

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. 2024–12232 Filed 6–3–24; 8:45 am]

BILLING CODE 4163–18–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day–24–0900; Docket No. CDC–2024–  
0045]

#### 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 Contact Investigation Outcome  
Reporting Forms. The project includes  
the contact investigation outcome  
reporting forms used to obtain data from  
State, local, and territorial public health  
professionals or conveyance operators  
and medical professionals on their  
contact tracing efforts to better assess  
the risk to individuals who may have  
been exposed to a confirmed case of a  
communicable disease of public health  
concern while traveling to or within the  
United States.

**DATES:** CDC must receive written  
comments on or before August 5, 2024.

**ADDRESSES:** You may submit comments,  
identified by Docket No. CDC–2024–  
0045 by either of the following methods:

- *Federal eRulemaking Portal:*  
[www.regulations.gov](http://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](http://www.regulations.gov).

*Please note:* Submit all comments  
through the Federal eRulemaking portal  
([www.regulations.gov](http://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](mailto: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

Contact Investigation Outcome  
Reporting Forms (OMB Control No.  
0920–0900, Exp. 8/31/2024)—  
Revision—National Center for Emerging  
and Zoonotic Infectious Diseases

(NCEZID), Centers for Disease Control  
and Prevention (CDC).

#### Background and Brief Description

This proposed information collection  
project includes the contact  
investigation outcome reporting  
information collection tools used by  
CDC to better assess the risk to interstate  
and arriving international travelers who  
may have been exposed to a confirmed  
case of a communicable disease of  
public health concern while traveling.  
Different forms are tailored for different  
diseases of public health concern that  
are tracked by the Division of Global  
Migration Health (DGMH/CDC). The  
information will be used to assist and  
collaborate with State, local, and  
territorial health departments,  
conveyance operators, air, maritime,  
and land port of entry partners, and  
international public health authorities  
to identify potential exposures and to  
determine the risk of infection, and  
whether future public health  
interventions are needed.

Methods used to collect the  
information are basic surveys of  
respondents that record information  
about the exposed traveler's location  
and activities on air or maritime  
conveyances or land border crossing,  
other potential exposures, signs/  
symptoms that may have occurred after  
their potential exposure, prior history of  
vaccination or disease, and other  
medical conditions that could influence  
the risk of infection or severity of  
illness.

Due to the COVID–19 pandemic, CDC  
modified how cruise ships report  
information on cases of influenza-like-  
illness. Since December 2023, CDC has  
been using a new surveillance system  
for cumulative acute respiratory illness  
(ARI) reporting (under OMB Control  
Number 0920–1335), which includes  
cases of influenza, COVID–19, and  
respiratory syncytial virus. One form  
(the Influenza Outbreak Enhanced Data  
Collection Form) is not used routinely  
now; however, this form may be used in  
limited circumstances in the future. The  
burden of outcome reporting forms has  
been adjusted based on more recent  
numbers, which notably excludes  
COVID–19 numbers because we are no  
longer requiring contact investigations  
for COVID–19 and thus these are lower  
than past estimates. This applies  
primarily to the General Contact  
Investigation Outcome Reporting Form.  
Other burden estimates have been  
adjusted to reflect current estimates  
reflecting 2021–2023 numbers.

CDC requests OMB approval for an  
estimated 50 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 response	Average burden per response	Total burden hours
Cruise Ship Physicians/Cargo Ship Managers.	Clinically Active TB Contact Investigation Outcome Reporting Form—Maritime.	17	1	20/60	6
Cruise Ship Physicians/Cargo Ship Managers.	Varicella Investigation Outcome Reporting Form.	74	1	20/60	25
Cruise Ship Physicians .....	Influenza Outbreak Enhanced Data Collection Form—Maritime.	20	1	20/60	7
State/Local/Territorial public health staff.	General Contact Investigation Outcome Reporting Form—Air.	8	1	5/60	1
State/Local/Territorial public health staff.	TB Contact Investigation Outcome Reporting Form—Air.	51	1	5/60	4
State/Local/Territorial public health staff.	Measles Contact Investigation Outcome Reporting Form—Air.	72	1	5/60	6
State/Local/Territorial public health staff.	Rubella Contact Investigation Outcome Reporting Form—Air.	5	1	5/60	1
State/Local/Territorial public health staff.	General Contact Investigation Outcome Reporting Form—Land.	2	1	5/60	0
<b>Total .....</b>	.....	.....	.....	.....	<b>50</b>

**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. 2024-12235 Filed 6-3-24; 8:45 am]

BILLING CODE 4163-18-P

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[60Day-24-1108; Docket No. CDC-2024-0041]

**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 Paul Coverdell National Acute Stroke Program (PCNASP). This data collection is designed to monitor trends in stroke and stroke care, with the ultimate mission of

improving the quality of care for stroke patients in the United States.

**DATES:** CDC must receive written comments on or before August 5, 2024.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2024-0041 by either of the following methods:

- *Federal eRulemaking Portal:* [www.regulations.gov](http://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](http://www.regulations.gov).

*Please note:* Submit all comments through the Federal eRulemaking portal ([www.regulations.gov](http://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](mailto: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.