• What factors are important to take into account when conducting risk assessments of safety and quality issues that occur with marketed medical devices? What principles best guide the risk assessment process to assure timely, consistent, and optimal results?

• Are there improvements that FDA and stakeholders could make to enhance risk assessments in recall and shortage situations with medical devices?

• Are there specific activities or issues related to postmarket quality, safety, or compliance activities where approaches used by FDA and industry currently differ enough to create confusion or delay or limit appropriate public health actions? Please identify them.

• In which activities and areas of postmarket quality, compliance, and safety would more detailed policies or guidance be most useful?

At this public workshop, participants will engage in open dialogue to discuss the responses to issues raised by the presenters and the questions in this **Federal Register** notice.

III. Reference

The following reference has been placed on display in the Division of Dockets Management (see **ADDRESSES**) and may be seen by interested persons between 9 a.m. and 4 p.m., Monday through Friday. We have verified all Web site addresses, but we are not responsible for subsequent changes to the Web sites after this document publishes in the **Federal Register.**

1. FDA, ''Quality System (QS) Regulation/Medical Device Good Manufacturing Practices,'' 2014, available at *http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/PostmarketRequirements/QualitySystemsRegulations/default.htm.*

Dated: March 13, 2015.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2015–06278 Filed 3–18–15; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Cancellation of Meeting

Notice is hereby given of the cancellation of the Center for Scientific Review Special Emphasis Panel, April 2, 2015, 1:00 p.m. to April 2, 2015, 2:00 p.m., National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD, 20892 which was published in the **Federal Register** on March 6, 2015, 80 FR 12185.

The meeting has been cancelled due to the reassignment of applications.

Dated: March 13, 2015.

Anna Snouffer,

Deputy Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2015–06274 Filed 3–18–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

[Docket ID: FEMA–2014–0033; OMB No. 1660–0132]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Federal Emergency Management Agency, DHS.

ACTION: Notice.

SUMMARY: The Federal Emergency Management Agency (FEMA) will submit the information collection abstracted below to the Office of Management and Budget for review and clearance in accordance with the requirements of the Paperwork Reduction Act of 1995. The submission will describe the nature of the information collection, the categories of respondents, the estimated burden (*i.e.,* the time, effort and resources used by respondents to respond) and cost, and the actual data collection instruments FEMA will use.

DATES: Comments must be submitted on or before April 20, 2015.

ADDRESSES: Submit written comments on the proposed information collection to the Office of Information and Regulatory Affairs, Office of Management and Budget. Comments should be addressed to the Desk Officer for the Department of Homeland Security, Federal Emergency Management Agency, and sent via electronic mail to *oira.submission@omb.eop.gov* or faxed to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Requests for additional information or copies of the information collection should be made to Director, Records Management Division, 500 C Street SW., Washington, DC 20472–3172, facsimile number (202) 212–4701, or email address *FEMA-Information-Collections-Management@fema.dhs.gov.*

SUPPLEMENTARY INFORMATION:

Collection of Information

Title: Level 1 Assessment Form, Level 3 Evaluation Form for Students, and Level 3 Evaluation Form for Supervisors.

Type of Information Collection: Revision of currently approved collection.

Form Titles and Numbers: FEMA Form 092–0–2, Level 1 Assessment Form; FEMA Form 092–0–2A, Level 3 Evaluation Form for Students; FEMA Form 092–0–2B, Level 3 Evaluation Form for Supervisors.

Abstract: The forms will be used to survey the Center for Domestic Preparedness (CDP) students enrolled in CDP courses and their supervisors. The surveys will collect information regarding quality of instruction, course material, and impact of training on their professional employment.

Affected Public: State, Local or Tribal Government.

Estimated Number of Respondents: 44,600.

Estimated Total Annual Burden Hours: 11,150.

Estimated Cost: 403,795.25.

Dated: March 13, 2015.

Terry Cochran,

Acting Director, Records Management Division, Mission Support, Federal Emergency Management Agency, Department of Homeland Security.

[FR Doc. 2015–06336 Filed 3–18–15; 8:45 am]

BILLING CODE 9111–53–P

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

[Docket No. FEMA–2015–0001; Internal Agency Docket No. FEMA–B–1301]

Proposed Flood Hazard Determinations

AGENCY: Federal Emergency Management Agency; DHS.

ACTION: Notice; correction.

SUMMARY: On April 4, 2013, FEMA published in the **Federal Register** a proposed flood hazard determination notice at 78 FR 20340 that contained a table which included a Web page address through which the Preliminary Flood Insurance Rate Map (FIRM), and where applicable, the Flood Insurance Study (FIS) report for the communities listed in the table could be accessed. The information available through the Web page address has subsequently been updated. The table provided here represents the proposed flood hazard