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**

Awardee Lead Profile Assessment (ALPA) (OMB Control No. 0920–1215, Exp. 5/31/2026)—Revision—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

The Centers for Disease Control and Prevention (CDC) is requesting Paperwork Reduction Act (PRA) Clearance for a three-year Revision of an information collection request (ICR) titled Awardee Lead Profile Assessment (ALPA) (OMB Control No. 0920–1215; Exp. 5/31/2026). The goal of this ICR is to build on CDC's existing childhood lead poisoning prevention program. CDC requires that CDC-funded Childhood Lead Poisoning Prevention Programs (CLPPPs), including those funded under the current Notice of Funding Opportunity Childhood Lead Poisoning Prevention and Surveillance of Blood Lead Levels in Children (CDC–RFA–EH21–2102), complete the ALPA annually.

CDC can use this information to inform guidance, resource development, and technical assistance to support eliminating lead exposure in children. Data will be aggregated by jurisdiction, shared with CDC-funded CLPPPs, and used in response to inquiries by the public, press, and Congress. The dissemination of results will support the ability for both funded and non-funded jurisdictions to: (1) identify policies and strategies that support or hinder childhood lead poisoning prevention efforts; and (2) develop and implement similar interventions to improve childhood lead poisoning prevention.

This program management information collection has been revised in several ways, including the addition of new response options and questions as well as simpler language and structure.

- The electronic data collection tool was updated from Epi Info to the REDCap platform to improve functionality and streamline data management.
- The section on local laws was removed to focus solely on programs receiving surveillance funding, ensuring greater relevance and consistency.
- Questions were revised to clarify those that were unclear or difficult to interpret.
- Redundant or overlapping questions and response choices were combined where appropriate for clarity and to reduce respondent burden.
- The previous alphanumeric question labels were replaced with a fully numeric system, creating a cleaner and more organized survey format.

The revisions on the survey will slightly affect the total time burden requested as the time to take the survey has increased from 47 minutes per response in 2021 to 53 minutes per response in 2025. This estimate is based on pilot tests of the revised survey among five respondents and includes the time needed to review the ALPA Training Manual. CDC requests OMB approval for an estimated 66 annual burden hours. 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) |
|---|---|---|---|---|
| State, Tribal, Local, and Territorial Governments | ALPA Web Survey | 75 | 1 | 53/60 |

**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. 2026–08668 Filed 5–4–26; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[30Day–26–0210]**

### 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 ''List of Ingredients Added to Tobacco in the Manufacture of Cigarette Products'' 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 9, 2025 to obtain comments from the public and affected entities. CDC received no public comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected entities' 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**

List of Ingredients Added to Tobacco in the Manufacture of Cigarette Products (OMB Control No. 0920–0210)—Reinstatement—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

Cigarette smoking is the leading preventable cause of premature death and disability in our Nation. Each year more than 480,000 deaths occur as the result of cigarette smoking–related diseases. The Centers for Disease Control and Prevention (CDC), Office on Smoking and Health (OSH) has the primary responsibility for the HHS smoking and health program. Since 1986, as required by the Comprehensive Smoking Education Act of 1984, which amended the Federal Cigarette Labeling and Advertising Act, 15 U.S.C. 1335a, CDC has collected information about the ingredients used in cigarette products. Respondents are commercial cigarette manufacturers, packagers, or importers (or their representatives), who are required by the CSEA to submit ingredient reports to HHS on an annual basis. Respondents are not required to submit specific forms; however, they are required to submit a list of all ingredients used in their products. CDC requires the ingredient report to be submitted by chemical name and Chemical Abstract Service (CAS) Registration Number, consistent with accepted reporting practices for other companies currently required to report ingredients added to other consumer products.

Ingredient reports are due annually on March 31. Information is submitted to CDC by mailing or faxing a written report on the respondent's letterhead. All faxed lists should be followed up with a mailed original. Electronic mail submissions are not accepted. Mail Annual Ingredient Submissions to Attention: FCLAA Program Manager, Office on Smoking and Health, National Center for Chronic Disease Prevention and Health Promotion, Centers for Disease Control and Prevention, 4770 Buford Highway, NE, MS S107–7, Atlanta, GA 30341–3717. Upon receipt and verification of the annual ingredient report, OSH issues a Certificate of Compliance to the respondent. CDC also uses the information to report to Congress (as deemed appropriate) discussing the health effects of these ingredients.

CDC requests OMB approval for a total estimated annualized burden of 358 hours. OMB approval is requested for three years. 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) |
|---|---|---|---|---|
| Business Entities | N/A | 55 | 1 | 6.5 |

**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. 2026–08667 Filed 5–4–26; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[60Day–26–0891; Docket No. CDC–2026–0727]**

### 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 World Trade Center Health Program Enrollment, Appeals & Reimbursement. This data collection is a federal limited benefit health care program providing medical monitoring and treatment benefits to eligible firefighters and related personnel, law enforcement officers, and rescue, recovery, and cleanup workers who responded to the September 11, 2001, attacks in New York City, at the Pentagon, and in Shanksville, Pennsylvania.

**DATES:** CDC must receive written comments on or before July 6, 2026.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC–2026–0727 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,