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Federal Register Notice (FRN) Publication Request Worksheet

BRIEF

Agency (Select one)

- [x] CDC
- [ ] ATSDR

Office of Management and Budget (OMB) Control Number: 0920-1092

9/30/2025
Expiration Date (mm/dd/yyyy)

Agency Information Collection Request (ICR) Tracking Number (CDC ID #): _____

Project Title: Sudden Death in the Young

Requesting CDC/ATSDR CIO and Program:

National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Division or Reproductive Health

Was this proposed information collection request package vetted through an internal CIO clearance process with relevant project and Paperwork Reduction Act oversight officials? [x] Yes [ ] No

FRN Type (Publication Requested) (Select one)

- [ ] 60-Day FRN
- [x] 30-Day FRN
- [ ] Emergency Review FRN
- [ ] Other

FRN Approval Category (Select one)

- [x] Routine
- [ ] Non-Routine

Non-routine = Urgent collections (either a public health emergency or soon to expire OMB approval). Routine = Everything else.

Use of Information collection (Select one)

- [ ] Application for Benefit
- [ ] Program Evaluation
- [ ] General Purpose Statistics
- [ ] Regulatory/Compliance
- [ ] Program Planning/Management
- [ ] Public Health/Emergency Response
- [ ] Research
- [x] Surveillance/Surveillance Core
- [ ] Functions Service Delivery/Customer
- [ ] Feedback Administrative
- [ ] Audit
- [ ] Other

Type of ICR (Select one)

- [ ] New collection (Request for a new OMB Control Number)
- [x] Extension without change of a currently approved collection
- [ ] Revision of a currently approved collection *[1]
- [ ] Reinstatement without change of a previously approved collection
- [ ] Reinstatement with change of a previously approved collection
- [ ] Existing collection in use without an OMB Control Number

[1] *For Revision Requests, in the Brief Summary section below, explain what has changed:

- Is there a change in the data collection instrument? Why? What caused the change?
- Is there an increase or decrease in respondents? From what (i.e., current approval) to what? What is the reason for the change?
- Is there an increase or decrease in burden? From what (i.e., current approval) to what? What is the reason for the change?

Requested Approval Period for proposed ICR (Select one)

- [x] Three years from approval date
- [ ] Two years from approval date
- [ ] One year from approval date
- [ ] Six months from approval date (Maximum for Emergency reviews)
- [ ] Other

The proposed data collection is in support of a: (Select one)

Provide the Title, Contract Number or Funding Announcement (FA) Number and Grant Number

- [ ] Contract:
- [x] Grant/CoAg: DP23-0006
- [ ] Other - Specify:

Who will collect the data? (Select all that apply)

- [ ] CDC
- [x] Grantees
- [ ] Public Health Partners
- [ ] Contractors
- [ ] Other

AFFECTED PUBLIC: Choose all that apply

- [ ] Individuals and Households
- [x] State, Local, or Tribal Governments
- [ ] Federal Government
- [x] Private Sector - If affected Public is Private Sector, check all the following that apply:

Is the proposed ICR related to the Affordable Care Act (PPACA, P.L. 111-148 &111-152)? [ ] Yes [x] No

Does the proposed collection pose burdens on practicing physicians or their patients? [x] Yes [ ] No

If yes, identify burden type below.

BURDEN TYPE

[x] Time [ ] Effort [ ] Financial Resources

LEGAL STATUTES

Authorizing Statute(s):
Public Health Service Act, as amended Section 301(a) and Section 317K 42 USC 241(a), 42 USC 247b-12
Sudden Death Data Enhancement and Awareness Act, Public Law Number 113-236 (enacted on 12/18/14)
Note: Authorizing Statuses include applicable Public Law, U.S. Code, Executive Orders, and Statuses

RULEMAKING

Associated Rulemaking Information: Yes [ ] No [x]

FR NOTICES / COMMENTS (For 30-Day FRN Requests)

60-day Notice: Federal Register Citation: Volume 90 No. 114 Page # 25288 Publication Date: June 16, 2025

Did the Agency receive public comments on the 60-day FRN? [ ] Yes [x]

No If yes, how many comments were received? _____

Of the comments received, how many did the CDC/ATSDR program consider substantive? _____

BRIEF SUMMARY OF INFORMATION COLLECTION

State information collection's purpose and the importance of collecting this information now:

The purpose of the Sudden Death in the Young (SDY) Case Registry is to improve and standardize case ascertainment so that funded jurisdictions can better their understanding of the incidence and risk factors for sudden death in youth.

State proposed use of collected data:

Data will inform our understanding of the epidemiology of SDY, including the incidence, and risk factors. These elements are used to inform prevention strategies as well as development of best practices for multi-state surveillance of SDY.

Provide location(s) of data collection activities:
The Cooperative Agreement currently funds 12 sites: South Carolina Philadelphia County, PA, Delaware, Ohio, Indiana, Minnesota, New Hampshire, Nevada, California, Utah, and Virginia.

Describe methods for collecting data: Information collection and reporting builds on existing, state-based procedures for Child Death Review (CDR). Through their existing CDR programs, awardees compile data for an estimated 606 cases per year, on a defined set of SDY questions and enter it into the existing SDY module that is part of the existing National Fatality Review-Case Reporting System (NFR-CRS) run through the Michigan Public Health Institute's (MPHI) National Center for Fatality Review and Prevention (NCFRP) program. Data is abstracted by states/jurisdictions from primary data sources, including medical examiner/coroner reports, death investigation reports, medical records, and child protective services records. For approximately 303 of these cases per year, awardee jurisdictions also convene an advanced review team with relevant clinical expertise, including state health personnel, pediatric cardiologists, pediatric neurologists or epileptologists, and forensic pathologists. The advanced review teams comprehensively review information from multiple data sources, discuss the information, and use the information to classify cases according to a standardized algorithm that differentiates causes of death. Additional information from these advanced case reviews is entered into the SDY module. Data is shared with CDC, which uses the data to provide technical assistance to improve completeness (including missing and unknown responses) and timeliness of completed cases.

Describe sampling plan:
CDC analyzes aggregated data and assess the completeness (including missing and unknown responses), timeliness, and case ascertainment of the data set. CDC shares findings and works with SDY awardees to provide technical assistance to improve the data over time. Data analyses beyond quality improvement analyses conducted by CDC are determined by the awardees. Awardees use their data to develop targeted prevention strategies to reduce the number of infant and childhood deaths.

Collaborative Efforts:

Completely describe collaborative efforts (names, dates, roles, where documented in ICR's justification, etc.):

Name	Date	Role	Where Documented in ICR's Justification

RESPONDENTS

Total number of data collection instruments: 1

Total number of Respondents: 48

Total number of Responses: 128

Total Burden Hours: 438

(Find specific information on respondent, response, and burden estimations in the Supporting Statement)

Provide any additional comments:

OBLIGATION TO RESPOND

- ☐ Mandatory
- ☒ Required to Obtain or Retain Benefits
- ☐ Voluntary

COSTS

Annual Cost to Federal Government: $179,796

Annual Cost to Respondents: $0

(Sum/total the "Estimated Annualized Burden Costs to Respondents" in Section A12 and "Estimates of Other Total Annual Cost Burden to Respondents or Record Keepers" in Section A13 of the Supporting Statement A of the Information Collection Request)

INCENTIVES

Will CDC/ATSDR offer incentives for proposed information collection project? Yes ☐ No ☒

If yes, what type(s) or kind(s) of incentive(s) will be offered? _____

Provide the incentive amounts that will be offered to information collection respondents/participants _____

Is the incentive offered within scope of Federal/Office of Management and Budget standards for incentives? Yes ☐ No ☒

FRN Publication Approval Needed by: _____

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