By direction of the Commission.

April J. Tabor,

Secretary.

[FR Doc. 2023–02383 Filed 2–3–23; 8:45 am]

BILLING CODE 6750–01–P

UNITED STATES AGENCY FOR GLOBAL MEDIA

USAGM Performance Review Board Members

AGENCY: United States Agency for Global Media.

ACTION: Notice.

SUMMARY: The United States Agency for Global Media (USAGM) announces the members of its SES Performance Review Board (PRB).

ADDRESSES: USAGM Office of Human Resources, 330 Independence Ave. SW, Washington, DC 20237.

FOR FURTHER INFORMATION CONTACT: Ellona Fritschie, Senior Advisor, at *efritschie@usagm.gov* or (202) 382–7500.

SUPPLEMENTARY INFORMATION: In accordance with 5 U.S.C. 4314, USAGM publishes this notice announcing the individuals who will serve as members of the PRB for a term of one year. The PRB is responsible for: (1) reviewing performance appraisals and ratings of Senior Executive Service and Senior Level members; and (2) making recommendations on other performance management issues, such as pay adjustments, bonuses, and Presidential Rank Awards. The names, position titles, and appointment types of each member of the PRB are set forth below:

1. Yolanda Lopez, Voice of America Director, Limited Term SES
2. Grant Turner, Chief Financial Officer, Career SES
3. James Reeves, Chief Information Officer, Career SES

Dated: January 9, 2023.

Armanda Matthews,

Program Support Specialist, U.S. Agency for Global Media.

[FR Doc. 2023–02396 Filed 2–3–23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60 Day–23–1027; Docket No. CDC–2023–0008]

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 Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery. This Generic Clearance is designed to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery.

DATES: CDC must receive written comments on or before April 7, 2023.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2023–0008 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

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (OMB Control No. 0920–1027, Exp. 8/31/2023)—Extension—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC is requesting a three-year Extension for the data collection titled Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (OMB Control No. 0920–1027). During the past three-year approval period, eight GenICs consisting of 750 responses have been submitted for approval. The collections included web-

based surveys, focus groups, and assessments. The information collection activities conducted under this extension will continue to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery.

By qualitative feedback, we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training, or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative, and actionable communications between CDC and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

This type of Generic Clearance for qualitative information will not be used for quantitative purposes that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: (1) the target population to which generalizations will be made; (2) the sampling frame; (3) the sample design (including stratification and clustering); (4) the precision requirements or power calculations that justify the proposed sample size; (5) the expected response rate; (6) the methods for assessing potential non-response bias; (7) the protocols for data collection; and (8) any testing procedures that were or will be undertaken prior fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other Generic mechanisms that are designed to yield quantitative results.

Respondents will be screened and selected from Individuals and Households, Businesses, Organizations, and/or State, Local or Tribal Government(s). The estimated annualized burden hours for this data collection activity are 9,690. There is no cost to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Type of collection	Number of respondents	Number of responses	Burden per response	Total burden
Individuals and Households, Businesses, Organizations, and/or State, Local or Tribal Government(s).	Online surveys	10,500	1	30/60	5,250
	Discussion Groups	280	1	2	560
	Focus groups	640	1	2	1,280
	Website/app usability testing.	2,000	1	30/60	1,000
	Interviews	800	1	2	1,600
Total					9,690

Jeffrey M. Zirger,

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

[FR Doc. 2023–02422 Filed 2–3–23; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day–23–0215]

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 ''Application Form and Related Forms for the Operation of the National Death Index (NDI)'' 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 November 16, 2022 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

Application Form and Related Forms for the Operation of the National Death Index (NDI) (OMB Control No. 0920–