DATES: Comments regarding this information collection are best assured of having their full effect if received by April 13, 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: *melissa.park@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 January 12, 2026 (Vol. 91, No. 7 FR 1192) 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: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (NCI), 0925–0642, Expiration Date 03/31/2026, EXTENSION, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: This activity collects qualitative customer and stakeholder feedback efficiently and timely, per the Administration's commitment to improving service delivery. This generic provides information about the National Cancer Institute's customer or stakeholder perceptions, experiences, and expectations; provides an early warning of service issues; or focuses on areas where communication, training, or operations changes might improve product or service delivery. It also allows feedback to contribute directly to the improvement of program management. Feedback collected under this generic clearance provides valuable information but will not yield data that can be generalized to the overall population.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 9,337 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
Surveys	Individuals	27,100	1	12/60	5,420
In-Depth Interviews (IDIs) or Small Discussion Groups.	Individuals	500	1	90/60	750
Focus Groups	Individuals	1,000	1	90/60	1,500
Website or Software Usability Tests	Individuals	5,000	1	20/60	1,667
Total			33,600		9,337

Dated: March 11, 2026.

Melissa M. Park,

Project Clearance Liaison, National Cancer Institute, National Institutes of Health.

[FR Doc. 2026–04979 Filed 3–12–26; 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: 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, contact the SAMHSA Reports Clearance Officer at *samhsapra@samhsa.hhs.gov.*

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.

Proposed Project: 2026–2029 National Survey on Drug Use and Health: Methodological Field Tests (Office of Management and Budget No. 0930–0290)—Extension

The National Survey on Drug Use and Health (NSDUH) is a survey of the U.S. civilian, non-institutionalized population aged 12 years old or older. The data are used to provide estimates of substance use and mental illness at the national, state, and substate levels. NSDUH data also help to identify the extent of substance use and mental illness among different subgroups, estimate trends over time, and determine the need for treatment services. The results are used by SAMHSA, the Office of National Drug Control Policy, federal government

agencies, and other organizations and researchers to establish policy, direct program activities, and better allocate resources.

Methodological tests will continue to examine the feasibility, quality, and efficiency of new procedures or revisions to existing survey protocol. Specifically, the tests will measure the reliability and validity of certain questionnaire sections and items through multiple measurements on a set of respondents; assess new methods for gaining cooperation and participation of respondents with the goal of increasing response and decreasing potential bias in the survey estimates; and assess the impact of new sampling techniques and technologies on respondent behavior and reporting. Research will involve focus groups, cognitive testing, and field tests. Prior to each methodological test, a separate clearance memo (under this generic clearance) will be presented to Office of Management and Budget for review.

These methodological tests will continue to examine ways to increase data quality, lower operating costs, and gain a better understanding of sources and effects of non-sampling error on NSDUH estimates. Particular attention will be given to minimizing the impact of design changes so survey data can be comparable over time. If findings suggest changes that might lead to improvements to the study, current procedures or data collection instruments may be revised.

The number of respondents to be included in each field test will vary, depending on the nature of the subject being tested and the target population. However, the total estimated response burden is 14,801 hours. The exact number of subjects and burden hours for each test are unknown at this time but will be clearly outlined in each individual submission. These estimated burden hours are distributed over 3 years as follows:

ESTIMATED TOTAL BURDEN FOR NSDUH METHODOLOGICAL FIELD TESTS

Activity	Number of respondents	Responses per respondent	Total number of responses	Average burden per response (hrs.)	Total burden (hrs.)
a. Focus Groups	378	1	378	2.0	756
b. Respondent screening for a	473	1	473	0.083	39
c. Cognitive testing	420	1	420	1.0	420
d. Respondent screening for c	800	1	800	0.083	66
e. Field Tests	12,000	1	12,000	1.0	12,000
f. Household screening for e	16,200	1	16,200	0.083	1,345
g. Screening Verification for e	804	1	804	0.067	54
h. Interview Verification for e	1,800	1	1,800	0.067	121
Total	32,875	.....	32,875	.....	14,801

Send comments to SAMHSA Reports Clearance Officer, 5600 Fisher Lane, Room 15E45, Rockville, MD 20852 *OR* email him a copy at *samhsapra@samhsa.hhs.gov.* Written comments should be received by May 12, 2026.

Carlos Graham,

Social Science Analyst.

[FR Doc. 2026–04982 Filed 3–12–26; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

[Docket No. FWS–HQ–NWRS–2025–1496; FVRS34510900000–XXX–FF09R50000; OMB Control Number 1018–0174]

Agency Information Collection Activities; U.S. Fish and Wildlife Service Preliminary Land Acquisition Process

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice of information collection; request for comment.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995, we, the U.S. Fish and Wildlife Service (Service), are proposing to renew a currently approved information collection without change.

DATES: Interested persons are invited to submit comments on or before May 12, 2026.

ADDRESSES: Send your comments on the information collection request (ICR) by one of the following methods (please reference Office of Management and Budget (OMB) Control No. 1018–0174 in the subject line of your comment):

- *Internet (preferred): https://www.regulations.gov.* Follow the instructions for submitting comments on Docket No. FWS–HQ–NWRS–2025–1496.
- *U.S. mail:* Service Information Collection Clearance Officer, U.S. Fish and Wildlife Service, 5275 Leesburg Pike, MS: PRB (JAO/3W); Falls Church, VA 22041–3803.

FOR FURTHER INFORMATION CONTACT: Madonna L. Baucum, Service Information Collection Clearance Officer, by email at *Info_Coll@fws.gov,* or by telephone at (703) 358–2503. Individuals in the United States who are deaf, deafblind, hard of hearing, or have a speech disability may dial 711 (TTY, TDD, or TeleBraille) to access telecommunications relay services. Individuals outside the United States should use the relay services offered within their country to make international calls to the point-of-contact in the United States. You may also view the ICR at *https://www.reginfo.gov/public/do/PRAMain.*

SUPPLEMENTARY INFORMATION: In accordance with the Paperwork Reduction Act (PRA; 44 U.S.C. 3501 *et seq.*) and its implementing regulations at 5 CFR part 1320, all information collections require approval under the PRA. We may not conduct or sponsor, and you are not required to respond to, a collection of information unless it displays a currently valid OMB control number.

As part of our continuing effort to reduce paperwork and respondent burdens, we are again inviting the public and other Federal agencies to comment on new, proposed, revised, and continuing collections of information. This helps us assess the impact of our information collection requirements and minimize the public's reporting burden. It also helps the public understand our information collection requirements and provide the requested data in the desired format.

We are especially interested in public comment addressing the following:

(1) Whether or not the collection of information is necessary for the proper