The One Health Harmful Algal Bloom System (OHHABS) was approved for data collection in 2016 and collects data on harmful algal blooms (HABs) and human and animal illnesses related to HAB exposures to support the understanding of HABs and the prevention of HAB-associated illnesses. As such, OHHABS is a centralized data source for voluntary public health surveillance of HAB events and HAB-associated illnesses using a One Health approach that takes into consideration information from the environment, animal cases, and human cases.

CDC requests OMB approval for an estimated 76 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) |
|---|---|---|---|---|---|
| OHHABS State Reporting Sites. | One Health Harmful Algal Bloom System (OHHABS) (electronic, year-round). | 57 | 4 | 20/60 | 80/60 |
| Total ..... | ..... | ..... | ..... | ..... | 76 |

**Jeffrey M. Zirger,**

*Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.*

[FR Doc. 2021–24993 Filed 11–15–21; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[30Day–22–0920]**

### 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 ''*Data Collection Through Web Based Surveys for Evaluating Act Against AIDS Social Marketing Campaign Phases Targeting Consumers,*'' 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 September 7, 2021 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**

Data Collection Through Web Based Surveys for Evaluating Act Against AIDS Social Marketing Campaign Phases Targeting Consumers (OMB Control No. 0920–0920, Exp. 11/30/2021)—Extension—National Center for HIV/AIDS, Viral Hepatitis, STD and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

In response to the continued HIV epidemic in our country, CDC launched the Let's Stop HIV Together campaign (formerly known as Act Against AIDS), a multifaceted communication campaign to reduce HIV incidence in the United States in 2009. CDC has released the campaign in phases, with some of the phases running concurrently. Each phase of the campaign uses mass media and direct-to-consumer channels to deliver messages. Some campaigns provide basic education and increase awareness of HIV/AIDS among the general public, whereas others emphasize HIV prevention and testing among specific subgroups or communities at greatest risk of infection. CDC will also develop new messages to address changes in prevention science and subpopulations affected by HIV. The proposed study will assess the effectiveness of these social marketing messages aimed at increasing HIV/AIDS awareness, increasing prevention behaviors, and improving HIV testing rates among consumers.

This Extension of an ongoing study will allow for continued evaluation of the effectiveness of Let's Stop HIV Together social marketing campaign through surveys with consumers. A total of 6,445 respondents were approved for the previously renewed Generic ICR (0920–0920) in 2018, and since the approval date, 1,000 respondents were surveyed under the GenIC, ''Development of Messages for the Let's Stop HIV Together National Campaign''. The information collected from this survey was used to evaluate the acceptability and potential effectiveness of proposed concepts, messages, and taglines for a component of the Let's

Stop HIV Together campaign focused on HIV prevention that promotes proven, effective prevention strategies, such as pre-exposure prophylaxis (PrEP) and treatment as prevention (TasP). We are requesting a one-year extension to continue surveying target audiences.

Through this extension, we plan to reach the remaining approved 5,445 respondents. To obtain the remaining respondents, we anticipate screening approximately 30,880 individuals. Depending on the target audience for the campaign phase, the study screener will vary. The study screener may address one or more of the following items: Race/ethnicity, sexual behavior, sexual orientation, gender identity, HIV testing history, HIV status, and injection drug use. Each survey will have a core set of items asked in all rounds, as well as a module of questions relating to specific Let's Stop HIV Together phases and activities.

Respondents will be recruited through national opt-in email lists, the internet, and external partnerships with community-based and membership organizations that work with or represent individuals from targeted populations (*e.g.,* National Urban League, the National Medical Association). Respondents will self-administer the survey at home on personal computers. The annual response burden is estimated at 3,751 hours. There is no cost to the 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 hr) |
|---|---|---|---|---|
| Individuals (male and female) aged 18 years and older | Study Screener | 30,880 | 1 | 2/60 |
| | Survey Module | 5,445 | 1 | 30/60 |

**Jeffrey M. Zirger,**

*Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.*

[FR Doc. 2021–24991 Filed 11–15–21; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[60Day–22–1318; Docket No. CDC–2021–0124]**

### 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 Requirement for Negative Pre-Departure COVID–19 Test Result or Documentation of Recovery from COVID–19 For All Airline or Other Aircraft Passengers Arriving into the United States from any Foreign Country and Requirement for Proof of COVID–19 Vaccination for Noncitizen, Nonimmigrant Air Passengers Arriving Into The United States From A Foreign Country. This proposed information collection is designed to ensure that CDC complies with Orders published October 25 and October 30, 2021, and to confirm that passengers who are coming into the United States via air travel have tested negative for or recently recovered from COVID–19, and that noncitizen, nonimmigrant passengers are fully vaccinated against COVID–19.

**DATES:** CDC must receive written comments on or before January 18, 2022.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC–2021–0124 by any 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,