

HISTORY:

86 FR 8645, February 8, 2021.

Terrell Dorn,

*Managing Director, Infrastructure Operations/
Chief Agency Privacy Officer Government
Accountability Office.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day-21-1054]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled, “Drug Overdose Response Investigation (DORI) Data Collections” to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on August 13, 2020 to obtain comments from the public and affected agencies. CDC did not receive comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) 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;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) 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 submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570. Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395-5806. Provide written comments within 30 days of notice publication.

Proposed Project

Drug Overdose Response Investigation (DORI) Data Collections (OMB Control No. 0920-1054, Exp. 03/31/2021)—Extension—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In 2015, CDC received OMB approval (OMB Control No. 0920-1054) for this Generic clearance for a three-year period to collect information to response to urgent requests from state and local health authorities to provide epidemiological information that allows for the selection of interventions to curb local epidemics of drug overdose. CDC seeks OMB approval for an Extension of this Generic clearance for a three-year period.

Drug Overdose Response Investigations (DORI) are to be conducted in response to urgent requests from state and local health authorities to provide epidemiological information that allows for the selection of interventions to curb local epidemics of drug overdose. Of particular interest is response to increasing trends in, or

changing characteristics of, overdose from prescription drugs (with a special interest in opioid analgesics such as oxycodone or methadone; benzodiazepines such as alprazolam) and/or illicit drugs (e.g., heroin). CDC’s National Center for Injury Prevention and Control (NCIPC) is frequently called upon to conduct DORIs at the request of state or local health authorities seeking support to respond to urgent public health problems resulting from drug use, misuse, addiction, and overdose. Such requests are typically, but not always, made through the Epi-Aid mechanism; in most investigations, CDC’s epidemiological response entails rapid and flexible collection of data that evolves during the investigation period.

Generic clearance is requested to ensure that timely information is collected during a DORI, which allows NCIPC to maintain critical mission function by working with state and local health authorities to protect the public’s health. During an unanticipated rise in nonfatal or fatal drug overdose where the substances responsible for the health event need to be identified, drivers and risk factors are undetermined, and/or subgroups at risk need to be identified, immediate action by CDC is necessary to minimize or prevent public harm. CDC must have the ability to rapidly deploy data collection tools to understand the scope of the problem and determine appropriate action. Procedures for each investigation, including specific data collection plans, depend on the time and resources available, number of persons involved, and other circumstances unique to the urgent conditions at hand. Data are collected by epidemiologists, psychologists, medical professionals, subject matter experts, and biostatisticians.

Data collected during a DORI are used to understand sudden increases in drug use and misuse associated with fatal and nonfatal overdoses, understand the drivers and risk factors associated with those trends, and identify the groups most affected. This allows CDC to effectively advise states on actions that could be taken to control the local epidemic. During a DORI, data are collected once, with the rare need for follow-up. The estimated annual burden hours are 2000, there are no costs 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 hours)
Drug Overdose Response Investigation Participants	DORI Data Collection Instruments	4,000	1	30/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, 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

[30Day-21-20QN]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled “Availability, Use, and Public Health Impact of Emergency Supply Kits among Disaster-Affected Populations” to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on August 28, 2020 to obtain comments from the public and affected agencies. CDC received one comment related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) 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;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) 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 submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570.

Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395-5806. Provide written comments within 30 days of notice publication.

Proposed Project

Availability, Use, and Public Health Impact of Emergency Supply Kits among Disaster-Affected Populations—New—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The National Center for Environmental Health (NCEH) is submitting a New Information Collection Request (ICR), for two-year approval. NCEH will conduct this cross-sectional study among two disaster-affected populations, at one site per year. NCEH will select geographic sites (e.g., city, town, region) for inclusion in the study after a disaster (e.g., hurricane, wildfire, flood, tornado) has occurred in the area. Parameters for site selection include a major or state-level disaster declaration for a natural disaster that affects a mid- to high-density area (e.g., population of 100,000 people) within the United States.

An all-of-society approach to disaster risk reduction emphasizes inclusion and engagement in preparedness activities.

A common recommendation is to promote household preparedness through the preparation of an emergency supply kit that can be used to shelter-in-place or during evacuation. Lack of household preparedness is a public health concern, especially in medically frail populations, because it consumes first responders’ time, taking them away from relief and recovery efforts, and can easily deplete community health resources. The Federal Emergency Management Agency (FEMA) states that individuals or households are prepared for a disaster if they have thought about and planned for the types of disaster for which they are at most risk, have developed a family communication and evacuation plan in the event of a disaster, and have assembled a complete disaster (emergency) supply kit. However, the prevalence of emergency supply kits across households in the United States ranges considerably from a community-level low of 10% to a regional high of 68%. This lack and variation of emergency supply kits across households makes household disaster preparedness a public health concern.

Self-sufficiency (defined as the ability to shelter-in-place without needing to leave your home or call for outside assistance for ~3 days following a disaster) can help reduce the demands placed on first responders during critical times, which has downstream public health impacts. Among persons with an existing physical or mental health condition at the time of the disaster, having an adequate supply of prescription and over-the-counter medications and medical supplies allows people to maintain treatment and prevent worsening or exacerbation of their existing condition or illness. It also can reduce their need for emergency medical services following a disaster. The FEMA definition of an emergency supply kit is one that can sustain each member of a household with food, water, and medication for up to three days. However, there are several knowledge gaps and challenges related to emergency supply kit use and effectiveness, including whether the current recommendations are adequate or need expansion. We identified the following gaps: