funded 33 state and local public health programs (funded SLHD). These recipients were selected through a competitive objective review process and are managed as CDC cooperative agreements. Awards are for five years and are renewed through an Annual Performance Report (APR)/Continuation Application. The Tracking Program collects data from recipients about their activities and progress for the purposes of program evaluation and monitoring (hereafter referenced as program data).

Environmental public health tracking is the ongoing collection, integration, analysis, and dissemination of health, exposure, and hazard data (hereinafter referenced as Tracking Network data) to inform public health actions that protect the population from harm resulting from exposure to environmental contaminants. The Tracking Network provides data from existing health, exposure, and hazard surveillance systems and supports ongoing efforts within the public health and environmental sectors to improve data collection, accessibility, and dissemination as well as analytic and response capacity. Data that were previously collected for different purposes and stored in separate state and local systems are now available in a nationally standardized format allowing programs to begin bridging the gap between health and the environment.

CDC is requesting approval for an increase of seven additional annual respondents from the 30 approved under the previous ICR and five-year NOFO (No. CDC–RFA–EH17–1702). In spring of 2022, under the new five-year NOFO (No. CDC–RFA–EH22–2202), the CDC's Tracking Program funded 33 state and local public health programs (funded SLHD). CDC is now requesting approval for up to 37 annual respondents. This number reflects the current 33 SLHD respondents plus four to allow for future funding of new SLHD or to collect voluntary responses from unfunded SLHD.

Data from recipients or other SLHD are submitted annually following standardized procedures. Tracking network data submitted annually by recipients and other SLHD to the Tracking Program include seven datasets and the metadata form, specifically (1) birth defects prevalence, (2) childhood blood lead levels, (3) drinking water monitoring, (4) emergency department visits, (5) hospitalizations, (6) radon testing, (7) biomonitoring, and (8) metadata. The Tracking Program will begin using Research Electronic Data Capture (REDCap) for its Electronic Data Capture System (EDCS) needs, which is an easy-to-use, free software tool that is useful for programmatic deliverable management and data capture. Using an EDCS significantly reduces the burden by optimizing the data capture method to eliminate the need for personnel to complete manual data cleaning and organization before using data for analysis and evaluation upon submission.

Based on the above changes, CDC requests OMB approval for an estimated 14,384 annualized burden hours. There is no cost to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
State and Local Health Department	Birth defects prevalence	30	1	40
	Childhood blood lead levels	37	1	40
	Drinking water monitoring	37	1	50
	Emergency department visits	37	1	40
	Hospitalizations	37	1	40
	Radon testing	25	1	50
	Biomonitoring	25	1	40
	Metadata records	37	2	20
	Work Plan Template	37	1	21
	Program Accomplishments and Public Health Actions Report.	37	2	20
	Performance Measures Report	37	1	20
	PHA Impact Follow Up Form	37	2	0.25
	Communications plan	37	1	2
	Web Stats Template	37	2	1

Jeffrey M. Zirger,

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10302]

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 July 27, 2023.

ADDRESSES: Written 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.

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/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786–4669.

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. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Collection Requirements for Compendia for Determination of Medically-accepted Indications for Off-label Uses of Drugs and Biologicals in an Anti-cancer Chemotherapeutic Regimen; *Use:* Section 182(b) of the Medicare Improvement of Patients and Providers Act (MIPPA) amended section 1861(t)(2)(B) of the Social Security Act (42 U.S.C. 1395x(t)(2)(B)) by adding at the end the following new sentence: 'On and after January 1, 2010, no compendia may be included on the list of compendia under this subparagraph unless the compendia has a publicly transparent process for evaluating therapies and for identifying potential conflicts of interest.' We believe that the implementation of this statutory provision that compendia have a ''publicly transparent process for evaluating therapies and for identifying potential conflicts of interests'' is best accomplished by amending 42 CFR 414.930 to include the MIPPA requirements and by defining the key components of publicly transparent processes for evaluating therapies and for identifying potential conflicts of interests.

All currently listed compendia will be required to comply with these provisions, as of January 1, 2010, to remain on the list of recognized compendia. In addition, any compendium that is the subject of a future request for inclusion on the list of recognized compendia will be required to comply with these provisions. No compendium can be on the list if it does not fully meet the standard described in section 1861(t)(2)(B) of the Act, as revised by section 182(b) of the MIPPA. *Form Number:* CMS–10302 (OMB control number: 0938–1078); *Frequency:* Annually; *Affected Public:* Business and other for-profits and Not-for-profit institutions; *Number of Respondents:* 845; *Total Annual Responses:* 900; *Total Annual Hours:* 5,135. (For policy questions regarding this collection contact Sarah Fulton at 410–786–2749.)

Dated: June 22, 2023.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10638]

Agency Information Collection Activities: Proposed Collection; 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 (the 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 60 days for public comment on the proposed action. Interested persons are invited to send 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 August 28, 2023.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. 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: Document Identifier/OMB Control Number: ____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.