- Agenda GBAC Fall Meeting
- Updates and Introductions
- Ethics Review
- Buy Clean Implications Task Group: Proposed Advice Letter
- AI and Federal Buildings Update
- Health and Wellbeing in Federal Buildings Update
- New Committee Topics and Directions
- Public Comment
- Next Steps and Closing Comments
- *Details:* This public meeting will serve as an annual review of GBAC activities. Members will have the opportunity to ask questions about ongoing Task Group work and suggest future topics they wish to investigate.

Procedures for Attendance and Public Comment

To register to observe any or all of these public meetings, please send the following information via email to *gbac@gsa.gov:* your first and last name, organization and email address, the meeting(s) you wish to attend, and whether you would like to provide public comment.

Requests to observe meetings must be received by 5 p.m. ET on the Tuesday before the meeting in question.

For all online meetings, web meeting attendance information will be provided following registration. Time will be provided at all meetings for public comment wherever possible.

GSA will be unable to provide technical assistance to any listener experiencing technical difficulties. Testing access to the web meeting site before the calls is recommended. To request an accommodation, such as closed captioning, or to ask about accessibility, please contact Mr. Bloom at *gbac@gsa.gov* at least five business days prior to the meeting to give GSA as much time as possible to process the request.

Background

The Administrator of GSA established the Committee on June 20, 2011 (76 FR 35894) pursuant to section 494 of the Energy Independence and Security Act of 2007 (EISA, 42 U.S.C. 17123). Under this authority, the Committee provides independent policy advice and recommendations to GSA to advance Federal building innovations in planning, design, and operations to reduce costs, enable agency missions, enhance human health and performance, and minimize environmental impacts.

Kinga Hydras,

Acting Director, Office of Federal High-Performance Green Buildings, General Services Administration.

[FR Doc. 2024–16493 Filed 7–25–24; 8:45 am]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–24–0943]

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 for the Residential Care Community and Adult Day Service Center Components of the National Post-acute and Long-term Care Study'' 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 May 7, 2024 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. 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 for the Residential Care Community and Adult Day Service Center Components of the National Post-acute and Long-term Care Study (OMB Control No. 0920–0943 Exp. 07/31/2025)—Revision—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The NPALS is designed to: (1) broaden NCHS' ongoing coverage of paid, regulated long-term care (LTC) providers; (2) present alongside existing administrative data on LTC providers and service users (*i.e.,* Centers for Medicare and Medicaid Services (CMS) data on inpatient rehabilitation facilities and patients, long-term care hospitals and patients, nursing homes and residents, home health agencies and patients, and hospices and patients); (3) update data more frequently on LTC providers and service users for which nationally representative administrative data do not exist; and (4) enable comparisons across LTC sectors and timely monitoring of supply and use of these sectors over time.

Data will be collected from two types of LTC providers in the 50 states and the District of Columbia: 11,600 Residential Care Communities (RCC) and 5,500 Adult Day Service Centers (ADSC). Data were collected in 2012, 2014, 2016, 2018, 2020, and 2022. The data to be collected in 2024 include the basic characteristics, services, staffing, and practices of RCCs and ADSCs, and aggregate-level distributions of the demographics, selected health conditions and health care utilization,

physical functioning, and cognitive functioning of RCC residents and ADSC participants.

Expected users of data from this collection effort include, but are not limited to CDC; other Department of Health and Human Services (DHHS) agencies, such as the Office of the Assistant Secretary for Planning and Evaluation, The Administration for Community Living, and the Agency for Healthcare Research and Quality; associations, such as LeadingAge, National Center for Assisted Living, American Seniors Housing Association, Argentum, and National Adult Day Services Association; universities; foundations; and other private sector organizations such as the Alzheimer's Association and the AARP Public Policy Institute.

Expected burden from data collection for eligible cases is 30 minutes per respondent. An estimated 5% of RCC and ADSC respondents will have an additional five minutes of burden to complete a data retrieval call. We calculated the burden based on a 100% response rate. A two-year clearance is requested to cover the collection of data. The burden for the collection is shown in Table below and totals 4,311 hours annually. 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)
RCC Director/Designated Staff Member	RCC Questionnaire	5,800	1	30/60
ADSC Director/Designated Staff Member	ADSC Questionnaire	2,750	1	30/60
RCC/ADSC Director/Designated Staff Member.	Data retrieval call	428	1	5/60

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. 2024–16490 Filed 7–25–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–24–24HP; Docket No. CDC–2024–0056]

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 information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Compliance Attestation Statement for the Framework for Nucleic Acid Synthesis Screening. The project aims to assist providers and manufacturers of synthetic nucleic acids and benchtop nucleic acid synthesis equipment (providers) in making an attestation that they have instituted a process to screen nucleic acid sequences of concern and verify customer legitimacy, in accordance with the requirements outlaid in the OSTP Framework for Nucleic Acid Synthesis Screening.

DATES: CDC must receive written comments on or before September 24, 2024.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0056 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

Compliance Attestation Statement for the Framework for Nucleic Acid