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,**

*Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.*

[FR Doc. 2025–10899 Filed 6–13–25; 8:45 am]

**BILLING CODE 4163–18–P**

## 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 Question Bank. | 2,880 | 1 | 5/60 | 240 |
| Healthcare and specialty audiences | Screening and Recruitment Question 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