(including the use of containment bags) for these devices.

On July 11, 2014, during the afternoon session, the committee will also be asked to discuss the regulatory classification of laparoscopic power morcellator devices when used to cut and extract tissue during gynecologic laparoscopic procedures and to assist FDA in determining the appropriate level of regulatory control necessary for this device type, including discussion of class II (special controls) or reclassification to class III (subject to premarket approval application (PMA)).

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.

The Center for Devices and Radiological Health (CDRH) plans to provide a live webcast of the July 10 and 11, 2014, meeting of the Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee. While CDRH is working to make webcasts available to the public for all advisory committee meetings held at the White Oak campus, there are instances where the webcast transmission is not successful; staff will work to reestablish the transmission as soon as possible. The link for the webcast is available at: *https://colaboration.fda.gov/obgyd/,* or further information regarding the webcast, including the Web address for the webcast, will be made available at least 2 days in advance of the meeting at the following Web site: *http://www.fda.gov/AdvisoryCommittees/CommitteesMeetingMaterials/MedicalDevices/MedicalDevicesAdvisoryCommittee/ObstetricsandGynecologyDevices/default.htm.* Select the link for 2014 Meeting Materials.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before June 24, 2014. Oral presentations from the public will be scheduled between approximately 9 a.m. and 10 a.m. for both days of this meeting. 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 June 16, 2014. 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 June 19, 2014.

FDA will work with the manufacturers of laparoscopic morcellators and containment bags who wish to make presentations to ensure that adequate time, separate from the approximate time slots for the general open public hearing session, is provided. Manufacturers interested in making formal presentations to the committee should notify the contact person on or before June 18, 2014. Manufacturers with common interests are urged to coordinate their oral presentations.

FDA is opening a docket for public comment on this document. The docket number is FDA–2014–N–0736. The docket will close on August 11, 2014. Interested persons are encouraged to use the docket to submit electronic or written comments regarding this meeting. Submit electronic comments to *http://www.regulations.gov.* Comments received on or before July 1, 2014, will be provided to the committee for their consideration. Comments received after July 1, 2014, will be taken into consideration by the Agency.

Submit written comments to the Division of Dockets Management, Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. It is only necessary to send one set of comments. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Divisions of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday and will be posted to the docket at *http://www.regulations.gov.*

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

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams, at *Annmarie.Williams@fda.hhs.gov,* or 301–796–5966, at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

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

Dated: June 3, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–13290 Filed 6–6–14; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (NCI)

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH), has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on March 18, 2014, Vol. #79, page 15133 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Cancer Institute (NCI), National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974, Attention: NIH Desk Officer.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments or request more information on the proposed project contact: Vivian Horovitch-Kelley, Office of Management Policy and Compliance, National Cancer Institute, 9609 Medical Center Drive, Bethesda, MD 20892–9760 or call non-toll-free number 240–276–6850 or Email your request including your address to: *horovitchkellv@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (NCI), 0925–0642, Expiration Date 9/31/2014, Extension, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: There are no changes being requested for this submission. The information collection activity will garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. This generic will provide information about the National Cancer Institute's customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. It will also allow feedback to contribute directly to the improvement of program management. Feedback collected under this generic clearance will provide useful information but it will not yield data that can be generalized to the overall population.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated burden hours are 8,750.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of collection	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Surveys	1000	1	30/60	500
In-Depth Interviews (IDIs) or Small Discussion Groups	500	1	90/60	750
Focus Groups	2000	1	90/60	3,000
Website or Software Usability Tests	3000	1	90/60	4,500

Dated: May 27, 2014.

Karla Bailey,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2014–13411 Filed 6–6–14; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Library of Medicine; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App), notice is hereby given of the meetings.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable materials, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Biomedical Library and Informatics Review Committee.

Date: November 6–7, 2014.

Time: November 6, 2014, 8 a.m. to 6 p.m.

Agenda: To review and evaluate grant applications.

Place: National Library of Medicine, Building 38, 2nd Floor, Board Room, 8600 Rockville Pike, Bethesda, MD 20892.

Time: November 7, 2014, 8 a.m. to 2 p.m.

Agenda: To review and evaluate grant applications.

Contact Person: Arthur A. Petrosian, Ph.D., Chief Scientific Review Officer, Division of Extramural Programs, National Library of Medicine, 6705 Rockledge Drive, Suite 301, Bethesda, MD 20892–7968, 301–496–4253, *petrosia@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program No. 93.879, Medical Library Assistance, National Institutes of Health, HHS)

Dated: June 3, 2014.

Michelle Trout,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2014–13328 Filed 6–6–14; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the National Cancer Advisory Board and NCI Board of Scientific Advisors, June 22, 2014, 4 p.m. to June 24, 2014, 12 p.m., National Institutes of Health, Building 31, C Wing, 6th Floor, 31 Center Drive, Conference Room 10, Bethesda, MD 20892 which was published in the **Federal Register** on May 7, 2014, 79 FR 26259.

The meeting notice is being amended to cancel the Subcommittee meeting on Planning and Budget on June 22, 2014. The meeting notice is also being amended to change the following open and closed session times of the Joint Board meeting. On June 23, 2014, the open session will be from 8:30 a.m. to 6 p.m. On June 24, 2014, the open session will be from 8:30 a.m. to 10:45 a.m. and the closed session will be from 10:45 a.m. to 12 p.m. The meeting is partially closed to the public.

Dated: June 3, 2014.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2014–13330 Filed 6–6–14; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as