## FEDERAL COMMUNICATIONS COMMISSION

**[OMB 3060–0392]**

### Information Collection Being Reviewed by the Federal Communications Commission Under Delegated Authority

**AGENCY:** Federal Communications Commission.

**ACTION:** Notice and request for comments.

**SUMMARY:** As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act (PRA) of 1995 (44 U.S.C. 3501–3520), the Federal Communications Commission (FCC or Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collections. Comments are requested concerning: Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees. The FCC may not conduct or sponsor a collection of information unless it displays a currently valid OMB control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid OMB control number.

**DATES:** Written PRA comments should be submitted on or before March 7, 2016. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

**ADDRESSES:** Direct all PRA comments to Nicole Ongele, FCC, via email *PRA@fcc.gov* and to *Nicole.Ongele@fcc.gov.*

**FOR FURTHER INFORMATION CONTACT:** For additional information about the information collection, contact Nicole Ongele at (202) 418–2991.

**SUPPLEMENTARY INFORMATION:**

*OMB Control Number:* 3060–0392.

*Title:* 47 CFR 1 Subpart J—Pole Attachment Complaint Procedures.

*Form Number:* N/A.

*Type of Review:* Extension of currently approved collection.

*Respondents:* Businesses or other for-profit, and State, local or tribal government.

*Number of Respondents and Responses:* 1,772 respondents; 1,772 responses.

*Estimated Time per Response:* 0.5 to 100 hours.

*Frequency of Response:* On occasion reporting requirement and third party disclosure requirement.

*Obligation to Respond:* Required to obtain or retain benefits. Statutory authority for this information collection is contained in 47 U.S.C. 224.

*Total Annual Burden:* 2,629 hours.

*Total Annual Cost:* $450,000.

*Privacy Act Impact Assessment:* No impact(s).

*Nature and Extent of Confidentiality:* There is no need for confidentiality. However, respondents may request materials or information submitted to the Commission be withheld from public inspection under 47 CFR 0.459 of FCC rules.

*Needs and Uses:* The rules and regulations contained in 47 CFR part 1 Subpart J provide complaint and enforcement procedures to ensure that telecommunications carriers and cable system operators have nondiscriminatory access to utility poles, ducts, conduits, and rights-of-way on rates, terms and conditions that are just and reasonable. They also provide complaint and enforcement procedures for incumbent local exchange carriers (as defined in 47 U.S.C. 251(h)) to ensure that the rates, terms, and conditions of their access to pole attachments are just and reasonable. The FCC will use the information collected under these rules to hear and resolve petitions for stay and complaints as mandated by Section 224 of the Communications Act of 1934, as amended. The information that is also filed is used to determine the merits of the petitions and complaints. Additionally, state certifications are used to make public notice of the states' authority to regulate rates, terms and conditions for pole attachments, and to determine the scope of the FCC's jurisdiction.

Federal Communications Commission.

**Sheryl D. Todd,**

*Deputy Secretary. Office of the Secretary.*

[FR Doc. 2015–33240 Filed 1–5–16; 8:45 am]

**BILLING CODE 6712–01–P**

## GENERAL SERVICES ADMINISTRATION

**[Notice–CSE–2016–01; Docket No. 2016–0002; Sequence No. 1]**

### Notice of the General Services Administration's Labor-Management Relations Council Meeting

**AGENCY:** Office of Human Resources Management (OHRM), General Services Administration (GSA).

**ACTION:** Notice of meeting.

**SUMMARY:** The General Services Administration's Labor-Management Relations Council (GLMRC), a Federal Advisory Committee established in accordance with the Federal Advisory Committee Act (FACA), 5 U.S.C., App., and Executive Order 13522, plans to hold one meeting that is open to the public.

**DATES:** The meeting will be held on Tuesday, January 26, 2016 from 9:30 a.m. to 4:30 p.m. and Wednesday, January 27, 2016 from 9:30 a.m. to 12:00 noon, Eastern Standard Time.

**ADDRESSES:** The meeting will be held in Room 6044 of the General Services Administration's Headquarters Building, 1800 F Street NW., Washington, DC 20405. This site is accessible to individuals with disabilities.

**FOR FURTHER INFORMATION CONTACT:** Ms. Paula D. Lucak, GLMRC Designated Federal Officer (DFO), OHRM, General Services Administration, at telephone 202–969–7110, or email at *gmlrc@gsa.gov.*

**SUPPLEMENTARY INFORMATION:**

#### Background

The GLMRC is a forum for managers and the exclusive representatives of the U.S. General Services Administration (GSA) employees, which are the two national labor unions. In this forum, managers and the Unions discuss Government operations to promote satisfactory labor relations and improve the productivity and effectiveness of GSA. The GLMRC serves as a complement to the existing collective bargaining process and allows managers and the Unions to collaborate in continuing to deliver the highest quality services to the public. The Council discusses workplace challenges and problems and recommends solutions that foster a more productive and cost-effective service to the taxpayer, through improving job satisfaction and employees' working conditions.

**Agenda**

The purpose of the meeting is for the GLMRC to discuss the Council's focus for the upcoming year and consider Agency initiatives. The topics to be discussed include Council metrics & GSA EVS results, GSA EEO program, and Council subcommittee updates.

**Meeting Access**

The meeting is open to the public. The meeting will be held in Room 6044 of the General Services Administration's Headquarters Building, 1800 F Street NW., Washington, DC 20405. This site is accessible to individuals with disabilities. In order to gain entry into the Federal building where the meeting is being held, public attendees who are Federal employees should bring their Federal employee identification cards, and members of the general public should bring their driver's license or other government-issued identification.

**Availability of Materials for the Meeting**

Please see the GLRMC Web site: *http://www.gsa.gov/portal/content/225831* for any materials available in advance of the meeting and for meeting minutes that will be made available after the meeting. Detailed meeting minutes will be posted within 90 days of the meeting.

**Procedures for Providing Public Comments**

The public is invited to submit written comments for the meeting until 5:00 p.m. Eastern Time on the Monday prior to the meeting, by either of the following methods:

*Electronic or Paper Statements:* Submit electronic statements to Ms. Paula Lucak, Designated Federal Officer, at *paula.lucak@gsa.gov;* or send paper statements in triplicate to Ms. Lucak at 1800 F Street NW., Suite 7003A, Washington, DC 20405. In general, public comments will be posted on the GLMRC Web site. All comments, including attachments and other supporting materials received, are part of the public record and subject to public disclosure.

Any comments submitted in connection with the GLMRC meeting will be made available to the public under the provisions of the Federal Advisory Committee Act.

Dated: December 30, 2015.

**Wade Hannum,**

*Office of Human Resources Management, OHRM Director, Office of HR Strategy and Services, Center for Talent Engagement (COE4), General Services Administration.*

[FR Doc. 2015–33302 Filed 1–5–16; 8:45 am]

**BILLING CODE 6820–34–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2015–N–0001]**

### Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee; Amendment of Notice

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing an amendment to the notice of meeting of the Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee. This meeting was announced in the **Federal Register** of November 23, 2015. The amendment is being made to reflect a change in the *Agenda* portion of the document. There are no other changes.

**FOR FURTHER INFORMATION CONTACT:** Sara Anderson, Center for Devices and Radiological Health, Food and Drug Administration, Bldg. 66, Rm. 1643, 10903 New Hampshire Ave., Silver Spring, MD 20993, *Sara.Anderson@fda.hhs.gov,* 301–796–7047, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). Please call the Information Line for up-to-date information on this meeting.

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of November 23, 2015, 80 FR 72971, FDA announced that a meeting of the Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee would be held on February 19, 2016. On page 72972, in the first column, the *Agenda* portion of the document is changed to read as follows:

The Committee will discuss, make recommendations, and vote on information regarding the premarket application (PMA) for the DIAM Spinal Stabilization System, sponsored by Medtronic Sofamor Danek USA. The DIAM Spinal Stabilization System is indicated for skeletally mature patients that have moderate low back pain (with or without radicular pain) with current episode lasting less than 1 year in duration secondary to lumbar degenerative disc disease (DDD) at a single symptomatic level from L2–L5. DDD is confirmed radiologically with one or more of the following factors: (1) Patients must have greater than 2 mm of decreased disc height compared to the adjacent level; (2) scarring/thickening of the ligamentum flavum, annulus fibrosis, or facet joint capsule; or (3) herniated nucleus pulposus. The DIAM device is implanted via a minimally invasive posterior approach.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link.

This notice is issued under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to the advisory committees.

Dated: December 30, 2015.

**Leslie Kux,**

*Associate Commissioner for Policy.*

[FR Doc. 2015–33262 Filed 1–5–16; 8:45 am]

**BILLING CODE 4164–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2015–N–4952]**

### Food and Drug Administration Safety and Innovation Act 907 Public Meeting: Progress on Enhancing the Collection, Analysis, and Availability of Demographic Subgroup Data; Request for Comments

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice of public meeting; request for comments.

**SUMMARY:** The Food and Drug Administration's (FDA or Agency) Office of Minority Health (OMH), Office of Women's Health (OWH), the Center for Biologics Evaluation and Research (CBER), the Center for Drug Evaluation and Research (CDER), and the Center for Devices and Radiological Health (CDRH)