

based surveys, focus groups, and assessments. The information collection activities conducted under this extension will continue to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving 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 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 CDC and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

This type of Generic Clearance for qualitative information will not be used for quantitative purposes that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: (1) the target population to which generalizations will be made; (2) the sampling frame; (3) the sample design (including stratification and clustering); (4) the precision requirements or power calculations that justify the proposed sample size; (5) the expected response

rate; (6) the methods for assessing potential non-response bias; (7) the protocols for data collection; and (8) any testing procedures that were or will be undertaken prior fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other Generic mechanisms that are designed to yield quantitative results.

Respondents will be screened and selected from Individuals and Households, Businesses, Organizations, and/or State, Local or Tribal Government(s). The estimated annualized burden hours for this data collection activity are 9,690. There is no cost to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Type of collection	Number of respondents	Number of responses	Burden per response	Total burden
Individuals and Households, Businesses, Organizations, and/or State, Local or Tribal Government(s).	Online surveys	10,500	1	30/60	5,250
	Discussion Groups	280	1	2	560
	Focus groups	640	1	2	1,280
	Website/app usability testing.	2,000	1	30/60	1,000
	Interviews	800	1	2	1,600
Total					9,690

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

[30 Day-23-0215]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled "Application Form and Related Forms for the Operation of the National Death Index (NDI)" to the Office of Management and Budget (OMB) for review and approval. CDC previously published a "Proposed Data Collection Submitted for Public Comment and Recommendations" notice on November 16, 2022 to obtain

comments from the public and affected agencies. CDC received one comment related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) 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;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) 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 submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570. 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. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395-5806. Provide written comments within 30 days of notice publication.

Proposed Project

Application Form and Related Forms for the Operation of the National Death Index (NDI) (OMB Control No. 0920-

0215, Exp. 3/31/2023)—Revision—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 306 of the Public Health Service (PHS) Act (42 U.S.C.), as amended, authorizes that the Secretary of Health and Human Services (DHHS), acting through NCHS, shall collect statistics on the extent and nature of illness and disability of the population of the United States.

The National Death Index (NDI) is a database containing identifying death record information submitted annually to NCHS by all the jurisdiction (states and territories) vital statistics offices, beginning with deaths in 1979. Searches

against the NDI file provide the jurisdictions and dates of death, and the death certificate numbers of deceased study subjects. Using the NDI Plus service, researchers have the option of also receiving cause of death information for deceased subjects, thus reducing the need to request copies of death certificates from the jurisdictions. The NDI Plus option currently provides the International Classification of Disease (ICD) codes for the underlying and multiple causes of death for the years 1979–2021. Health researchers must complete administrative forms in order to apply for NDI services and submit records of study subjects for computer matching against the NDI file.

A three-year revision request is submitted to continue the use of the two

administrative forms (the application form and transmittal form) utilized in the operation of the National Death Index (NDI) program, along with worksheets used to calculate related fees. These forms are submitted by NDI users when applying for use of the NDI and when actually using the service. In addition, this request includes the electronic versions that replace the three paper documents, one of which will include a minor reduction in the number of data collection items.

The total estimated annual burden hours are 1,276. This represents an increase of 489 hours from 787, due primarily to the increase in applications, and transmittal forms. There is no cost to respondents except for their time.

ESTIMATES OF ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Researcher	Application Form—Electronic	282	1	150/60
Researcher	Transmittal Form—Paper/Electronic	400	3	18/60
Researcher	Early Transmittal Form—Paper/Electronic	100	3	18/60
Researcher	Fee Worksheet	450	1	15/60
Researcher	Early Release Fee Worksheet	100	1	5/60

Jeffrey M. Zirger,

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

[FR Doc. 2023-02421 Filed 2-3-23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—DP23-002, Improving Health Outcomes for Patients With Inflammatory Bowel Disease; Amended Notice of Closed Meeting

Notice is hereby given of a change in the meeting of the Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—DP23-002, Improving Health Outcomes for Patients with Inflammatory Bowel Disease; March 8, 2023, 11:00 a.m.–3:00 p.m., EST, Teleconference, in the original FRN. The meeting was published in the **Federal Register** on December 9, 2022, Volume 87, Number 236, page 75632.

The meeting is being amended to change the meeting time and should read as follows:

Date: March 8, 2023.

Time: 10:00 a.m.–3:00 p.m., EST.

The meeting is closed to the public.

FOR FURTHER INFORMATION CONTACT:

Catherine Barrett, Ph.D., Scientific Review Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 4770 Buford Highway, Mailstop S107-3, Atlanta, Georgia 30341-3717; Telephone: (404) 718-7664; Email: CBarrett@cdc.gov.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit,
Office of the Chief Operating Officer, Centers
for Disease Control and Prevention.

[FR Doc. 2023-02476 Filed 2-3-23; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day–23–22HK]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled “Surveillance of HIV-related service barriers among Individuals with Early or Late HIV Diagnoses (SHIELD)” to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on July 22, 2022, to obtain comments from the public and affected agencies. CDC received no comments to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that: