

**SUPPLEMENTARY INFORMATION:**

**Registration:** To attend the public workshop, you must register before February 2, 2017, by visiting <https://healthpolicy.duke.edu/events/ninth-annual-sentinel-initiative-public-workshop>. You may also register for the live Webcast by visiting this Web page. There will be no onsite registration. When registering, please provide the following information: Your name, title, company or organization (if applicable), postal address, telephone number, and email address. Those without Internet access should contact Carlos Bell to register (See **FOR FURTHER INFORMATION CONTACT**). There is no registration fee for the public workshop. However, registration will be on a first-come, first-served basis because seating is limited. Therefore, early registration is recommended. Upon registering, attendees will receive an confirmatory email, containing event materials. A 1-hour lunch break is scheduled, but food will not be provided. There are multiple restaurants within walking distance of the Barbara Jordan Conference Center at the Kaiser Family Foundation.

If you need special accommodations due to a disability, please contact Joanna Higgison at the Duke-Margolis Center for Health Policy (phone: 908-432-4872, email: [joanna.higgison@duke.edu](mailto:joanna.higgison@duke.edu)) at least 7 days in advance.

**Streaming Webcast of the Public Workshop:** This public workshop will also be Webcast (archived video footage will be available following the workshop at <https://healthpolicy.duke.edu/events/ninth-annual-sentinel-initiative-public-workshop>). Persons interested in viewing the live Webcast must register online by February 1, 2017, at 5 p.m. EST. Early registration is recommended because Webcast connections are limited. Organizations are requested to register all participants, and view the workshop using one connection per location whenever possible. Webcast participants will be sent technical system requirements upon registering. Prior to joining the streaming Webcast of the public workshop, it is recommended that you review these technical system requirements.

**Meeting Materials:** All event materials will be sent to registered attendees via email before the workshop. The event materials will also be available to view on the Duke-Margolis Web site at <https://healthpolicy.duke.edu/events/ninth-annual-sentinel-initiative-public-workshop>.

**Transcripts:** Please be advised that transcripts will not be available.

Dated: November 3, 2016.

**Leslie Kux,**

*Associate Commissioner for Policy.*

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**BILLING CODE 4164-01-P**

## **DEPARTMENT OF HEALTH AND HUMAN SERVICES**

### **Health Resources and Services Administration**

#### **Agency Information Collection Activities: Proposed Collection: Public Comment Request; The Advanced Education Nursing Traineeship (AENT) Program Specific Data Collection Forms**

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

**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 January 9, 2017.

**ADDRESSES:** Submit your comments to [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or mail the HRSA Information Collection Clearance Officer, Room 14N39, 5600 Fishers Lane, Rockville, MD 20857.

**FOR FURTHER INFORMATION CONTACT:** To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or call the HRSA Information Collection Clearance Officer at (301) 443-1984.

**SUPPLEMENTARY INFORMATION:** When submitting comments or requesting information, please include the information request collection title for reference.

**Information Collection Request Title:** The Advanced Education Nursing Traineeship (AENT) Program Specific Data Collection Forms.

OMB No. 0915-0375, Revision

**Abstract:** The Advanced Nursing Education Workforce (ANEW) Program is a new program that incorporates elements of the AENT and the Advanced Nursing Education Programs. The current OMB approved Program Specific Data Collection Forms for the

former AENT Program will be simplified and used for the ANEW program.

HRSA provides advanced education nursing grants to educational institutions to increase the numbers of advanced education nurses through the ANEW Program. The ANEW Program is authorized by Title VIII, Section 811(a)(2) of the Public Health Service Act, (42 U.S.C. 296j(a)(2)), as amended by Section 5308 of the Patient Protection and Affordable Care Act of 2010, Public Law 111-148. This renewal with revision request includes the Project Abstract, Program Narrative, Attachments, and Tables. The proposed ANEW tables are very similar to the previous AENT tables and include information on program participants such as the projected number of enrollees/trainees receiving traineeship support, projected number of graduates receiving traineeship support for the previous fiscal year, the types of programs they are enrolling into and/or from which enrollees/trainees are graduating, and the distribution of primary care nurse practitioners, primary care clinical nurse specialists, and nurse-midwives who plan to practice in rural and underserved settings. To reduce the reporting burden for applicants, HRSA simplified the tables to focus on the types of providers and practice settings that are included in the statute to determine whether applicants qualify for the preference or special consideration in making awards for this program.

**Need and Proposed Use of the Information:** The Project Abstract is often distributed to provide information to the public and Congress. HRSA will use this information in determining the eligibility for the statutory funding preference and special consideration and to succinctly capture data for the number of projected students for subsequent years in the project period.

**Likely Respondents:** Likely respondents are potential applicants for the ANEW program. Eligible applicants for the ANEW program include entities that provide registered nurses with primary care nurse practitioner (NP), primary care clinical nurse specialist (CNS), and nurse-midwife education. Such programs may include accredited schools of nursing, nursing centers, academic health centers, state or local governments, and other public or private nonprofit entities authorized by the Secretary of HHS to confer degrees to RNs for primary care NP, primary care CNS, or nurse-midwife education. Federally recognized Indian Tribal Government and Native American Organizations as well as faith-based or

community-based organizations may apply if they are otherwise eligible.

Eligible state government entities include the 50 states, the District of Columbia, Guam, the Commonwealth of Puerto Rico, the Northern Mariana Islands, American Samoa, the U.S. Virgin Islands, the Federated States of Micronesia, the Republic of the Marshall Islands, and the Republic of Palau.

**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 Burden Hours:*

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
ANEW Application including the ANEW Program Specific Tables and Attachments .....	236	1	236	7	1,652
Total .....	236	1	236	7	1,652

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.

**Jason E. Bennett,**

*Director, Division of the Executive Secretariat.*

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**BILLING CODE 4165-15-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### **Proposed Collection; 60-Day Comment Request; NCI Genomic Data Commons (GDC) Data Submission Request Form (National Cancer Institute)**

**AGENCY:** National Institutes of Health.

**ACTION:** Notice.

**SUMMARY:** In compliance with the requirement of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

**DATES:** Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

**FOR FURTHER INFORMATION CONTACT:** To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Louis M. Staudt, MD, Ph.D., Director, Center for Cancer Genomics, National Cancer Institute, Building 10 Room 5A02, 10 Center Drive, Bethesda MD 20814 or call non-toll-free number 301-402-1892 or Email your request, including your address to: *lstaudt@mail.nih.gov*. Formal requests for additional plans and instruments must be requested in writing.

**SUPPLEMENTARY INFORMATION:** Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of

information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

**Proposed Collection Title:** NCI Genomic Data Commons (GDC) Data Submission Request Form, 0925-NEW, National Cancer Institute (NCI), National Institutes of Health (NIH).

**Need and Use of Information Collection:** The purpose of the NCI Genomic Data Commons (GDC) Data Submission Request Form is to provide a vehicle for investigators to request submission of their cancer genomic data into the GDC in support of data sharing. The purpose is to also provide a mechanism for the GDC Data Submission Review Committee to review and assess the data submission request for applicability to the GDC mission. The scope of the form involves obtaining information from investigators that: (1) Would like to submit data about their study into the GDC, (2) are affiliated with studies that adhere to GDC data submission conditions. The benefits of the collection are that it provides the needed information for investigators to understand the types of studies and data that the GDC supports, and that it provides a standard mechanism for the GDC to assess incoming data submission requests.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 50 hours.