(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.

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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,

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

World Trade Center Health Program Enrollment, Appeals & Reimbursement (OMB Control No. 0920–0891, Exp. 5/31/2028)—Revision—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The National Institute for Occupational Safety and Health (NIOSH) seeks OMB approval for a revision of an ongoing information collection, World Trade Center Health Program Enrollment, Appeals & Reimbursement (OMB Control No. 0920–0891, Exp. 5/31/2028). In accordance with the James Zadroga 9/11 Health and Compensation Act of 2010, individuals seeking enrollment in the World Trade Center (WTC) Health Program as responders or survivors may apply to the Program. Title I of the Zadroga Act (Pub. L. 111–347, as amended by Pub. L. 114–113 and Pub. L. 116–59), added Title XXXIII to the Public Health Service Act (PHS Act), establishing the World Trade Center (WTC) Health Program within the Department of Health and Human Services (HHS). The Director of NIOSH serves as the Administrator of the WTC Health Program for most purposes, with certain payment functions carried out by the Centers for Medicare & Medicaid Services (CMS).

As established by the Zadroga Act, the WTC Health Program 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 (responders), and to eligible persons who were present in the dust or dust cloud on September 11, 2001, or who worked, resided, or attended school, childcare, or adult daycare in the New York City disaster area (survivors). The WTC Health Program has been authorized for 75 years (through 2090).

In December 2022, Congress amended the Public Health Service Act (42 U.S.C. 300mm–51(c)) to include a Research Cohort for Emerging Health Impacts on Youth (individuals who were 21 years of age or younger on September 11, 2001). This act instructs the WTC Health Program Administrator to establish this research cohort. In this Revision, NIOSH requests OMB approval of revisions to the web-based Youth Research Cohort (YRC) portal, which will be used to engage potential future cohort participants and facilitate Program's efforts to educate potential participants about the YRC. In this Revision, total annualized burden will decrease from 14,332 hours to 11,332 (−3,000 hours). This Revision decreases the amount of time expected for potential YRC participants to provide the requested information by replacing the previous Youth Research Cohort Registration Portal form with a new Youth Research Cohort Information Form. Burden hours are further reduced by eliminating the previous Youth Research Cohort Registration HIPAA Authorization Form which previously asked potential YRC participants to provide protected health information.

CDC requests OMB approval for an estimated 11,332 annual burden hours. There are no additional costs to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number responses per respondent	Average burden per response (in hours)	Total burden hours
FDNY Responder	World Trade Center Health Program FDNY Responder Application for Enrollment.	140	1	30/60	70
General Responder	World Trade Center Health Program Responder Application for Enrollment (Other than FDNY).	6,215	1	30/60	3,108
Pentagon/Shanksville Responder.	World Trade Center Health Program Pentagon/Shanksville Responder Application for Enrollment.	742	1	30/60	371
WTC Survivor	World Trade Center Health Program Survivor Application for Enrollment (all languages).	9,240	1	30/60	4,620
General responder	Clinic Selection Postcard for new general responders in NY/NJ to select a clinic.	3,830	1	15/60	958
Interested Party	Petition for the addition of health conditions	35	1	1	35
Program Applicants or Members.	Designated Representative Appointment Form	1,300	1	15/60	325

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondent	Form name	Number of respondents	Number responses per respondent	Average burden per response (in hours)	Total burden hours
Program Applicants or Members.	Designated Representative HIPAA Release Form.	1,300	1	15/60	325
General Public	WTC Health Program HIPAA Authorization for Deceased Individuals.	30	1	15/60	8
Program Applicants or Members.	WTC Health Program General HIPAA Authorization to Third Parties.	30	1	15/60	8
Program Applicants or Members.	Designated Representative Appointment Form	15	1	15/60	4
Youth Research Cohort Potential Participants.	Youth Research Cohort Information Form	6,000	1	15/60	1,500
Total					11,332

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. 2026–08669 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-26–0910; Docket No. CDC–2026–0728]

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 continuing information collection project titled Message Testing for Tobacco Communication Activities (MTTCA). The MTTCA clearance is designed to collect information about adult smokers' and nonsmokers' attitudes and perceptions towards tobacco, and to pretest draft messages and materials for clarity, salience, appeal, and persuasiveness.

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

ADDRESSES: You may submit comments, identified by Docket No. CDC–2026–0728 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

Message Testing for Tobacco Communication Activities (MTTCA) (OMB Control No. 0920–0910, Exp. 09/30/26)—Extension—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In 2012, CDC's Office on Smoking and Health obtained OMB approval of a Generic Clearance to support the development and testing of tobacco-related health messages, including messages disseminated through multiple phases of a media campaign (Message Testing for Tobacco Communication Activities (MTTCA), OMB Control No. 0920–0910, Exp. 1/31/2015). In 2015, OSH obtained approval