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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–25–0213; Docket No. CDC–2025–
0027]

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 Vital
Statistics Report Form. This data
collection is used to report annual
counts of marriages and divorces/
annulments to the Federal government
in support of the National Vital
Statistics System.

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

ADDRESSES: You may submit comments,
identified by Docket No. CDC–2025–
0027 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

National Vital Statistics Report Form
(OMB Control No. 0920–0213, Exp. 07/

31/2025)—Extension—National Center
for Health Statistics (NCHS), Centers for
Disease Control and Prevention (CDC).

Background and Brief Description

The compilation of national vital
statistics by the Federal government
dates to the beginning of the 20th
century. To administer these functions,
the National Office of Vital Statistics
was established in the Public Health
Service in April 1953. In August of
1960, the National Office of Vital
Statistics was reorganized as the
Division of Vital Statistics in the newly
created National Center for Health
Statistics (NCHS), which is now part of
the Centers for Disease Control and
Prevention (CDC).

One of the functions of NCHS is to
plan and administer a program to
provide statistics on births, deaths, fetal
deaths, marriages, and divorces reported
in the National Vital Statistics System.
This includes promoting the uniform
collection of data on these events and
providing technical assistance to the
registration areas; conducting follow
back surveys to expand the scope of
national vital statistics beyond the data
available from vital records; preparing
life tables and analyses of life table
phenomena; and investigating the
quality and reliability of data and
methodology.

One part of this function is to provide
national final counts of marriage, and
divorce occurrences following the end
of each year. The collection of the data
is authorized by 42 U.S.C. 242k.
Provisional counts of marriages and
divorces are disseminated
electronically. This form is the sole
source of final counts for these two
events. These data have been published
since 1937 and are the sole source of
this information at the national level.
The data are used by the Department of
Health and Human Services (HHS) and
by other government, academic, and
private research organizations in
tracking changes in trends of vital
events. The counts of events requested
on the form are necessary to the
administration of this portion of the
program.

CDC requests OMB approval for an
estimated 46 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)
State, Territory, and New Mexico County Officials.	Annual Vital Statistics Occurrence Report.	91	1	30/60	46
Total					46

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–25–0600; Docket No. CDC–2025–0015]

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, CDC Model Performance Evaluation Program (MPEP) for *Mycobacterium tuberculosis* Drug Susceptibility Testing. CDC is requesting a three-year approval for extension of the currently approved project used to monitor and evaluate performances and practices among national laboratories' *M. tuberculosis* susceptibility testing.

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

ADDRESSES: You may submit comments, identified by Docket No. CDC–2025–0015 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

CDC Model Performance Evaluation Program (MPEP) for *Mycobacterium tuberculosis* Drug Susceptibility Testing (OMB Control No. 0920–0600, Exp. 9/30/2025)—Extension—National Center for HIV, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The CDC is requesting an Extension of a currently approved information collection request (ICR) titled CDC Model Performance Evaluation Program (MPEP) for *Mycobacterium tuberculosis* (TB) Drug Susceptibility Testing for a period of three years. The Extension submitted for this ICR will not require changes in the scope of the project.

As part of the Extension, CDC is requesting a non-substantive change to the title of the data collection to CDC Model Performance Evaluation (MPEP) for *Mycobacterium tuberculosis* Drug Susceptibility Testing to reflect that nontuberculous mycobacteria are no longer included in the program.

While the overall number of cases of TB in the U.S. has remained fairly stable, rates still remain high among foreign-born persons, persons in correctional facilities, homeless populations, and individuals infected with HIV in major metropolitan areas. To reach the goal of eliminating TB, the Model Performance Evaluation Program for *Mycobacterium tuberculosis* drug susceptibility testing is used to monitor and evaluate performance and practices among U.S. laboratories performing *M. tuberculosis* susceptibility testing. Participation in this program is one way