The 2021 rotating sample adult core will include questions that were previously fielded in the 2019 NHIS including items on chronic pain, preventive screening tests and aspirin use. New rotating core include items on allergies and psychological distress, both of which were fielded in the pre-redesigned NHIS. New sponsored content includes items on epilepsy, myalgic encephalomyelitis/chronic fatigue syndrome (ME/CFS), insulin affordability, diabetes distress, A1C testing, colorectal cancer, prostate cancer, cervical cancer and breast cancer screening, occupational health, life satisfaction, hepatitis A and B vaccination coverage, COVID–19 vaccination coverage, and loss of the sense of taste and smell. New sponsored cancer control content that focuses on cancer screenings uses questions similar to those used in the 2019 NHIS.

The 2021 rotating sample child core will include items on stressful life events previously fielded in 2019 and on allergies, fielded in the pre-redesigned NHIS. New content included for analyses in conjunction with the adolescent follow-back study (see below) includes items on social and emotional support, bullying, health care utilization and life satisfaction.

Beginning around July 1, interviewers will ask the respondents for sample children aged 12–17 (usually the parent or guardian) for permission to contact the adolescent by web, phone, or mail and to ask follow-up questions about topics (1) already included in the sample child NHIS and (2) topics added to the sample child specifically related this follow-back. The adolescent questionnaire will be conducted web phone, or mail and include items on general health and well-being, height and weight, health care utilization, content of care in past year (or at last wellness visit), health care access, use of complementary and alternative health, physical activity, sleep, screen time, cognition, concussions, behavior, depression and anxiety, sexual orientation and gender identity, mental health care use and unmet need, social support, stressful life events, bullying, everyday discrimination, and demographics. Items on the survey environment and experience with the survey will also be asked.

Like in past years, and in accordance with the 1995 initiative to increase the integration of surveys within the DHHS, respondents to the 2021 NHIS will serve as the sampling frame for the Medical Expenditure Panel Survey conducted by the Agency for Healthcare Research and Quality. In addition, a subsample of NHIS respondents and/or members of commercial survey panels may be identified to participate in short, web-based methodological and cognitive testing activities to evaluate the questionnaire and/or inform the development of new rotating and sponsored content using web and/or mail survey tools. In the future, a subsample of NHIS respondents may also be re-contacted for a brief health exam.

There is no cost to the respondents other than their time. Clearance is sought for three years, to collect data for 2021–2023. The total annualized burden is estimated to be 42,845 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Adult Household Member	Household Roster	36,000	1	4/60
Sample Adult	Adult Questionnaire	30,000	1	48/60
Adult Family Member	Child Questionnaire	10,000	1	19/60
Adult Family Member	Methodological Projects	15,000	1	20/60
Sample Child	Adolescent follow-back Survey	1,200	1	16/60
Sample Adult	Health Exam	10,000	1	45/60
Adult Family Member	Reinterview Survey	5,500	1	5/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–21–1071; Docket No. CDC–2020–0108]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery. This Generic Information Collection enables the CDC to garner customer and stakeholder feedback on service delivery through routine surveys, focus groups, usability testing, and customer comment cards.

DATES: CDC must receive written comments on or before December 21, 2020.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2020–0108 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov*. Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *Regulations.gov*.

Please note: Submit all comments through the Federal eRulemaking portal (regulations.gov) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and

instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329; phone: 404–639–7118; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

2. Evaluate 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. Enhance the quality, utility, and clarity of the information to be collected; and

4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses.

5. Assess information collection costs.

Proposed Project

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (OMB Control No. 0920–1071, Exp. 02/28/2021)—Extension—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC/NCEZID is seeking a three-year extension of OMB control No. 0920–1071 to continue collecting routine customer feedback on agency service delivery. 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 Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention (CDC) (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.

Since the previous renewal in 2018, NCEZID has utilized 0920–1071 on 10 different occasions. The total number of responses was 15,585. The total number of burden hours was 2,525.

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 Question 16 will be followed);
- Information gathered will not be used for the purpose of substantially informing influential policy decisions; [1]
- 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 (Attachment C) will be submitted to OMB along with supporting documentation.

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)

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. CDC requests approval for an estimated 3,850 annual burden hours. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
General public	Online surveys	1,500	1	30/60	750
	Focus groups	800	1	2	1,600
	In-person surveys	1,000	1	30/60	500
	Usability testing	1,500	1	30/60	750
	Customer comment cards	1,000	1	15/60	250
Total					3,850

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-21–0199; Docket No. CDC–2020–0107]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Application for Permit to Import Biological Agents and Vectors of Human Disease into the United States, Application for Permit to Import or Transport Live Bats, and Application for Permit to Import Infectious Human Remains into the United States (OMB Control No. 0920–0199). The purpose of this data collection is to support Section 361 of the Public Health Service (PHS) Act and to prevent the introduction, transmission, or spread of communicable diseases from foreign countries into the States or possessions, or from one State or possession into any other State or possession.

DATES: CDC must receive written comments on or before December 21, 2020.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2020–0199 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov*. Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *Regulations.gov*.

Please note: Submit all comments through the Federal eRulemaking portal (*regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329; phone: 404–639–7118; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
2. Evaluate 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. Enhance the quality, utility, and clarity of the information to be collected; and
4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses.
5. Assess information collection costs.

Proposed Project

Import regulations for infectious biological agents, infectious substances, and vectors (42 CFR 71.54) (OMB Control No. 0920–0199, Exp. 4/30/