- (3) Women who care for a young child (children ages 12 and younger; English speaking),
- (4) Women who care for a young child (children ages 12 and younger; Spanish speaking),
- (5) Women who care for an aging parent 65+ (English speaking),
- (6) Women who care for an aging parent 65+ (Spanish speaking),
- (7) Men aged 65+ with one or more chronic conditions (English speaking), and
- (8) Healthy adults 65+ (English speaking).

HCP audiences include:

(1) Emergency Medical Services personnel (English speaking),

(2) Nurse Practitioners and Physician Assistants who work at urgent care clinics (English speaking),

(3) Emergency Department triage nurses (English speaking),

(4) General medical ward staff (English speaking),

(5) Primary care physicians (English speaking),

(6) Long-term care (LTC) nurses (English speaking), and

(7) LTC medical technicians and sitters (English speaking).

This program evaluation will assist CDC in determining if the media campaign, along with partner outreach, was successful in changing awareness, knowledge, and behaviors of consumers and HCPs in select target markets. The data collected will also be used to inform future refinement and implementation of the campaign (materials and tactics).

CDC requests OMB approval for an estimated 68 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)
Consumer	Get Ahead of Sepsis Consumer Pre-test	50	1	20/60 20/60	17
Consumer HCPs	Get Ahead of Sepsis Consumer Post-test Get Ahead of Sepsis HCP Campaign Pre-	50 50		20/60	17
HCPs	test.  Get Ahead of Sepsis HCP Campaign Posttest.	50	1	20/60	17
Total					68

#### 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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BILLING CODE 4163-18-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Centers for Disease Control and Prevention

[60Day-22-0978; Docket No. CDC-2022-0012]

# 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 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 Emerging Infections Program (EIP). EIP is a population-based surveillance system designed to collect information via active, laboratory case finding that is used for detecting, identifying, and monitoring emerging pathogens.

DATES: CDC must receive written comments on or before April 1, 2022.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2022-0012 by either of the following methods:

- Federal eRulemaking Portal: 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 Regulations.gov.

Please note: Submit all comments through the Federal eRulemaking portal (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; phone: 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**

Emerging Infections Program (EIP) (OMB Control No. 0920–0978, Exp. 4/30/2022)—Revision—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Emerging Infections Programs (EIPs) are population-based centers of excellence established through a network of state health departments collaborating with academic institutions; local health departments; public health and clinical laboratories; infection control professionals; and healthcare providers. EIPs assist in local, state, and national efforts to prevent, control, and monitor the public health impact of infectious diseases.

Activities of the EIPs fall into the following general categories: (1) Active surveillance; (2) applied public health epidemiologic and laboratory activities; (3) implementation and evaluation of pilot prevention/intervention projects; and (4) flexible response to public health emergencies. Activities of the EIPs are designed to: (1) Address issues that the EIP network is particularly suited to investigate; (2) maintain sufficient flexibility for emergency

response and new problems as they arise; (3) develop and evaluate public health interventions to inform public health policy and treatment guidelines; (4) incorporate training as a key function; and (5) prioritize projects that lead directly to the prevention of disease.

A Revision is being submitted to make existing collection instruments clearer and to add several new forms specifically surveying laboratory practices. These forms will allow the EIP to better detect, identify, track changes in laboratory testing methodology, gather information about laboratory utilization in the EIP catchment area to ensure that all cases are being captured, and survey EIP staff to evaluate program quality.

CDC requests OMB approval for an estimated burden of 61,956 hours. There is no cost to respondents other than their time.

#### ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Avg. burden per response (in hours)	Total burden (in hours)
State Health Department	ABCs Case Report FormABCs Invasive Pneumococcal Disease in Children and Adults Case Report Form.	10 10	809 127	20/60 10/60	2,697 212
	ABCs <i>H. influenzae</i> Neonatal Sepsis Expanded Surveillance Form.	10	6	10/60	10
	ABCs Severe GAS Infection Supplemental Form.	10	136	20/60	453
	ABCs Neonatal Infection Expanded Tracking Form.	10	37	20/60	123
	FoodNet Campylobacter	10	970	21/60	3,395
	FoodNet Cyclospora	10	42	10/60	70
	FoodNet Listeria monocytogenes	10	16	20/60	53
	FoodNet Salmonella	10	855	21/60	2,993
	FoodNet Shiga toxin producing E. coli.	10	290	20/60	967
	FoodNet Shigella	10	234	10/60	390
	FoodNet Vibrio	10	46	10/60	77
	FoodNet Yersinia	10	55	10/60	92
	FoodNet Hemolytic Uremic Syndrome Case Report Form.	10	10	1	100
	FoodNet Clinical Laboratory Practices and Testing Volume.	10	70	20/60	233
	FluSurv-NET Influenza Hospitalization Surveillance Network Case Report Form.	10	764	25/60	3,183
	FluSurv-NET Influenza Hospitalization Surveillance Project Vaccination Phone Script Consent Form (English).	10	333	5/60	278
	FluSurv-NET Influenza Hospitalization Surveillance Project Vaccination Phone Script (Spanish).	10	333	5/60	278
	Influenza Hospitalization Surveil- lance Project Provider Vaccination History Fax Form (Children/ Adults).	10	333	5/60	278
	FluSurv-NET Laboratory Survey	10	16	10/60	26

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Type of respondent	Form name	Number of respondents	Number of responses per respondent	Avg. burden per response (in hours)	Total burden (in hours)
	HAIC—MuGSI Case Report Form for Carbapenem-resistant Enterobacteriaceae (CRE) and Acinetobacter baumannii (CRAB).	10	500	28/60	2,333
	HAIC—MuGSI Extended-Spectrum Beta-Lactamase-Producing Enterobacteriaceae (ESBL/iEC).	10	4200	25/60	17,500
	HAIC—Invasive Methicillin-resistant Staphylococcus aureus (MRSA) Infection Case Report Form.	10	340	28/60	1,587
	HAIC—Invasive Methicillin-sensitive Staphylococcus aureus (MSSA) Infection Case Report Form.	10	584	28/60	2,725
	HAIC—CDI Case Report and Treatment Form.	10	1650	38/60	10,450
	HAIC Candidemia Case Report	10	200	30/60	1,134
	HAIC—Annual Survey of Laboratory Testing Practices for C. difficile Infections.	10	16	19/60	51
	HAIC—CDI Annual Surveillance Officers Survey.	10	1	15/60	3
	HAIC—Emerging Infections Program <i>C. difficile</i> Surveillance Nursing Home Telephone Survey (LTCF).	10	45	5/60	38
	HAIC—Invasive Staphylococcus aureus Laboratory Survey.	10	11	20/60	37
	HAIC—Invasive Staphylococcus aureus Supplemental Surveillance Officers Survey.	10	1	10/60	17
	HAIC—Laboratory Testing Practices for Candidemia Questionnaire.	10	20	12/60	40
	HAIC MuGSI CA CP-CRE Health interview (new).	100	10	30/60	50
	HAIC MuGSI Supplemental Surveil- lance Officer Survey (new).	10	1	15/60	3
	HAIC Death Ascertainment Variables.	10	8	1440/60	10,080
Total					61,956

### 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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BILLING CODE 4163-18-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Centers for Disease Control and Prevention

[30Day-22-22AD]

## 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 Research Data Center Proposal (RDC) Proposal for Access to Confidential Data for the National Center for Health Statistics 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 October 25, 2021 to obtain comments from the public and affected agencies. CDC received one nonsubstantive 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