The docket will close 30 days after those documents are posted.

Transcripts: Please be advised that as soon as a transcript is available, it will be accessible at in the docket at *http://www.regulations.gov.* It may be viewed at the Division of Dockets Management (see **ADDRESSES**). A transcript will also be available in either hardcopy or on CD–ROM, after submission of a Freedom of Information request. The Freedom of Information office address is available on the Agency's Web site at *http://www.fda.gov.* A link to the transcripts will also be available approximately 45 days after the public workshop on the Internet at *http://www.fda.gov/ForIndustry/UserFees/MedicalDeviceUserFee/ucm454039.htm.* (Select this meeting from the posted events list).

Dated: October 3, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016–24237 Filed 10–6–16; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request; Data Use Agreement and Supplement for 2014 Health Center Patient Survey

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

ACTION: Notice.

SUMMARY: In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, HRSA submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period.

DATES: Comments on this ICR should be received no later than November 7, 2016.

ADDRESSES: Submit your comments, including the ICR title, to the desk officer for HRSA, either by email to *OIRA_submission@omb.eop.gov* or by fax to 202–395–5806.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email the HRSA Information Collection Clearance Officer at *paperwork@hrsa.gov* or call (301) 443–1984.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: Data Use Agreement and Supplement for 2014 Health Center Patient Survey.

OMB No.: 0906–xxxx—NEW.

Abstract: The Health Center Patient Survey (HCPS), sponsored by HRSA's Bureau of Primary Health Care (BPHC), surveyed patients who use health centers funded under Section 330 of the Public Health Service Act. HCPS collects data on health center patients' sociodemographic characteristics, health conditions, health behaviors, access to and utilization of health care services, and satisfaction with health care. Survey results come from in-person, one-on-one interviews with patients and are nationally representative of the Health Center program patient population. To inform BPHC and HHS policy, funding, and planning decisions, the survey investigated how well HRSA-supported sites meet health care needs of the medically underserved and assessed how patients perceive the quality of their care.

The HCPS is unique because it focuses on comprehensive patient-level data. These and other features of the data will provide researchers and policymakers the capacity to empirically explore policy relevant topics relevant to the Health Center program using up-to-date information.

Prior to releasing this information, BPHC will request prospective users to fill out a ''Data Use Agreement'' (DUA). BPHC uses DUAs as legal binding agreements when an external entity (*e.g.,* contractor, private industry, academic institution, other federal government agency, or state agency) requests the use of BPHC personally/organizationally identifiable data that is covered by the Privacy Act of 1974. The agreement delineates the confidentiality requirements of the Privacy Act, security safeguards, and BPHC's data use policies and procedures. The DUA will serve as both a means of informing data users of these requirements and a means of obtaining their agreement to abide by these requirements.

Need and Proposed Use of the Information: Before allowing access to unrestricted data that contains sensitive grantee and patient information that is protected by the Privacy Act of 1974, prospective users will submit a signed DUA and describe what proposed research they intend to undertake in using the dataset. A BPHC workgroup will determine whether the project is an appropriate and legitimate use of the data. The criteria to determine admissible projects will include: (1) Relevance of the topic of study to BPHC/HHS policy; (2) feasibility of the project given the parameters described in DUA supplemental; and (3) the proposed end-use of the research that will be undertaken.

Likely Respondents: Prospective researchers in academia, private contractors, and Primary Care Associations/Health Center grantee organizations.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. 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. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
DUA	20	1	20	0.25	5
DUA Supplemental	20	1	20	1.25	25
Total	40	.....	40	.....	30

Jason E. Bennett,

Director, Division of the Executive Secretariat.

[FR Doc. 2016–24300 Filed 10–6–16; 8:45 am]

BILLING CODE 4165–15–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 amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings 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 material, 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: Center for Scientific Review Special Emphasis Panel; PAR13–309–311: Translational Research in Pediatric and Obstetric Pharmacology and Therapeutics.

Date: November 1, 2016.

Time: 10:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Dianne Hardy, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6175, MSC 7892, Bethesda, MD 20892, 301–435–1154, *dianne.hardy@nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; RFA AI016–025: Non-vaccine Biomedical Prevention (nBP) of HIV Acquisition/Transmission.

Date: November 1, 2016.

Time: 1:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Jingsheng Tuo, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5207, Bethesda, MD 20892, 301–451–8754, *tuoj@nei.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Biomedical Research Shared Instrumentation.

Date: November 1, 2016.

Time: 2:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Kathryn Kalasinsky, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5158 MSC 7806, Bethesda, MD 20892, 301–402–1074, *kalasinskyks@mail.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Topics in Infectious Diseases.

Date: November 1, 2016.

Time: 2:00 p.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: John C Pugh, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3114, MSC 7808, Bethesda, MD 20892, (301) 435–2398, *pughjohn@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR Panel: Small Business: Serious STEM Games for Pre-College and Informational Science Education Audiences.

Date: November 2, 2016.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Residence Inn Bethesda, 7335 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Allen Richon, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6184, MSC 7892, Bethesda, MD 20892, 301–379–9351, *allen.richon@nih.hhs.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict Bioengineering Sciences #2.

Date: November 2, 2016.

Time: 2:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Joseph Thomas Peterson, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4118, MSC 7814, Bethesda, MD 20892, 301–408–9694, *petersonjt@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Shared Instrumentation: Confocal Microscopy and Imaging.

Date: November 3–4, 2016.

Time: 8:00 a.m. to 11:30 a.m.

Agenda: To review and evaluate grant applications.

Place: Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road, NW., Washington, DC 20015.

Contact Person: Thomas Beres, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Dr. Rm. 5201, MSC 7840, Bethesda, MD 20892, 301–435–1175, *berestm@mail.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Shared Instrumentation: Electron Microscopy.

Date: November 4, 2016.

Time: 1:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road, NW., Washington, DC 20015.

Contact Person: Thomas Beres, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Dr. Rm. 5201, MSC 7840, Bethesda, MD 20892, 301–435–1175, *berestm@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: October 3, 2016.

Sylvia L. Neal,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–24254 Filed 10–6–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the NIH Clinical Center Research Hospital Board scheduled for October 21, 2016, 9:00 a.m. to 5:00 p.m., in Conference Room 6C6, Building 31, National Institutes of Health, Bethesda, MD 20892, which was published in the **Federal Register** on September 16, 2016, 81 FR 63778.

A discussion of the reports from the Anonymous Safety Hotline developed by NIH for Clinical Center staff, patients, or visitors to report any concerns about care or unsafe conditions has been added to the closed portion of the meeting. This portion will be closed to the public in accordance with the provisions set forth in section 552b(c)(9)(B) and 552b(c)(6), Title 5 U.S.C., as amended. The premature disclosure of the laboratories or units and staff involved in the individual reports could significantly limit the Hotline's purpose by compromising anonymity. These actions would frustrate NIH's use of this resource as it strives to improve the overall safety and quality of care at the Clinical Center.

There are no changes for the open portion of the meeting that was advertised on September 16, 2016.