Office Building, Docket Library, Room 10102, 725 17th Street NW., Washington, DC 20503. Comments sent to OMB by U.S. postal mail, however, are subject to delays due to heightened security precautions. Thus, comments instead should be sent by facsimile to (202) 395–5167.

David C. Shonka,

Acting General Counsel.

[FR Doc. 2017–04929 Filed 3–13–17; 8:45 am]

BILLING CODE 6750–01–P

GULF COAST ECOSYSTEM RESTORATION COUNCIL

Notice of Proposed Subaward Under a Council-Selected Restoration Component Award

AGENCY: Gulf Coast Ecosystem Restoration Council.

ACTION: Notice.

SUMMARY: The Gulf Coast Ecosystem Restoration Council (Council) publishes notice of a proposed subaward from the National Oceanic and Atmospheric Administration National Centers for Coastal Ocean Science (NOAA) to the Gulf of Mexico Alliance (GOMA), a nonprofit organization, for the purpose of supporting the Council Monitoring and Assessment Program (CMAP). The Council and NOAA have entered an interagency agreement for NOAA to carry out CMAP, as approved in the Council's Initial Funded Priorities List.

FOR FURTHER INFORMATION CONTACT: Please send questions by email to *raams_pgmsupport@restorethegulf.gov.*

SUPPLEMENTARY INFORMATION: Section 1321(t)(2)(E)(ii)(III) of the RESTORE Act (33 U.S.C. 1321(t) and *note*) and Treasury's implementing regulation at 31 CFR 34.401(b) require that, for purposes of awards made under the Council-Selected Restoration Component, a State or Federal award recipient may make a grant or subaward to or enter into a cooperative agreement with a nongovernmental entity that equals or exceeds 10 percent of the total amount of the award provided to the State or Federal award recipient only if certain notice requirements are met. Specifically, at least 30 days before the State or Federal award recipient enters into such an agreement, the Council must publish in the **Federal Register** and deliver to specified Congressional Committees the name of the recipient and subrecipient; a brief description of the activity, including its purpose; and the amount of the award. This notice accomplishes the **Federal Register** publication requirement.

Description of Proposed Action

As specified in the Initial Funded Priorities List, which is available on the Council's Web site at *https://www.restorethegulf.gov/council-selected-restoration-component/funded-priorities-list,* RESTORE Act funds will support the Council Monitoring and Assessment Program (CMAP). Administered jointly by NOAA and the Department of the Interior's United States Geological Survey (USGS), CMAP will build the foundational components for Gulf region-wide monitoring in order to measure impacts of investments in restoration. Through collaboration with the Gulf States, federal and local partners, academia, non-governmental/non-profit organizations, and industry, the program will use a Monitoring Community of Practice coordinated by the Gulf of Mexico Alliance to leverage existing resources, capacities, and expertise and build on existing monitoring programs. These existing programs will be coordinated into a network, to provide efficiency in monitoring and collaborative cross-program review of performance with other Gulf ecosystem recovery efforts.

The program will: (1) Create an inventory of the existing monitoring programs, data, protocols and standards; (2) determine the minimum monitoring elements needed to evaluate the performance of restoration projects; (3) evaluate monitoring program suitability; (4) combine data from the suitable existing programs into searchable databases for Council use; (5) examine the inventory to determine what data are missing (*i.e.* identify information gaps) that would be required for the RESTORE Council; (6) document existing baseline assessments of habitat and water quality conditions; and (7) provide recommendations to the Council to supplement and refine the existing monitoring programs to fill-in the information gaps where possible.

Will D. Spoon,

Program Analyst, Gulf Coast Ecosystem Restoration Council.

[FR Doc. 2017–04937 Filed 3–13–17; 8:45 am]

BILLING CODE 6560–58–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2017–0008]

Proposed Data Collection Submitted for Public Comment and Recommendations: Survey of Engineered Nanomaterial Occupational Safety and Health Practices; Extension of Public Comment Period

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Extension of public comment period.

SUMMARY: On February 10, 2017, the Centers for Disease Control and Prevention (CDC), located within the Department of Health and Human Services (HHS) published a notice in the **Federal Register** requesting public comment on the proposed information collection entitled ''Survey of Engineered Nanomaterial Occupational Safety and Health Practices''. Written and electronic comments were to be received on or before April 11, 2017. Because of an improper docket opening, CDC is extending the comment period to allow the public a full 60 days to provide comment on this docket. In consideration of this public access issue, HHS/CDC is extending the comment period to May 12, 2017.

DATES: Written comments must be received on or before May 11, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2017–0008 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov*. Follow the instructions for submitting comments.
- *Mail:* Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *Regulations.gov*, including any personal information provided. For access to the docket to read background documents or comments received, go to *Regulations.gov*.

Please note: All public comment should be submitted through the Federal eRulemaking portal (*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 the Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329; phone: 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 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 OMB for approval.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) The accuracy of the agency's estimate of the burden of the proposed collection of information; (c) Ways to enhance the quality, utility, and clarity of the information to be collected; (d) Ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology; and (e) Estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information.

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2017–04942 Filed 3–13–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–1062]

Joint Meeting of the Anesthetic and Analgesic Drug Products Advisory Committee and the Drug Safety and Risk Management 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 Anesthetic and Analgesic Drug Products Advisory Committee and the Drug Safety and Risk Management Advisory Committee. The general function of the committees is to provide advice and recommendations to the Agency on FDA's regulatory issues. At least one portion of the meeting will be closed to the public. FDA is establishing a docket for public comment on this document.

DATES: The meeting will be held on April 5, 2017, from 8 a.m. to 5 p.m.

ADDRESSES: Tommy Douglas Conference Center, the Ballroom, 10000 New Hampshire Ave., Silver Spring, MD 20903. The conference center's telephone number is 240–645–4000. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.* You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2017–N–1062 for ''Joint Meeting of the Anesthetic and Analgesic Drug Products Advisory Committee and the Drug Safety and Risk Management Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments.'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The