

expansion of the San Ysidro LPOE in three independent phases to improve overall capacity and operational efficiency at the LPOE. The San Ysidro LPOE is located along Interstate 5 (I-5) at the U.S.-Mexico border in the San Ysidro community of the City of San Diego, California.

GSA is proposing the following changes to the Approved Project: A redesign of the proposed pedestrian plaza on the east side of the LPOE. The pedestrian plaza would be expanded to the north to include an additional parcel adjacent to the LPOE. GSA proposes acquisition of the adjacent 0.24-acre parcel to the north that contains two commercial buildings and incorporation of this parcel (Additional Land Area) into the pedestrian plaza. In addition to these proposed changes to the Approved Project, the Revised Project also includes the other components of the Approved Project that have not changed.

The changed circumstances associated with the Approved Project include new information regarding the condition of existing structures adjacent to the LPOE that affect the ability of GSA to implement the Approved Project. The Approved Project anticipated that construction of the pedestrian plaza would require demolition of the existing Milo Building within the LPOE. During final design, it was discovered that two existing buildings adjacent to the Milo Building on the Additional Land Area would likely collapse when the Milo Building is removed. The condition of these adjacent buildings was not known at the time the 2009 Final EIS or 2014 Final SEIS were prepared and this changed circumstance has bearing on the ability to implement the Approved Project.

Due to the changed circumstances and changes to the Approved Project, GSA made the decision to prepare an SEIS for the Revised Project. The purpose of the Revised Project is the same as the Approved Project that was identified in the 2009 Final EIS and 2014 Final SEIS. The purpose of the Revised Project is to improve operational efficiency, security, and safety for cross-border travelers and federal agencies at the San Ysidro LPOE.

The Draft SEIS analyzes two alternatives of the Revised Project, as well as the No Action Alternative. Both of the Action Alternatives include the proposed modifications described above, as well as the other improvements originally proposed as part of the Approved Project.

Alternative 1 would include demolition of the two existing buildings within the Additional Land Area that would be added to the LPOE and incorporated

into the pedestrian plaza. Alternative 2 would involve renovation/adaptive reuse of the existing buildings on the Additional Land Area that would be added to the LPOE and incorporated into the design of the pedestrian plaza and LPOE. Under the No Action Alternative, GSA would continue to implement the Approved Project except that the Milo Building would not be demolished.

#### Public Meeting

The public meeting will be conducted in open house format, where project information will be presented and distributed. Comments must be received by November xx, 2018, and emailed to [osmah.kadri@gsa.gov](mailto:osmah.kadri@gsa.gov), or sent to the address listed above.

Dated: September 12, 2018.

**Matthew Jear**,

*Director, Portfolio Management Division, Pacific Rim Region, Public Buildings Service.*

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#### GENERAL SERVICES ADMINISTRATION

[Notice-CX-2018-01; Docket No. 2018-0002; Sequence No. 24]

#### Office of Human Resources Management; SES Performance Review Board

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

**ACTION:** Notice.

**SUMMARY:** Notice is hereby given of the appointment of new members to the General Services Administration Senior Executive Service Performance Review Board. The Performance Review Board assures consistency, stability, and objectivity in the performance appraisal process.

**DATES:** These appointments are effective September 24, 2018.

**FOR FURTHER INFORMATION CONTACT:** Ms. Shonna James, Director, Executive Resources Division, Office of Human Resources Management, General Services Administration, 1800 F Street NW, Washington, DC 20405, 202-230-7005.

**SUPPLEMENTARY INFORMATION:** Section 4314(c)(1) through (5) of title 5 U.S.C requires each agency to establish, in accordance with regulation prescribed by the Office of Personnel Management, one or more SES performance review board(s). The board is responsible for making recommendations to the appointing and awarding authority on

the performance appraisal ratings and performance awards for the Senior Executive Service employees.

The following have been designated as members of the Performance Review Board of GSA:

- Allison Brigati, Deputy Administrator, Office of the Administrator—Chair.
- Giancarlo Brizzi, Regional Commissioner, Public Buildings Service, Greater Southwest Region.
- Mary Davie, Deputy Commissioner, Federal Acquisition Service.
- Michael Gelber, Deputy Commissioner, Public Buildings Service.
- Antonia Harris, Chief Human Capital Officer, Office of Human Resources Management.
- Tiffany Hixon, Regional Commissioner, Federal Acquisition Service, Northwest, Arctic Region.
- Jack St. John, General Counsel.
- Alan Thomas, Commissioner, Federal Acquisition Service.

Dated: September 18, 2018.

**Emily W. Murphy**,

*Administrator, General Services Administration.*

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#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

##### Food and Drug Administration

[Docket No. FDA-2018-N-3405]

##### Patient Engagement Advisory Committee; Notice of Meeting

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing a forthcoming public advisory committee meeting of the Patient Engagement Advisory Committee (the Committee). The general function of the Committee is to provide advice to the Commissioner, or designee, on complex issues relating to medical devices, the regulation of devices, and their use by patients. The meeting will be open to the public.

**DATES:** The meeting will be held on November 15, 2018, from 8 a.m. to 5 p.m.

**ADDRESSES:** Hilton Washington DC North/Gaithersburg, Grand Ballroom, 620 Perry Pkwy., Gaithersburg, MD 20877. The hotel's telephone number is 301-977-8900; additional information available online at: <https://>

[www3.hilton.com/en/hotels/maryland/hilton-washington-dc-north-gaithersburg-GAIGHHF/index.html](http://www3.hilton.com/en/hotels/maryland/hilton-washington-dc-north-gaithersburg-GAIGHHF/index.html).

Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: <https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm>.

**FOR FURTHER INFORMATION CONTACT:**

Letise Williams, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5441, Silver Spring, MD 20993-0002, [letise.williams@fda.hhs.gov](mailto:letise.williams@fda.hhs.gov), 301-796-8398, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's website at <https://www.fda.gov/AdvisoryCommittees/default.htm> and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

**SUPPLEMENTARY INFORMATION: Agenda:**

On November 15, 2018, the Committee will discuss and make recommendations on the topic "Connected and Empowered Patients: E-Platforms Potentially Expanding the Definition of Scientific Evidence." The recommendations will address how FDA can leverage patient-driven platforms, such as social media and registries, to better engage patients and consumers as empowered partners in the work of protecting public health and promoting responsible innovation. Social media and other web platform enablers are facilitating the growth of virtual patient communities. Increasingly, patients and health care consumers are using these platforms to share their health experiences and seek information from other patients and consumers, rather than their health care providers alone. Novel approaches and methodologies are being used to tap into some of these platforms as potentially rich sources of patient-generated health data, which could be used as relevant and reliable real-world evidence (<https://www.fda.gov/ucm/groups/fdagov-public/@fdagov-meddev-gen/documents/document/ucm513027.pdf>).

This meeting will help advance FDA's objective to assure the needs, experiences, and perspectives of

patients are included as part of FDA's deliberations involving the regulation of medical devices and their use by patients. For this meeting, FDA is seeking input from the Committee and the public on whether and how FDA can harness the emerging potential of these patient platforms to better engage patients and consumers as empowered partners in the work of protecting public health and promoting responsible innovation. In addition, FDA is seeking recommendations from the Committee on ways to leverage these platforms to disseminate as well as potentially collect and evaluate health information to and from patients and consumers.

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 website 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 website after the meeting. Background material is available at <https://www.fda.gov/AdvisoryCommittees/Calendar/default.htm>. Scroll down to the appropriate advisory committee-meeting link.

*Procedure:* Interested persons may present data, information, or views, orally or in writing, on issues pending before the Committee. Oral presentations from the public will be scheduled between approximately 11:15 a.m. to 12:15 p.m. on November 15, 2018. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before October 15, 2018. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by October 17, 2018. Individuals who do not wish to speak at the open public hearing session but would like their comments to be heard by the Committee may send written submissions to the contact person on or before October 23, 2018.

FDA welcomes the attendance of the public at its advisory committee

meetings and will make every effort to accommodate persons with disabilities. If you require accommodations due to a disability, please contact AnnMarie Williams at [Annmarie.Williams@fda.hhs.gov](mailto:Annmarie.Williams@fda.hhs.gov), or 301-796-5966 at least 7 days in advance of the meeting.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our website at <https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm11462.htm> for procedures on public conduct during advisory committee meetings. Please be advised that, for the round table portion of the meeting, FDA will prepare a summary of the discussion in lieu of detailed transcripts.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: September 18, 2018.

**Leslie Kux,**

*Associate Commissioner for Policy.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Health Resources and Services Administration**

**Agency Information Collection Activities: Proposed Collection: Public Comment Request; Information Collection Request Title: National Health Service Corps Scholar/Students to Service Travel Worksheet, OMB No. 0915-0278—Extension**

**AGENCY:** Health Resources and Services Administration (HRSA), Department of Health and Human Services.

**ACTION:** Notice.

**SUMMARY:** In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR. **DATES:** Comments on this ICR should be received no later than November 23, 2018.

**ADDRESSES:** Submit your comments to [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or mail the HRSA