adulthood in the healthiest possible way.

CDC will use a web-based system to collect data on the approaches that LEAs are using to meet their goals. Approaches include helping LEAs and priority schools deliver sexual health education emphasizing HIV and other STD prevention; increasing adolescent access to key sexual health services; and establishing safe and supportive environments for students and staff. To track LEA progress and evaluate the effectiveness of program activities, CDC will be collecting data using a mix of process and outcome measures. Process measures to be completed by all LEAs will assess the extent to which planned program activities have been implemented and lead to feasible and sustainable programmatic outcomes. Process measures include items on school health policy and practice assessment and training and technical assistance received from non-governmental partner organizations. Outcome measures, which will be completed by local education agencies, assess whether funded activities at each site are leading to intended outcomes including public health impact of systemic change in schools. These measures drove the development of questionnaires that have been tailored to each of the LEAs' strategies (*i.e.,* SHE, SHS, SSE).

Respondents are 25 LEAs that have been funded under PS18–1807. Local education agencies will complete the questionnaires semi-annually using the Program Evaluation and Reporting System (PERS), an electronic web-based interface specifically designed for this data collection. Each LEA will receive a unique log-in to the system and technical assistance to ensure they can use the system easily. The dates when data are requested reflect the Office of Financial Resources (OFR) deadlines to provide timely feedback to LEAs and CDC staff for accountability and optimal use of funds. CDC anticipates that semi-annual information collection will begin in February 2020 and will describe activities conducted during the period August 2019–July 2022. The estimated burden per response is approximately 2–26 hours. This estimate includes time for local education agencies to gather information at the district and school-levels. Annualizing this collection over three years results in an estimated annualized burden of 1,750 hours per year and 5,250 for three years across all funded local education agencies.

LEAs are required to allocate at least 6% of their NOFO award on evaluation ranging from $15,000 to $21,000. Grantees may use these discretionary funds for collection of process and outcome measures, including time to gather and enter data into the online and evaluation reporting system. There is no cost to the respondents other than their time. The total annual burden hours are 1,750.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
LEA	Funded District Questionnaire	25	2	2
	Priority School Questionnaire	25	2	26
	District Assistance Questionnaire	25	2	7

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Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30-Day–19–1202]

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 Survey of Engineered Nanomaterial Occupational Safety and Health Practices 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 April 23, 2019, 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 or send an email to *omb@cdc.gov.* 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

Survey of Engineered Nanomaterial Occupational Safety and Health Practices (OMB Control No. 0920–1202, Exp. 10/31/2019)—Revision—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

As mandated in the Occupational Safety and Health Act of 1970 (Pub. L. 91–596), the mission of the National

Institute for Occupational Safety and Health (NIOSH) is to conduct research and investigations on work-related disease and injury and to disseminate information for preventing identified workplace hazards (Sections 20(a)(1) and (d)). This dual responsibility recognizes the need to translate research into workplace application if it is to impact worker safety and well-being. The goal of this project is to assess the relevance and impact of NIOSH's contribution to guidelines and risk mitigation practices for safe handling of engineered nanomaterials in the workplace. The intended use of this data is to inform NIOSH's research agenda to enhance its relevance and impact on worker safety and health in the context of engineered nanomaterials.

The research under this project will survey companies who manufacture, distribute, fabricate, formulate, use or provide services related to engineered nanomaterials. The analysis will describe the survey sample, response rates, and types of company by industry and size. Further analysis will focus on identifying the types of engineered nanomaterials being used in industry and the types of occupational safety and health practices being implemented. The analysis will be used to develop a final report which evaluates the influence of NIOSH products, services, and outputs on industry occupational safety and health practices.

Under this project, the following activities and data collections will be conducted:

(1) Company Pre-calls. Sampled companies will be contacted to identify the person who will complete the survey and to ascertain whether or not the company handles engineered nanomaterials.

(2) Survey. A web-based questionnaire, with a mail option, will be administered to companies. The purpose of the survey is to learn directly from companies about their use of NIOSH materials and their occupational safety and health practices concerning engineered nanomaterials.

A sample of 600 companies will be compiled from lists of industry associations, research reports, marketing databases, and web-based searches. Of the 600 selected companies we anticipate that 500 will complete the survey. The company pre-call is expected to require five minutes to complete. The survey is expected to require 20 minutes to complete; including the time it may take respondents to look-up and retrieve needed information. The estimated annualized burden hours for the respondents' time to participate in this information collection is 108 hours. There are no costs to the responders other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Avg. burden per response (in hrs.)
Receptionist	Pre-call	300	1	5/60
Occupational health and safety specialists	Survey	100	1	20/60
Industrial Production Managers	Survey	75	1	20/60
Natural Sciences Managers	Survey	75	1	20/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers CMS–10261, CMS–10556, CMS–R–305, CMS–10328 and CMS–10079]

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

AGENCY: Centers for Medicare & Medicaid 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 October 11, 2019.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–5806 *OR* Email: *OIRA_submission@omb.eop.gov*.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html*.

1. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov*.

2. Call the Reports Clearance Office at (410) 786–1326.

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