

consultants such as survey methodologist, academic researchers, clinicians or other health care providers; NHANES data or website users; members of the general public or individuals abroad who would be part of a collaborative development project

or projects between NCHS and related public health agencies in the U.S. and/or abroad. The type of participant involved in a given developmental project would be determined by the nature of the project. The details of each

project will be included in the specific GenIC submissions.
There is no cost to respondents other than their time. A three-year clearance is requested. The estimated annualized burden hours for this generic data collection is 59,465.

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)
Individuals or Households	Developmental Projects & Focus Group documents.	35,000	1	1.5	52,500
Volunteers	Developmental Projects & Focus Group documents.	300	1	1.5	450
Individuals or households, Volunteers, NHANES Participants.	24-hour developmental projects	200	1	25	5,000
NHANES participants	Developmental Projects	1,000	1	1.5	1,500
Subject Matter Experts	Focus Group/Developmental Project Documents.	15	1	1	15
Total	59,465

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-20-1011]

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 Emergency Epidemic Investigation Data Collections (OMB control number 0920-1011), for review and approval. CDC previously published a "Proposed Data Collection Submitted for Public Comment and Recommendations" notice on September 4, 2019 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 or send an email to omb@cdc.gov. 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

Emergency Epidemic Investigation Data Collections (OMB Control No. 0920-1011, Exp. 01/31/2020)—Extension—Center for Surveillance, Education, and Laboratory Services

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

Background and Brief Description

CDC previously conducted Emergency Epidemic Investigations (EELs) under Office of Management and Budget (OMB) Control Number 0920-0008. In 2013, CDC received OMB approval (OMB Control Number 0920-1011) for a new OMB generic clearance for a three-year period to collect vital information during EELs in response to urgent outbreaks or events (i.e., natural, biological, chemical, nuclear, radiological) characterized by undetermined agents, undetermined sources, undetermined transmission, or undetermined risk factors. This generic clearance was approved for a three-year extension, which expires on 1/31/2020. CDC seeks OMB approval for an extension of this generic clearance for an additional three-year period.

Supporting effective emergency epidemic investigations is one of the most important ways that CDC protects the health of the public. CDC is frequently called upon to conduct EELs at the request of local, state, or international health authorities seeking support to respond to urgent outbreaks or urgent public health-related events. In response to external partner requests, CDC provides necessary epidemiologic support to identify the agents, sources, modes of transmission, or risk factors to effectively implement rapid prevention and control measures to protect the public's health. Data collection is a critical component of the epidemiologic support provided by CDC; data are

analyzed to determine the agents, sources, modes of transmission, or risk factors so that effective prevention and control measures can be implemented. During an unanticipated outbreak or event, immediate action by CDC is necessary to minimize or prevent public harm. The legal justification for EEIs are found in the Public Health Service Act (42 U.S.C. Sec. 301 [241]) (a).

Successful investigations are dependent on rapid and flexible data collection that evolves during the investigation and is customized to the unique circumstances of each outbreak or event. Data collection elements will be those necessary to identify the agents, sources, mode of transmission, or risk factors. Examples of potential

data collection methods include telephone or face-to-face interview; email, web or other type of electronic questionnaire; paper-and-pencil questionnaire; focus groups; medical record review; laboratory record review; collection of clinical samples; and environmental assessment. Respondents will vary depending on the nature of the outbreak or event; examples of potential respondents include health care professionals, patients, laboratorians, and the general public. Participation in EEIs is voluntary and there are no anticipated costs to respondents other than their time. CDC will use the information gathered during EEIs to rapidly identify and effectively

implement measures to minimize or prevent public harm.

CDC projects 60 EEIs in response to outbreaks or events characterized by undetermined agents, undetermined sources, undetermined transmission, or undetermined risk factors annually. The projected average number of respondents is 200 per EEI, for a total of 12,000 respondents. CDC estimates the average burden per response is 0.5 hours and each respondent will be asked to respond once. Therefore, the total estimated annual burden hours are 6,000. These estimates are based on the reported burden for EEIs that have been performed during the previous two years. OMB approval is requested for three years.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Emergency Epidemic Investigation Participants.	Emergency Epidemic Investigation Data Collection Instruments.	12,000	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-20-1154]

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 CDC/ATSDR Formative Research and Tool Development 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 23, 2019 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 or send an email to omb@cdc.gov. 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

CDC/ATSDR Formative Research and Tool Development—(OMB Control No. 0920-1154, Exp. 1/31/2020)—Extension—Office of Science (OS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC) requests approval for an extension of a generic clearance for CDC/ATSDR Formative Research and Tool Development. This information collection request is designed to allow CDC to conduct formative research information collection activities used to inform many aspects of surveillance, communications, health promotion, and research project development at CDC. Formative research is the basis for developing effective strategies including communication channels, for influencing behavior change. It helps researchers identify and understand the characteristics—interests, behaviors and needs—of target populations that influence their decisions and actions.

Formative research is integral in developing programs as well as improving existing and ongoing programs. Formative research looks at the community in which a public health intervention is being or will be implemented and helps the project staff