This request is to expand the UDS data reporting resource to the BHW NMHC grantees and IPCP program cooperative agreement awardees. Calendar year data would be submitted annually to enable BHW to track clinical practice and patient outcome data. The data collection is limited to NMHC and IPCP grantees and cooperative agreement awardees because of the similarities these care models share with health centers; therefore, the use of the pre-existing infrastructure will enable HRSA to populate the data set with additional sources, making the resource more robust.

Need and Proposed Use of the Information: HRSA collects UDS data which are used to ensure compliance with legislative and regulatory requirements, improve grantee and cooperative agreement awardee performance and operations, and report overall program accomplishments. BHW proposes to collect core data elements that include patient demographics, healthcare services, clinical indicators and outcomes, provider utilization, and costs. BHW will use the patient and provider-level data to determine the impact of healthcare services on patient outcomes. The data will also enable BHW to establish or expand targeted programs and identify effective services and interventions to improve the health of underserved communities and vulnerable populations. In addition, the UDS data are useful to BHW grantees and cooperative agreement awardees for performance and operations improvement, patient forecasts, identification of trends/patterns, implication of access barriers, and cost analysis to support long-term sustainability.

Likely Respondents: The respondents will be HRSA BHW Nurse Managed Health Clinic (NMHC) grantees and Interprofessional Collaborative Practice (IPCP) program cooperative agreement awardees.

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 Information Collection Request are summarized in the table below.

Total estimated annualized hours: Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Universal Report	81	1	81	170	13,770
Grant Report	81	1	81	22	1,782
Total	162				15,552

HRSA specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Jackie Painter,

Director, Division of the Executive Secretariat.

[FR Doc. 2015–17552 Filed 7–16–15; 8:45 am]

BILLING CODE 4165–15–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

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Health Resources and Services Administration (HRSA) has 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 August 17, 2015.

ADDRESSES: Submit your comments, including the Information Collection Request 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: Rural Access to Emergency Devices Grant Program OMB No. 0915–xxxx—NEW.

Abstract: This program is authorized by the Public Health Improvement Act Title IV—Cardiac Arrest Survival Act of 2000, Subtitle B—Rural Access to Emergency Devices, Section 413, (42 U.S.C. 254c (Note) and the Consolidated and Further Continuing Appropriations Act (Pub. L. 113–235). The purpose of this grant program is to: (1) Purchase automated external defibrillators (AEDs) that have been approved, or cleared for marketing, by the Food and Drug Administration; (2) provide defibrillator and basic life support training in AED usage through the American Heart Association, the American Red Cross, or other nationally recognized training courses; and (3) place the AEDs in rural communities with local organizations.

Need and Proposed Use of the Information: For this program, performance measures were drafted to provide data useful to the program and to enable HRSA to provide aggregate program data required by Congress under the Government Performance and Results Act (GPRA) of 1993 (Pub. L. 103–62). These measures cover the principal topic areas of interest to the Federal Office of Rural Health Policy, including: (a) The number of counties served by the program; (b) the number of AEDs purchased and placed and the

locations of the placements; (c) the number of training sessions and the number of individuals trained; (d) the number of times an AED is used and the outcome; and (e) the number of lay persons and first responders who administer CPR or use an AED on an individual. These measures will speak to the Federal Office of Rural Health Policy's progress toward meeting the set goals.

A 60-day **Federal Register** notice was published February 20, 2015 (80 FR 9270–9271). There were no comments.

Likely Respondents: Rural Access to Emergency Devices Grant Program award recipients.

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
Rural Access to Emergency Devices Grant Program	12	1	12	5.5	66
Total	12				66

Jackie Painter,

Director, Division of the Executive Secretariat.

[FR Doc. 2015–17550 Filed 7–16–15; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

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 meeting.

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 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: National Institute of Allergy and Infectious Diseases Special Emphasis Panel; NIAID SBIR Phase II Clinical Trial Implementation Cooperative Agreement and Clinical Trial Planning Grants.

Date: August 27, 2015.

Time: 12:00 p.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 5601 Fisher Lane, Rockville, MD 20892, (Telephone Conference Call).

Contact Person: Paul A. Amstad, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, Room 3G41, NIAID/NIH/DHHS, 5601 Fishers Lane, Bethesda, MD 20892–7616, 240–669–5067, *pamstad@niaid.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)

Dated: July 14, 2015.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015–17595 Filed 7–16–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases Notice of Closed Meeting

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 meeting.

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 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: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; Ancillary Studies on IBD.

Date: August 10, 2015.

Time: 4:30 p.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Maria E. Davila-Bloom, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 758, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–7637, *davila-bloomm@extra.niddk.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: July 14, 2015.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015–17594 Filed 7–16–15; 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 Meeting

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 meeting.