. 91, No. 113, FR 35694) 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 Cancer Institute (NCI), 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 Title: Specimen Resource Locator (NCI), 0925–0703: Expiration Date 08/31/2026, EXTENSION, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: Department of Health and Human Services (DHHS), National Institutes of Health (NIH), and National Cancer Institute (NCI) seek to obtain OMB approval to extend the Specimen Resource Locator (SRL) collection for an additional three (3) years. The availability of specimens and associated data is critical to increase our knowledge of cancer biology and to translate important research discoveries into clinical applications. The discovery and validation of cancer prevention markers require access, by researchers, to quality clinical biospecimens. In response to this need, the National Cancer Institute's (NCI) Cancer Diagnosis Program has developed and is expanding a searchable database: Specimen Resource Locator (SRL). The SRL allows scientists in the research community and the NCI to locate specimens needed for their research. The SRL lists non-commercial, either NCI or non-NCI-supported human biorepositories and their links. This administrative submission is an online form that collects information to manage and improve a program and its resources for the use of all scientists. This submission does not involve any hypothesis-driven analysis or research; only descriptive program-management metrics are used.

OMB approval is requested for three years. There are no costs to respondents other than their time. The estimated annualized burden hours are 107.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hour
Private Sector	Initial Request (Attachment 2)	70	1	5/60	6
State Government		70	1	5/60	6
Federal Government		60	1	5/60	5
Private Sector	Enter Resource (Attachment 3)	30	1	7/60	4
State Government		30	1	7/60	4
Federal Government		20	1	7/60	2
Private Sector	Enter Collection (Attachment 4)	30	3	18/60	27
State Government		30	3	18/60	27
Federal Government		20	3	18/60	18
Private Sector	Annual Update (Attachment 5)	30	1	5/60	3
State Government		30	1	5/60	3
Federal Government		20	1	5/60	2
Total			600		107