contract file and become a matter of record. FedBizOpps is an electronic method for publicizing contract opportunities. The number of presolicitation notices issued government-wide in the last 365 days (as of August 11, 2017) per FedBizOpps was 7952.

We estimate that three respondents on average would reply to a presolicitation notice. Based on this info, we feel that the estimated respondents would number 23,856. We also estimate that each respondent on average would reply to three presolicitation notices a year. Time required to read and prepare information is estimated at 5 minutes per completion.

The burden has increased from the one in **Federal Register** Notice 79 FR 49316 dated August 20, 2014 due to the additional number of presolicitation notices issued by the Federal Government in FedBizOpps. This had the effect of expanding the number of respondents who would be affected by the collection.

B. Annual Reporting Burden

Respondents: 16,699.

Responses per Respondent: 3.

Annual Responses: 50,097.

Hours per Response: .08.

Total Burden Hours: 4,008.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the FAR, and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0037, Presolicitation Notice and Response, in all correspondence.

Dated: August 16, 2017.

Lorin S. Curit,

Director, Federal Acquisition Policy Division, Office of Government-wide Acquisition Policy, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2017–17830 Filed 8–22–17; 8:45 am]

BILLING CODE 6820–EP–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0250; Docket No. 2017–0001; Sequence 7]

Information Collection; General Services Administration Acquisition Regulation; Zero Burden Information Collection Reports

AGENCY: Office of the Chief Acquisition Officer, General Services Administration (GSA).

ACTION: Notice of request for public comments regarding an extension to an existing OMB information collection.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division (MVCB) will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement regarding Zero Burden Information Collection Reports.

DATES: Submit comments on or before: October 23, 2017.

ADDRESSES: Submit comments identified by Information Collection 3090–0250, Zero Burden Information Collection Reports, by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by searching the OMB control number 3090–0250. Select the link ''Comment Now'' that corresponds with ''Information Collection 3090–0250, Zero Burden Information Collection Reports''. Follow the instructions provided on the screen. Please include your name, company name (if any), and ''Information Collection 3090–0250, Zero Burden Information Collection Reports'' on your attached document.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Sosa/IC 3090–0250, Zero Burden Information Collection Reports.

Instructions: Please submit comments only and cite Information Collection 3090–0250, Zero Burden Information Collection Reports, in all correspondence related to this collection. Comments received generally will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two to three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

FOR FURTHER INFORMATION CONTACT: Ms. Dana Munson, Procurement Analyst, General Services Acquisition Policy, at telephone 202–357–9652 or via email to *dana.munson@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

This information requirement consists of reports that do not impose collection burdens upon the public. These collections require information which is already available to the public at large or that is routinely exchanged by firms during the normal course of business. A general control number for these collections decreases the amount of paperwork generated by the approval process.

GSA has a published rule in the **Federal Register** that falls under information collection 3090–0250. The rule that prescribed clause 552.238–70 ''Identification of Electronic Office Equipment Providing Accessibility for the Handicapped'' was published at 56 FR 29442, June 27, 1991, titled ''Implementation of Public Law 99–506'', with an effective date of July 8, 1991.

B. Annual Reporting Burden

None.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate and based on valid assumptions and methodology; and ways to enhance the quality, utility, and clarity of the information to be collected.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 3090–0250, Zero

Burden Information Collection Reports, in all correspondence.

Jeffrey A. Koses,

Director, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2017–17831 Filed 8–22–17; 8:45 am]

BILLING CODE 6820–61–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2017–0072, NIOSH–300]

Draft—National Occupational Research Agenda for Manufacturing

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

ACTION: Request for comments.

SUMMARY: As steward of the National Occupational Research Agenda (NORA), the National Institute for Occupational Safety and Health of the Centers for Disease Control and Prevention announces the availability of the draft National Occupational Research Agenda for Manufacturing for public comment. Written by the NORA Manufacturing Sector Council, the Agenda identifies the most important occupational safety and health research needs for the next decade, 2016–2026. A copy of the draft Agenda is available at *http://www.regulations.gov* (search Docket Number CDC–2017–0072).

DATES: Electronic or written comments must be received by October 23, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2017–0072 and docket number NIOSH–300, by any of the following methods:

- *Federal eRulemaking Portal: http://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 must include the agency name and Docket Number [CDC–2017–0072; NIOSH–300]. All relevant comments received will be posted without change to *http://www.regulations.gov,* including any personal information provided. For access to the docket to read background documents or comments received, go to *http://www.regulations.gov.*

FOR FURTHER INFORMATION CONTACT: Emily Novicki (*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.

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.

The National Occupational Research Agenda for Manufacturing is intended to identify the research, information, and actions most urgently needed to prevent occupational injuries and illnesses in the manufacturing sector. The National Occupational Research Agenda for Manufacturing provides a vehicle for industry 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. The first National Occupational Research Agenda for Manufacturing was published in 2010 for the second decade of NORA (2006–2016). This draft is an updated agenda for the third decade of NORA (2016–2026). The revised agenda 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 Manufacturing. A copy of the draft Agenda is available at *http://www.regulations.gov* (see Docket Number CDC–2017–0072, NIOSH–300).

Dated: August 17, 2017.

Frank Hearl,

Chief of Staff, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2017–17786 Filed 8–22–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–4835]

Peripheral and Central Nervous System Drugs Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Peripheral and Central Nervous System Drugs Advisory Committee. The general function of the committee is to provide advice and recommendations to the Agency on FDA's regulatory issues. The meeting will be open to the public. FDA is establishing a docket for public comment on this document.

DATES: The public meeting will be held on September 28, 2017, from 9 a.m. to 4:30 p.m.

ADDRESSES: Tommy Douglas Conference Center, The Ballroom, 10000 New Hampshire Ave., Silver Spring, MD 20903. Answers to commonly asked questions about FDA Advisory Committee meetings may be accessed at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.* Information about the Tommy Douglas Conference Center may be accessed at: *http://www.tommydouglascenter.com/.*

FDA is establishing a docket for public comment on this meeting. The docket number is FDA–2017–N–4835. The docket will close on September 27, 2017. Submit either electronic or written comments on this public meeting by September 27, 2017. Late, untimely filed comments will not be considered. Electronic comments must be submitted on or before September 27, 2017. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of September 27, 2017. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Comments received on or before September 14, 2017, will be provided to the committee. Comments received after that date will be taken into consideration by the Agency.