CDC requests OMB approval for three years and an estimated 46 annual burden hours. There are no changes to the content of the information collection instrument or the estimated burden per response.

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) |
|---|---|---|---|---|---|
| PHHS Block Grant Coordinator or Designee. | PHHS Block Grant Assessment ..... | 61 | 1 | 45/60 | 46 |
| Total ..... | ..... | ..... | ..... | ..... | 46 |

**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. 2024–28688 Filed 12–5–24; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### Notice of Closed Meeting

Pursuant to 5 U.S.C. 1009(d), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), title 5 U.S.C., as amended, and the Determination of the Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, pursuant to Public Law 92–463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—CE25–032, Formative Research and Pilot-Testing of Community-led Primary Prevention Approaches to Address Elevated Risk of Intimate Partner Violence & Sexual Violence.

*Dates:* April 1–2, 2025.

*Times:* 10 a.m.–5 p.m., EDT.

*Place:* Web Conference.

*Agenda:* To review and evaluate grant applications.

*For Further Information Contact:* Carlisha Gentles, Pharm.D., B.C.P.S., C.D.C.E.S., Scientific Review Officer, National Center for Injury Prevention and Control, Centers for Disease Control and Prevention, 4770 Buford Highway NE, Mailstop S106–9, Atlanta, Georgia 30341. Telephone: (770) 488–1504; Email: *CGentles@cdc.gov.*

The Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

**Kalwant Smagh,**

*Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.*

[FR Doc. 2024–28512 Filed 12–5–24; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### Notice of Closed Meeting

Pursuant to 5 U.S.C. 1009(d), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of the Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, pursuant to Public Law 92–463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—CE25–029, Grants to Support New Investigators in Conducting Research Related to Preventing Interpersonal Violence Impacting Children and Youth.

*Dates:* March 11–12, 2025.

*Times:* 10 a.m.–5 p.m., EDT.

*Place:* Web Conference.

*Agenda:* To review and evaluate grant applications.

*For Further Information Contact:* Aisha L. Wilkes, M.P.H., Scientific Review Officer, National Center for Injury Prevention and Control, Centers for Disease Control and Prevention, 4770 Buford Highway NE, Mailstop S106–9, Atlanta, Georgia 30341. Telephone: (404) 639–6473; Email: *AWilkes@cdc.gov.*

The Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

**Kalwant Smagh,**

*Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.*

[FR Doc. 2024–28511 Filed 12–5–24; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[30Day–25–23FN]**

### 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 ''Menthol-

Flavored Tobacco Product Policy Evaluation'' 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 3, 2024 to obtain comments from the public and affected agencies. CDC received two 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**

Menthol-Flavored Tobacco Product Policy Evaluation—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

The Centers for Disease Control and Prevention (CDC) is submitting this New information collection request (ICR) for an evaluation of local policies restricting the sale of menthol-flavored tobacco products (hereafter known as menthol tobacco products). This evaluation will assess the differences in outcomes of interest between cities with policies restricting the sale of menthol tobacco products and cities without these policies. Outcomes of interest include commercial tobacco product use and related behavior, perceived access to menthol tobacco products, and menthol tobacco product-related beliefs and perceptions. This study focuses on the general population and groups who use commercial menthol tobacco products at higher rates including people who identify as lesbian, gay, bisexual, transgender, queer, or another sexual or gender minority (LGBTQ+) and racial and ethnic minority groups (persons identifying as American Indian or Alaska Native, Asian, Black or African American, Native Hawaiian or Other Pacific Islander, Hispanic or Latino, or biracial). Understanding how policies restricting the sale of menthol tobacco products impact tobacco use, behavior, and attitudes may help to inform public health activities and decisions regarding tobacco control. Although local tobacco policies have been shown to be effective at limiting the availability of policy-restricted products, no study to date has collected this type of web panel and focus group data among disparately affected population groups across jurisdictions after policy implementation. There are no other evaluation data collection efforts conducted on this topic to date, nor does the information to be collected exist in any existing centralized data source. While some questions in the web panel survey were sourced from pre-existing validated survey questions, each data collection tool proposed within this package has been designed to understand the effect of local menthol tobacco sales restriction policies on both the general population and those disproportionately impacted by menthol tobacco product use.

CDC is requesting OMB approval for three years. The total estimated annualized burden is 1,750 hours. There are no costs 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) |
|---|---|---|---|---|
| Adults in the general population who currently use or formerly used tobacco living in the cities of interest. | Survey Screener Questionnaire | 9,800 | 1 | 2/60 |
| Adults in the general population who currently use or formerly used tobacco living in the cities of interest. | Community Web-Panel Survey | 5,366 | 1 | 15/60 |
| Adults in the general population who currently use or formerly used menthol tobacco living in the intervention cities of interest. Racial and ethnic minority adults and LGBTQ+ adults meeting the above criteria identified from the Qualtrics panel. | Focus Group Screener Questionnaire | 200 | 1 | 2/60 |
| Adults in the general population who currently use or formerly used menthol tobacco living in the intervention cities of interest. Racial and ethnic minority adults and LGBTQ+ adults meeting the above criteria identified from the Qualtrics panel. | Community Focus Group | 75 | 1 | 1 |

**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. 2024–28689 Filed 12–5–24; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

**[Document Identifier: CMS–10398 #84]**

### Medicaid and Children's Health Insurance Program (CHIP) Generic Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** On May 28, 2010, the Office of Management and Budget (OMB) issued Paperwork Reduction Act (PRA) guidance related to the ''generic'' clearance process. Generally, this is an expedited process by which agencies may obtain OMB's approval of collection of information requests that are ''usually voluntary, low-burden, and uncontroversial collections,'' do not raise any substantive or policy issues, and do not require policy or methodological review. The process requires the submission of an overarching plan that defines the scope of the individual collections that would fall under its umbrella. This **Federal Register** notice seeks public comment on one or more of our collection of information requests that we believe are generic and fall within the scope of the umbrella. Interested persons are invited to submit comments regarding our burden estimates or any other aspect of this collection of information, including: the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments must be received by December 20, 2024.

**ADDRESSES:** When commenting, please reference the applicable form number (CMS–10398 #84) and the OMB control number (0938–1148). To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: CMS–10398 #84/OMB control number: 0938–1148, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/medicare/regulations-guidance/legislation/paperwork-reduction-act-1995/pra-listing.*

**FOR FURTHER INFORMATION CONTACT:** William N. Parham at 410–786–4669.

**SUPPLEMENTARY INFORMATION:** Following is a summary of the use and burden associated with the subject information collection(s). More detailed information can be found in the collection's supporting statement and associated materials (see **ADDRESSES**).

**Generic Information Collection**

1. *Title of Information Collection:* PACE Medicaid Capitation Rate Setting Guide; *Type of Information Collection Request:* New information collection request information request; *Use:* The Program of All-inclusive Care for the Elderly (PACE) is a fully integrated Medicare program and Medicaid state plan option that provides community-based care and services to individuals aged 55 or older who meet a state's nursing home level of care criteria. PACE organizations must provide all Medicare and Medicaid covered services. The financing of this model is accomplished through prospective capitation of both Medicare and Medicaid payments.

The PACE Medicaid Capitation Rate Setting Guide provides technical assistance to states for their PACE rate setting, and the information to include when submitting rate packages to CMS for review and approval. The guide also includes a template cover sheet to be used by states for their rate package submissions as streamlined submission forms which will improve the efficiency of CMS reviews.

*Form Number:* CMS–10398 #84 (OMB control number: 0938–1148); *Frequency:* Annual and on occasion; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 34; *Total Annual Responses:* 34; *Total Annual Hours:* 17. (For policy questions regarding this collection contact: Angela Cimino at 410–786–2638.)

**William N. Parham, III,**

*Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.*

[FR Doc. 2024–28699 Filed 12–5–24; 8:45 am]

**BILLING CODE 4120–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

**[Document Identifier: CMS–10767]**

### Agency Information Collection Activities: Submission for OMB Review; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments on the collection(s) of information must be received by the OMB desk officer by January 6, 2025.

**ADDRESSES:** Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this