

The burden hours are estimated based on limited pilot testing conducted internally using the survey instrument. In these pilot tests, the amount of time for instruction review, collection of mock information, and the survey completion was between 2–4 minutes. The median time of three minutes was used to estimate annual burden hours. Currently, the total number of clinics which will be using this system in the

United States is unknown. However, the total number of employed miners in the metal/non-metal industry is known, with 255,702 employed in 2023. MSHA estimated in their regulatory documents that anywhere between 25%–75% of metal/non-metal miners will participate in this program, leading to an annual average number of radiographs submitted to be 13,500. If we take the total number of clinics to be at least

double the number of clinics offering NIOSH-approved radiography listed on NIOSH's website (169), then at least 338 clinics will participate.

CDC requests OMB approval for three years, with an estimated 462 annual burden hours. There are no costs to respondents other than their time to participate.

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)
Clinics and staff	Request to Access X-ray Classification Submission.	338	1	1/60	6
Clinics and staff	X-ray classification submission for metal and non-metal miners.	338	40	2/60	451
Total					462

Jeffrey M. Zirger,

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Office of Public Health Ethics and
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Disease Control and Prevention.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–25–1432; Docket No. CDC–2025–0012]

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 continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled NCEZID Rapid Message Testing & Development System. This program will enable the National Center for Emerging and Zoonotic Infectious Diseases (NCEZID) to test health messages and gather

information to inform the development of health messages.

DATES: CDC must receive written comments on or before August 15, 2025.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2025–0012 by either of the following methods:

- **Federal eRulemaking Portal:** www.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 H21–8, Atlanta, Georgia 30329.

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

Please note: Submit all comments through the Federal eRulemaking portal (www.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 H21–8, Atlanta, Georgia 30329; telephone: 404–639–7570; 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;
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; and
5. Assess information collection costs.

Proposed Project

NCEZID Rapid Message Testing & Development System (OMB Control No.

0920–1432, Exp. 5/31/2025)—
Extension—National Center for
Emerging and Zoonotic Infectious
Diseases (NCEZID), Centers for Disease
Control and Prevention (CDC).

Background and Brief Description

CDC’s National Center for Emerging
and Zoonotic Infectious Diseases
(NCEZID) offers numerous powerful
resources to anticipate, prevent, and
address outbreaks of infectious diseases.
From researchers to emergency
responders; from laboratories to
surveillance of mobile populations;
from collaborations at the Federal level
to partnerships at the local level,
NCEZID keeps people safe from threats
like anthrax, Ebola virus, Zika virus,
sepsis, mpox, and foodborne illnesses
like *Salmonella*. These efforts are vital
to protect and save lives.

The ability to effectively
communicate with the public about
these threats is one of NCEZID’s most
vital roles. Particularly during an
outbreak, it is critical that the public
understands what is happening and
why, and trusts and follows public
health leaders’ guidance. Recent public
health responses to COVID–19 and
mpox have underscored the need to
improve the speed and content of health
communications, particularly among
populations at higher risk for zoonotic
and infectious diseases. This Rapid
Message Testing & Message
Development System enables NCEZID to
collect information vital to the
development of clear, salient, relevant,
appealing, and persuasive messages
related to outbreaks and other emerging
and zoonotic diseases. The system also
allows for the relatively rapid testing of
messages when the need arises within

the Center, prior to the dissemination of
those messages and associated
communications materials. The data
collection is intended to ensure NCEZID
messages are clear, salient, appealing,
and persuasive to target audiences. Data
will guide revisions to existing or draft
messages, inform the development of
new messages, and otherwise enable
message developers to make optimal
decisions about message content,
format, and dissemination so that
NCEZID’s messages effectively reach
and resonate with their intended
audiences.

Data collection methods proposed for
this system include in-depth interviews,
online or in-person focus groups, and
online surveys. CDC requests OMB
approval for an estimated 3,431 annual
burden hours. There is no cost to
respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)	Total burden (in hr)
General public	Surveys Question Bank	10,000	1	10/60	1,667
General public	Screening and Recruitment Ques- tion Bank.	2,880	1	5/60	240
Healthcare and specialty audiences	Screening and Recruitment Ques- tion Bank.	3,600	1	5/60	300
General public	Focus Groups Question Bank	288	1	2	576
Healthcare and specialty audiences	Focus Groups Question Bank	288	1	2	576
Healthcare and specialty audiences	In-Depth Interviews Question Bank	72	1	1	72
Total					3,431

Jeffrey M. Zirger,
Lead, Information Collection Review Office,
Office of Public Health Ethics and
Regulations, 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–25–1317; Docket No. CDC–2025–
0023]

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 continuing information collection, as
required by the Paperwork Reduction
Act of 1995. This notice invites
comment on a proposed information
collection project titled National
Healthcare Safety Network (NHSN)
Respiratory Data. This data collection is
designed to standardize the data
elements collected across the country
regarding the impact of respiratory
viruses on healthcare facilities.

DATES: CDC must receive written
comments on or before August 15, 2025.

ADDRESSES: You may submit comments,
identified by Docket No. CDC–2025–
0023 by either of the following methods:

- *Federal eRulemaking Portal:*
www.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 H21–8, Atlanta,
Georgia 30329.

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

Please note: Submit all comments
through the Federal eRulemaking portal
(*www.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
H21–8, Atlanta, Georgia 30329;
Telephone: 404–639–7570; 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