296. Caroline Moore, Longview, Texas, Court of Federal Claims No: 25–2135V
297. William Olaschinez, Marmora, New Jersey, Court of Federal Claims No: 25–2137V
298. Dominique Taylor on behalf of P.T., Tampa, Florida, Court of Federal Claims No: 25–2138V
299. Lucy Dorego, Fall River, Massachusetts, Court of Federal Claims No: 25–2139V
300. Timur Sarac, Columbus, Ohio, Court of Federal Claims No: 25–2140V
301. Erica Love, Aubrey, Texas, Court of Federal Claims No: 25–2142V
302. Julie Bard, Washington, District of Columbia, Court of Federal Claims No: 25–2144V
303. Thomas Carr, Oswego, New York, Court of Federal Claims No: 25–2146V
304. Patricia Hannum, Charlotte, North Carolina, Court of Federal Claims No: 25–2147V
305. Julie Canterbury, Overland Park, Kansas, Court of Federal Claims No: 25–2148V
306. Audrey Kim Griffin, Centennial, Colorado, Court of Federal Claims No: 25–2149V
307. Susan Friedlander, Boston, Massachusetts, Court of Federal Claims No: 25–2150V
308. Denise Bolton-Cromwell, Wynnewood, Pennsylvania, Court of Federal Claims No: 25–2152V
309. Melissa Amstadt, Rio Rancho, New Mexico, Court of Federal Claims No: 25–2153V
310. Julie Johnson, Downers Grove, Illinois, Court of Federal Claims No: 25–2154V
311. Nancy Crozier, Columbia, Tennessee, Court of Federal Claims No: 25–2155V
312. Sergey Pirogov, West Springfield, Massachusetts, Court of Federal Claims No: 25–2157V
313. Rosemane Joseph, Miami, Florida, Court of Federal Claims No: 25–2159V
314. Sarah Keegans, Chicago, Illinois, Court of Federal Claims No: 25–2164V
315. May Lei, Brookfield, Wisconsin, Court of Federal Claims No: 25–2165V
316. Khadija Gilkey, Clarksville, Tennessee, Court of Federal Claims No: 25–2166V
317. Ellen Burton, Galveston, Texas, Court of Federal Claims No: 25–2167V
318. Amanda Pittman, Washington, District of Columbia, Court of Federal Claims No: 25–2168V
319. Katelyn Brooks, Katy, Texas, Court of Federal Claims No: 25–2171V
320. Ryan Lee Beasley on behalf of the estate of Tommy Lee Beasley, Deceased, High Point, North Carolina, Court of Federal Claims No: 25–2172V
321. Stanley Hill, Palm Springs, California, Court of Federal Claims No: 25–2173V
322. April Wright, Galesburg, Illinois, Court of Federal Claims No: 25–2175V
323. Elizabeth Boston, Cincinnati, Ohio, Court of Federal Claims No: 25–2180V
324. Lisa Osterhoudt, Lebanon, Tennessee, Court of Federal Claims No: 25–2182V
325. Khalif Quran, Bonifay, Florida, Court of Federal Claims No: 25–2183V
326. Paul James Pope Sr., Hampstead, Maryland, Court of Federal Claims No: 25–2184V
327. William Korang, Forest Hills, New York, Court of Federal Claims No: 25–2187V
328. Gregory Centaro, East Orange, New Jersey, Court of Federal Claims No: 25–2188V
329. Suzanne Rankin, Cocoa Beach, Florida, Court of Federal Claims No: 25–2189V
330. Erik Halbig, Nashville, Tennessee, Court of Federal Claims No: 25–2190V

[FR Doc. 2026–01963 Filed 1–30–26; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Public Comment Request; Information Collection Request Title: Rural Northern Border Region Outreach Program Performance Measures Report

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

ACTION: Notice.

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on this ICR should be received no later than April 3, 2026.

ADDRESSES: Submit your comments to *paperwork@hrsa.gov* or mail the HRSA Information Collection Clearance Officer, Room 13N82, 5600 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email *paperwork@hrsa.gov* or call Samantha Miller, the HRSA Information Collection Clearance Officer, at (301) 443–3983.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the ICR title for reference.

Information Collection Request Title: Rural Northern Border Region Outreach Program Performance, OMB No. 0915–xxxx–[New]

Abstract: The Rural Northern Border Region Outreach Program (RNBR–OP) is authorized under 42 U.S.C. 254c(e) (Section 330A(e) of the Public Health Service Act) to promote the delivery of health care services to rural underserved populations in Northern Border Regional Commission counties in Maine, New Hampshire, New York, and Vermont. This authority permits HRSA to ''award grants to eligible entities to promote rural health care services outreach by improving and expanding the delivery of health care services to include new and enhanced services in rural areas through community engagement and evidence-based or innovative, evidence-informed models.'' 42 U.S.C. 254c(e).

Need and Proposed Use of the Information: The purpose of the proposed data collection is to assess RNBR–OP awardees' progress toward meeting RNBR–OP goals (as stated in the authorizing statute). Additionally, HRSA will be able to monitor and assess the impact of the RNBR–OP program, and identify improvements made by RNBR–OP awardees in specific topic areas. RNBR–OP grantees will submit annual reports to HRSA on performance measures covering the following topic areas: (1) capacity/organizational information; (2) workforce training; (3) access/population demographics; (4) health status and/or quality; and (5) sustainability.

Likely Respondents: The respondents will be recipients of the RNBR–OP funding.

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
RNBR–OP Performance Measures Report	13	1	13	17	221
Total	13	1	13	17	221

HRSA specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2026–02067 Filed 1–30–26; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; NCI Genomic Data Commons (GDC) Data Submission Request Form (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 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 April 3, 2026.

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 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: *melissa.park@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: NCI Genomic Data Commons (GDC) Data Submission Request Form, 0925–0752, Expiration Date 04/30/2026, EXTENSION. National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of the NCI Genomic Data Commons (GDC) Data Submission Request Form is to continue to provide a vehicle for investigators to request the submission of their cancer genomic data into the GDC in support of data sharing. The purpose is also to provide a mechanism for the GDC Data Submission Review Committee to review and assess the data submission request for applicability to the GDC mission. The scope of the form involves obtaining information from investigators that: (1) would like to submit data about their study into the GDC, (2) are affiliated with studies that adhere to GDC data submission conditions. The benefits of the collection are that it provides the needed information for investigators to understand the types of studies and data that the GDC supports and that it provides a standard mechanism for the GDC to assess incoming data submission requests. The only change requested in this Extension is a reduction in the number of respondents from 200 to 100, resulting in a reduction in the total annual burden hours from 50 to 25. There are no other substantive changes to this submission other than the cost-of-living changes to the federal and labor costs.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 25 hours.