**ACTION:** Notice of availability.

**SUMMARY:** NIOSH announces the availability of the final *National Occupational Research Agenda for Public Safety.* 

**DATES:** The final document was published March 6, 2019 on the CDC website.

**ADDRESSES:** The document may be obtained at the following link: *https://www.cdc.gov/nora/councils/pubsaf/* research agenda.html.

### FOR FURTHER INFORMATION CONTACT:

Emily Novicki, M.A., M.P.H, (*NORACoordinator@cdc.gov*), National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, Mailstop E–20, 1600 Clifton Road NE, Atlanta, GA 30329, phone (404) 498–2581 (not a toll free number).

**SUPPLEMENTARY INFORMATION:** On April 12, 2018, NIOSH published a request for public review in the **Federal Register** [83 FR 15838] of the draft version of the *National Occupational Research Agenda for Public Safety.* All comments received were reviewed and addressed where appropriate.

Dated: March 7, 2019.

Frank J. Hearl,

Chief of Staff, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention. [FR Doc. 2019–04460 Filed 3–11–19; 8:45 am]

BILLING CODE 4163-19-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[Docket Number CDC-2019-0018, NIOSH-328]

# Draft—National Occupational Research Agenda for Healthy Work Design and Well-being

**AGENCY:** National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Request for comment.

**SUMMARY:** The National Institute for Occupational Safety and Health of the Centers for Disease Control and Prevention announces the availability of a draft NORA Agenda entitled National Occupational Research Agenda for Healthy Work Design and Well-being for public comment. To view the notice and related materials, visit https:// www.regulations.gov and enter CDC- 2019–0018 in the search field and click "Search."

# Table of Contents

- DATES:
- ADDRESSES:
- FOR FURTHER INFORMATION CONTACT:
- SUPPLEMENTARY INFORMATION:BACKGROUND:

**DATES:** Electronic or written comments must be received by May 13, 2019. **ADDRESSES:** You may submit comments, identified by CDC–2019–0018 and docket number NIOSH–328, by any of the following methods:

• Federal eRulemaking Portal: https://www.regulations.gov Follow the instructions for submitting comments.

• *Mail:* National Institute for Occupational Safety and Health, NIOSH Docket Office, 1090 Tusculum Avenue, MS C-34, Cincinnati, Ohio 45226–1998.

Instructions: All submissions received in response to this notice must include the agency name and docket number [CDC-2019-0018; NIOSH-328]. All relevant comments received will be posted without change to https:// www.regulations.gov, including any personal information provided. For access to the docket to read background documents or comments received, go to https://www.regulations.gov. All information received in response to this notice will also be available for public examination and copying at the NIOSH Docket Office, 1150 Tusculum Avenue, Room 155, Cincinnati, OH 45226–1998.

FOR FURTHER INFORMATION CONTACT: Emily Novicki, M.A., M.P.H., (*NORACoordinator@cdc.gov*), National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, Mailstop E–20, 1600 Clifton Road NE, Atlanta, GA 30329, phone (404) 498–2581 (not a toll free number).

**SUPPLEMENTARY INFORMATION:** The National Occupational Research Agenda (NORA) is a partnership program created to stimulate innovative research and improved workplace practices. The national agenda is developed and implemented through the NORA sector and cross-sector councils. Each council develops and maintains an agenda for its sector or cross-sector.

Background: The National Occupational Research Agenda for Healthy Work Design and Well-being is intended to identify the research, information, and actions most urgently needed to prevent occupational injuries. This research agenda provides a vehicle for stakeholders to describe the most relevant issues, gaps, and safety and health needs for the sector. Each NORA research agenda is meant to guide or promote high priority research efforts on a national level, conducted by various entities, including: Government, higher education, and the private sector.

This is the first Healthy Work Design and Well-Being Agenda, developed for the third decade of NORA (2016–2026). It was developed considering new information about injuries and illnesses, the state of the science, and the probability that new information and approaches will make a difference. As the steward of the NORA process, NIOSH invites comments on the draft National Occupational Research Agenda for Healthy Work Design and Well-Being. Comments expressing support or with specific recommendations to improve the agenda are requested. A copy of the draft agenda is available at https:// www.regulations.gov (see Docket Number CDC-2019-0018).

Dated: March 7, 2019.

#### Frank J. Hearl,

Chief of Staff, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention. [FR Doc. 2019–04498 Filed 3–11–19; 8:45 am]

BILLING CODE 4163-19-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier CMS-10325]

### 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 April 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 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: Disclosure and **Recordkeeping Requirements for** Grandfathered Health Plans under the Affordable Care Act; Use: Section 1251 of the Affordable Care Act provides that certain plans and health insurance coverage in existence as of March 23, 2010, known as grandfathered health plans, are not required to comply with certain statutory provisions in the Act. The final regulations titled "Final Rules under the Affordable Care Act for Grandfathered Plans, Preexisting Condition Exclusions, Lifetime and Annual Limits, Rescissions, Dependent Coverage, Appeals, and Patient Protections" (80 FR 72192, November 18, 2015) require that, to maintain its status as a grandfathered health plan, a plan must maintain records documenting the terms of the plan in effect on March 23, 2010, and any other documents that are necessary to verify, explain or clarify status as a grandfathered health plan. The plan must make such records available for examination upon request by participants, beneficiaries, individual policy subscribers, or a state or federal agency official.

A grandfathered health plan is also required to include a statement in any summary of benefits under the plan or health insurance coverage, that the plan or coverage believes it is a grandfathered health plan within the meaning of section 1251 of the Affordable Care Act. and providing contact information for participants to direct questions and complaints. In addition, a grandfathered group health plan that is changing health insurance issuers is required to provide the succeeding health insurance issuer (and the succeeding health insurance issuer must require) documentation of plan terms (including benefits, cost sharing, employer contributions, and annual limits) under the prior health insurance coverage sufficient to make a determination whether the standards of paragraph §147.140(g)(1) of the final regulations are exceeded. It is also required that, for an insured group health plan (or a multiemployer plan) that is a grandfathered plan, the relevant policies, certificates, or contracts of insurance, or plan documents must disclose in a prominent and effective manner that employers, employee organizations, or plan sponsors, as applicable, are required to notify the issuer (or multiemployer plan) if the contribution rate changes at any point during the plan year. Form Number: CMS-10325 (OMB Control Number: 0938–1093; Frequency: Occasionally; Affected Public: Private Sector, State,

Local or Tribal governments; *Number of Respondents:* 20,973; *Total Annual Responses:* 3,831,484; *Total Annual Hours:* 114. (For policy questions regarding this collection contact Usree Bandyopadhyay at 410–786–6650.)

Dated: March 7, 2019.

# William N. Parham, III,

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

[FR Doc. 2019–04494 Filed 3–11–19; 8:45 am] BILLING CODE 4120–01–P

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10052 and CMS-10629]

# Agency Information Collection Activities: Proposed Collection; 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 (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 May 13, 2019.

**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://*