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**

Validated Interview and Survey of Outpatient Providers on Antibiotic Stewardship Interventions (OMB Control No. 0920–1308)— Reinstatement—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

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 Validated Follow-up Interview of

Clinicians on Outpatient Antibiotic Stewardship Interventions. This collection aims to perform an interview of outpatient clinicians regarding the acceptability and perceived clinicianlevel barriers associated with our yearlong implementation of interventions designed around the Core Elements of Outpatient Antibiotic Stewardship.

Data will be collected through semistructured, in-person interviews with a sample of 40 clinicians, including nine clinicians from our original qualitative study to determine changes in perceptions over time from this baseline. In addition, we are proposing to sample an additional 31 clinicians to ensure that we do not introduce unnecessary bias and limit generalizability of the deep contextual information that would put our results at risk with a smaller sample size. Data will also be collected through a validated survey disseminated to clinicians employed by Intermountain. CDC expects about 250 clinicians to respond to our survey. Information gained from respondents to the two methods of collection will be used to refine, enhance and improve our stewardship program while allowing us to more deeply understand the unique environment and barriers found in

CDC requests OMB approval for an estimated 123 annual burden hours. There is no cost to respondents other than their time to participate.

## ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Urgent Care Clinician	Interview Guide	40 250	1 1	1 20/60

### 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. 2023–12937 Filed 6–15–23; 8:45 am]

BILLING CODE 4163-18-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Centers for Disease Control and Prevention

[60Day-23-1331; Docket No. CDC-2023-0050]

### 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 Heat-related Changes in Cognitive Performance. This data collection is designed to evaluate and assess the cognitive impacts of heat exposure on workers.

**DATES:** CDC must receive written comments on or before August 15, 2023.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2023-0050 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**

Heat-related Changes in Cognitive Performance (OMB Control No. 0920– 1331, Exp. 3/31/2024)—Revision— National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

NIOSH, under Public Law 91-173 as amended by Public Law 95 -164 (Federal Mine Safety and Health Act of 1977), and Public Law 109-236 (Mine Improvement and New Emergency Response Act of 2006) has the responsibility to conduct research to improve working conditions and to prevent accidents and occupational diseases in U.S. mines. Heat strain is one of these occupational diseases and is an increasing problem among many industries, including mining. As mines expand into deeper and hotter environments, and as heat waves occur with increasing frequency and severity, heat strain among underground and surface miners is likely to increase. Not only can heat strain lead to heat illness, but studies have demonstrated associations between heat exposure and work injuries. Although the underlying mechanism between heat exposure and injury is not known, reduced cognitive function is likely contributory. Despite the increasing importance of heat strain in mining, few studies have focused on heat strain among U.S. miners. The few studies that are available have demonstrated that miners often exceed a core body temperature of 38 °C during work activities, which is above the recommended threshold, but more information on frequency, duration, and intensity of elevated core body temperatures is needed to focus future heat strain research to better serve the mining industry.

In addition to determining the patterns of duration and intensity of heat strain among U.S. miners, investigating the additional effects of heat strain beyond the risk of heat illness is an important step in improving miner health and safety. Studies have demonstrated associations between heat stress and cognitive deficits, but substantial inter- and intraindividual variability exists in the physiologic and cognitive responses to heat exposure. More information is needed about the most important factors (e.g., age, sex, chronic disease, fitness

level, hydration) contributing to individual variability as well as interactions between these factors, because individual variability likely affects the usefulness of one-size-fits-all heat stress indices that are currently used in mining. It is also unclear which characteristics of core body temperature (e.g., absolute temperature thresholds vs. rising or falling temperatures vs rate of temperature change) are most associated with cognitive dysfunction. A better understanding of how individual variability and core body temperature relate to cognitive deficits would assist in developing strategies for screening and monitoring miners to mitigate or prevent heat strain. Therefore, this study aims to assess the following objectives: (1) Whether a core body temperature threshold exists at which cognitive performance begins to decline, (2) What factors most contribute to individual variability in cognitive and physiologic responses to heat, and (3) What patterns of duration and intensity of heat strain are most common among U.S. surface and underground miners.

To study these objectives, a dual-arm field and laboratory study will be conducted. The field study will be conducted at surface and underground mines. Data will be collected from miners working in warm or hot areas of participating mines. Participants will swallow temperature pills to measure core body temperature and will wear bio-harnesses to measure heart rate. Two six-minute assessments will be taken during each shift. The assessments include questions on sleepiness and work tasks and a Psychomotor Vigilance Test (PVT) to assess vigilant attention and reaction time. An initial screening questionnaire as well as post-shift questionnaires will be used to obtain information on risk factors for heat strain and cognitive deficits. The purpose of collecting data at the field sites is to evaluate the frequency, duration, and intensity of heat strain by monitoring core body temperature and heart rate throughout two complete shifts, as well as to assess associations between core body temperature and cognitive deficits.

The laboratory study will be conducted in an environmental chamber, in which environmental conditions can be highly controlled. Data will be collected from miners, construction workers, and firefighters. These three groups were chosen because of their risk of heat exposure and their proximity to the NIOSH laboratory where the study will be conducted. Participants will perform alternating resistance and aerobic exercises

followed by brief surveys to evaluate sleepiness (Karolinska Sleepiness Scale), affect (Positive and Negative Affect Schedule), and fatigue. Following these surveys, two cognitive tests (PVT and N-back, which measures vigilance, working memory, and complex tracking) will be administered. Testing will occur at room temperature and in hot conditions to compare cognitive test results between conditions. Participants will swallow temperature pills and wear bio-harnesses to enable the collection of real-time core body temperature and heart rate data. An initial health screening questionnaire as well as additional questionnaires administered prior to each test will be used to ensure that participants are able to withstand the physical demands of testing and to

provide information on factors that affect individual variability to heat tolerance. Additionally, a physical examination and fingerstick blood tests will be used for health screening. The purpose of collecting data in the environmental chamber is to compare physiologic and cognitive measurements at different core body temperatures to evaluate factors contributing to individual variability in cognitive and physiologic responses to heat and to evaluate whether core body temperature thresholds exist above which cognitive deficits are observed.

NIOSH is requesting a Revision for this study, because the COVID pandemic substantially delayed the ability to begin data collection. We are also making minor changes to data

collection instruments. These questions were revised to improve flow and clarity, which will likely decrease the amount of time spent on questionnaires and decrease the interruptions required of field participants. The total estimated burden hours requested are 109 for the field study and 77 for the environmental chamber study. There are no costs to respondents other than their time. All data collection activities will be conducted in full compliance with the CDC regulations to maintain the privacy of data obtained on persons and to protect the rights and welfare of human subjects. Consistent with Section 301(d) of the Public Health Service Act, a Certificate of Confidentiality (CoC) applies to this research.

### **ESTIMATED TOTAL BURDEN HOURS**

Type of respondent	Form name	Number of respondents	Number responses per respondent	Average burden per response (hours)	Total burden (hours)
	Field stud	ly			
Miners	Informed consent form (field) Initial health screening questionnaire (field).	59 59	1 1	30/60 30/60	30 30
Miners	Mid-shift field questionnaire	59 59 59	4 5 2	1/60 5/60 10/60	4 25 20
	Chamber st	udy			
Miners/firefighters/construction workers.	Informed consent form (chamber)	30	1	30/60	15
Miners/firefighters/construction workers.	Physical examination form	30	1	10/60	5
Miners/firefighters/construction workers.	Initial health screening questionnaire (chamber).	30	1	30/60	15
Miners/firefighters/construction workers.	Release of information form	5	1	1/60	1
Miners/firefighters/construction workers.	TSS and RPE	30	5	1/60	3
Miners/firefighters/construction workers.	PANAS and KSS	30	5	2/60	5
Miners/firefighters/construction workers.	Cognitive test: PVT	30	5	10/60	25
Miners/firefighters/construction workers.	Cognitive test: N-back	30	5	1/60	3
Miners/firefighters/ constructionworkers.	Pre-testing health questionnaire	30	2	5/60	5
Total					186

### Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.

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