Supporting Statement A

Generic Clearance for the Collection of Qualitative

Feedback on Agency Service Delivery (NIAID)

**OMB Number: 0925-0668, Expiration Date: 07/31/2025**

This is an extension to the original submission and all changes throughout this document are in yellow highlight

**Date**

June 2025

Check off which applies:

* New
* Revision
* Reinstatement with Change
* Reinstatement without Change

X Extension

* Emergency
* Existing

Name: Brandie Taylor
Address: 5601 Fishers Lane, Rockville, Maryland 20892

Telephone: 240.669.2096
Fax: FAX 301-480-5752

Email: taylorbr@mail.nih.gov

National Institute of Allergy and Infectious Diseases

National Institutes of Health

**Table of contents**

A. ABSTRACT

A.1 Circumstances Making the Collection of Information Necessary

A.2. Purpose and Use of the Information COLLECTION

A.3 Use of Information Technology and Burden Reduction

A.4 Efforts to Identify Duplication and Use of Similar Information

A.5 Impact on Small Businesses or Other Small Entities

A.6 Consequences of Collecting the Information Less Frequently

A.7 Special Circumstances Relating to the Guidelines of 5 CFR 1320.5

A.8 Comments in Response to the Federal Register Notice and Efforts to Consult Outside Agency

A.9 Explanation of Any Payment of Gift to Respondents

A.10 Assurance of Confidentiality Provided to Respondents

A.11 Justification for Sensitive Questions

A.12 Estimates of Hour Burden Including Annualized Hourly Costs

A.13 Estimate of Other Total Annual Cost Burden to Respondents or Record keepers

A.14 Annualized Cost to the Federal Government

A.15 Explanation for Program Changes or Adjustments

A.16 Plans for Tabulation and Publication and Project Time Schedule

A.17 Reason(s) Display of OMB Expiration Date is Inappropriate

A.18 Exceptions to Certification for Paperwork Reduction Act Submissions

 **Attachments**

1. Sub-Study Template Submission Form
2. List of Sub-study Approvals

**A. Justification**

Abstract: This extension collection of information is necessary to enable the Agency to garner customer and stakeholder feedback in an efficient, timely manner, in accordance with our commitment to improving service delivery. The information collected from our customers and stakeholders will help ensure that users have an effective, efficient, and satisfying experience with the Agency’s programs. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

**A.1 Circumstances Making the Collection of Information Necessary**

Executive Order 12862 directs Federal agencies to provide service to the public that matches or exceeds the best service available in the private sector. In order to work continuously to ensure that our programs are effective and meet our customers’ needs, the National Institute of Allergy and Infectious Diseases of the National Institutes of Health (hereafter “the Agency”) seeks to obtain OMB approval of a generic clearance to collect qualitative feedback on our service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions but are not statistical surveys that yield quantitative results that can be generalized to the population of study.

This collection of information is necessary to enable the Agency to garner customer and stakeholder feedback in an efficient, timely manner, in accordance with our commitment to improving service delivery. The information collected from our customers and stakeholders will help ensure that users have an effective, efficient, and satisfying experience with the Agency’s programs. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative, and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

**A.2 Purpose and Use of the Information Collection**

Improving agency programs requires ongoing assessment of service delivery, by which we mean systematic review of the operation of a program compared to a set of explicit or implicit standards, as a means of contributing to the continuous improvement of the program. The Agency will collect, analyze, and interpret information gathered through this generic clearance to identify strengths and weaknesses of current services and make improvements in service delivery based on feedback. The solicitation of feedback will target areas such as: timeliness, appropriateness, accuracy of information, courtesy, efficiency of service delivery, and resolution of issues with service delivery. Responses will be assessed to plan and inform efforts to improve or maintain the quality of service offered to the public. If this information is not collected, vital feedback from customers and stakeholders on the Agency’s services will be unavailable.

The Agency will only submit a collection for approval under this generic clearance if it meets the following conditions:

* Information gathered will be used only internally for general service improvement and program management purposes and is not intended for release outside of the agency (if released, procedures outlined in Section A.16 will be followed);
* Information gathered will not be used for the purpose of substantially informing influential policy decisions [[1]](#footnote-2);
* Information gathered will yield qualitative information; the collections will not be designed or expected to yield statistically reliable results or used as though the results are generalizable to the population of study ;
* The collections are voluntary;
* The collections are low-burden for respondents (based on considerations of total burden hours, total number of respondents, or burden-hours per respondent) and are low-cost for both the respondents and the Federal Government;
* The collections are non-controversial and do not raise issues of concern to other Federal agencies;
* Any collection is targeted to the solicitation of opinions from respondents who have experience with the program or may have experience with the program in the near future; and
* With the exception of information needed to provide renumeration for participants of focus groups and cognitive laboratory studies, personally identifiable information (PII) is collected only to the extent necessary and is not retained.

If these conditions are not met, the Agency will submit an information collection request to OMB for approval through the normal PRA process.

To obtain approval for a collection that meets the conditions of this generic clearance, a standardized form will be submitted to OMB along with supporting documentation (e.g., a copy of the focus group guide). The submission will have automatic approval, unless OMB identifies issues within 5 business days.

The types of collections that this generic clearance covers include, but are not limited to:

* Customer comment cards/complaint forms
* Small discussion groups
* Focus Groups of customers, potential customers, delivery partners, or other stakeholders
* Cognitive laboratory studies, such as those used to refine questions or assess usability of a website;
* Qualitative customer satisfaction surveys (e.g., post-transaction surveys; opt-out web surveys)
* In-person observation testing (e.g., website or software usability tests)

There have been 12 projects approved under this generic clearance since its approval three years ago, all contributing significantly to the mission of NIAID. The projects have consisted of a variety of customer satisfaction and feedback surveys. Attachment 2 provides a list of the information collections (sub-studies) that have been previously approved in the past three years, and indicates which sub-studies are still actively collecting data. Attachment 2 also provides a list of the information collections (sub-studies) that were approved under this generic clearance prior to its approval three years ago, and indicates which are still actively collecting data. We would like to continue collecting data for those studies that are still actively collecting data.

The Agency has established a manager/managing entity to serve for this generic clearance and will conduct an independent review of each information collection to ensure compliance with the terms of this clearance prior to submitting each collection to OMB.

**A.3 Use of Information Technology and Burden Reduction**

If appropriate, agencies will collect information electronically and/or use online collaboration tools to reduce burden.

## **A.4 Efforts to Identify Duplication and Use of Similar Information**

No similar data are gathered or maintained by the Agency or are available from other sources known to the Agency.

## **A.5 Impact on Small Businesses or Other Small Entities**

Small business or other small entities may be involved in these efforts, but the Agency will minimize the burden on them of information collections approved under this clearance by sampling, asking for readily available information, and using short, easy-to-complete information collection instruments.

## **A.6 Consequences of Collecting the Information Less Frequently**

Without these types of feedback, the Agency will not have timely information to adjust its services to meet customer needs.

## **A.7 Special Circumstances Relating to the Guidelines of 5 CFR 1320.5**

There are no special circumstances. The information collected will be voluntary and will not be used for statistical purposes.

## **A.8.1 Comments in Response to the Federal Register Notice**

In accordance with 5 CFR 1320.8(d), on November 29, 2024, a 60-day notice for public comment was published in the *Federal Register* (89 FR 94750). One public comment was received.

## **A.8.2 Efforts to Consult Outside Agency**

In addition to public comments, NIAID solicits input from stakeholders through feedback mechanisms such as those already approved by OMB, reports, annual meetings and other venues.

**A.9 Explanation of Any Payment of Gift to Respondents**

The “default” for much of this work will be not to offer incentives. However, when deemed necessary, for instance, when individuals are recruited to travel to interviewing site, respondents may be eligible for an incentive. Some participants will receive a standard nominal incentive, number of follow-up visits required, etc. The decision to provide incentive and amount provided is in keeping with standard federal and institutional guidance.

Incentive levels will vary between $15-75 USD per interaction, dependent upon the duration of the interaction (survey vs. IDI vs. home visits), target audience, geographic location, and number of visits. Inadequate respondent recruitment limits the effectiveness of the questionnaire evaluation. Requests and justification for incentives will be included in each individual collection submission.

**A.10 Assurance of Confidentiality Provided to Respondents**

If a confidentiality pledge is deemed useful and feasible, the Agency will only include a pledge of confidentiality that is supported by authority established in statute or regulation, that is supported by disclosure and data security policies that are consistent with the pledge, and that does not unnecessarily impede sharing of data with other agencies for compatible confidential use. If the agency includes a pledge of confidentiality, it will include a citation for the statute or regulation supporting the pledge. Personally Identifiable Information (PII) will not be collected.

**A.11 Justification for Sensitive Questions**

No questions will be asked that are of a personal or sensitive nature.

**A.12.1 Estimates of Hour Burden Including Annualized Hourly Costs**

A variety of instrument’s and platforms will be used to collect information from respondents. The annual burden hours requested (2,511) are based on the number of collections we expect to conduct over the requested period for this clearance. The estimated burden over the three years is 7,200 hours.

Table 12-1 Estimated Annualized Burden Hours

|  |
| --- |
| Estimated Annual Reporting Burden |
| Type of Collection | No. of Respondents | Annual Frequency per Response | Hours per Response | Total Hours |
| Customer satisfaction surveys | 4000 | 1 | 30/60 | 2000 |
| In-Depth Interviews (IDIs) or Small Discussion Groups | 20 | 1 | 90/60 | 30 |
| Individual Brief Interviews | 40 | 1 | 15/60 | 10 |
| Focus Groups | 30 | 1 | 2 | 60 |
| Pilot testing surveys | 20 | 1 | 30/60 | 10 |
| Conferences and Training Pre- and Post-surveys | 500 | 1 | 30/60 | 250 |
| Website or Software Usability Tests | 20 | 1 | 2 | 40 |
|  **Total** | **4630** | **4630** |  | **2400** |

**A.12-2 Annual Cost to respondent**

No costs are anticipated except for the respondents’ time to participate in these activities are anticipated. Estimates are based on both the historical numbers of respondents from past projects as well as projections of projects to be conducted over the next three years. The total cost burden over 3 years is estimated to be $288,000. NIAID estimates the burden of this collection of information as follows:

 Table 12-2 Annualized Cost to Respondents

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
|  | Type of Respondents | Annual Burden Hours | Total Annual Burden | Hourly Respondent Wage Rate\* | Respondent Cost |
| Customer satisfaction surveys | General Public | 1000 | 2000 | $31 | $31,000 |
| Health Professionals  | 1000 | $49 | $49,000 |
| In-Depth Interviews (IDIs) or Small Discussion Groups | General Public | 15 | 30 | $31 | $465  |
| Health Professionals  | 15 | $49 | $735 |
| Individual Brief Interviews | General Public | 5 | 10 | $31 | $155 |
| Health Professionals  | 5 | $49 | $245 |
| Focus Groups | General Public | 30 | 60 | $31 | $930 |
| Health Professionals  | 30 | $49 | $1,470 |
| Pilot testing surveys | General Public | 5 | 10 | $31 | $155 |
| Health Professionals  | 5 | $49 | $245 |
| Conferences and Training Pre- and Post-surveys | General Public | 125 | 250 | $31 | $3,875 |
| Health Professionals  | 125 | $49 | $6,125 |
| Website or Software Usability Tests | General Public | 20 | 40 | $31 | $620 |
| Health Professionals  | 20 | $49 | $980 |
| **TOTAL** |  | 2400 |  |  | $96,000 |

\*Bureau of Labor Statistics: The General Public rate was obtained from: https://www.bls.gov/oes/current/oes\_nat.htm

The Health Professionals wage rate was obtained from <https://www.bls.gov/oes/current/oes_nat.htm#29-0000>

Occupation title “Healthcare Practitioners and Technical Occupations”, occupation code 29-0000.

**A.13 Estimate of Other Total Annual Cost Burden to Respondents or Record Keepers**

There are no other costs, including capital and start-up, or operation, maintenance, and purchase costs, associated with this generic.

## **A.14 Annualized Cost to the Federal Government**

The anticipated cost to the Federal Government is approximately $127,280 annually, for a total of approximately $381,840 over the period of three years. These costs are comprised of: operational expenses (e.g., equipment, overhead, printing, postage and support staff), contractor payments and any other expense that is necessary to collect the information approved under this generic clearance.

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Cost Descriptions** | **Grade/Step** | **Salary** | **% of Effort** | **Fringe (if applicable)** | **Total Cost to Gov’t** |
| **Federal Oversight** |  |  |  |  |  |
| NIAID PRA/OMB Liaison | 14-9 | $180,484 | 5% |  | $9,024 |
| Assistant NIAID PRA/OMB Liaison | 13-3 | $127,313 | 30% |  | $38,194 |
| Contractor Staff (Project Director, Senior Researcher, Analyst, project support) |  | $702,368 | 10% |  | $70,237 |
| Operational Costs for Data Collection Activities (e.g., printing, postage, equipment), non-labor |  |  |  |  | $12,000 |
|  |  |  |  |  |  |
| **Contractor Cost** |  |  |  |  |  |
|  |  |  |  |  |  |
| Travel  |  |  |  |  |  |
| Other Cost |  |  |  |  |  |
|  |  |  |  |  |  |
| **Total** |  |  |  |  | **$129,455** |

\*the Salary in table above is cited from: https://www.opm.gov/policy-data-oversight/pay-leave/salaries-wages/salary-tables/pdf/2025/DCB.pdf and https://www.opm.gov/policy-data-oversight/pay-leave/salaries-wages/salary-tables/pdf/2025/BOS.pdf

**A.15 Explanation for Program Changes or Adjustments**

This is a request for an extension without change for 0925-0668 – Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (NIAID). There are no changes to the purpose/scope of this submission from the previously approved submission. During the last 3 years, NIAID utilized 4,475 burden hours. However, NIAID has continued to increase awareness of the customer service generic through presentations at multiple Institute-wide forums and utilization of the *Inside*NIAID electronic newsletter. Based on the interest generated and anticipated requests we expect using 7,200 burden hours at the end of the 3 years.

## **A.16 Plans for Tabulation and Publication and Project Time Schedule**

Feedback collected under this generic clearance provides useful information, but it does not yield data that can be generalized to the overall population. Findings will be used for general service improvement but are not for publication or other public release.

Although the Agency does not intend to publish its findings, the Agency may receive requests to release the information (e.g., congressional inquiry, Freedom of Information Act requests). The Agency will disseminate the findings when appropriate, strictly following the Agency's "Guidelines for Ensuring the Quality of Information Disseminated to the Public.” and will include specific discussion of the limitation of the qualitative results discussed above.

**A.17 Reason(s) Display of OMB Expiration Date is Inappropriate**

We are requesting no exemption.

**A.18 Exceptions to Certification for Paperwork Reduction Act Submissions**

 None

1. As defined in OMB and agency Information Quality Guidelines, “influential” means that “an agency can reasonably determine that dissemination of the information will have or does have a clear and substantial impact on important public policies or important private sector decisions.” [↑](#footnote-ref-2)