

examine 1115 Medicaid waivers that have expanded eligibility to include specifically people living with HIV/AIDS (PLWH) who are not otherwise eligible for Medicaid services. Since 1990, the Ryan White HIV/AIDS Program (RWHAP) has provided funding for primary care, medications, and support services for PLWH, helping fill the health care and service gap for those who are uninsured or ineligible for Medicaid.

As part of this project, case studies will be conducted in eight states that have implemented 1115 Medicaid waivers to expand Medicaid eligibility for PLWH. The case studies will include site visits and discussions with the state Medicaid programs and with RWHAP grantees and service providers to examine the waivers and their impact on PLWH. In addition, the studies will explore whether and how the 1115

Medicaid waivers have helped states and RWHAP grantees and providers prepare for implementation of the Affordable Care Act, including providing insights into Medicaid expansion.

**Need and Proposed Use of the Information:** Given the important role of the RWHAP and Medicaid in meeting the health care needs of PLWH, there is a need to understand better, how Medicaid expansion and the 1115 Medicaid waivers will affect the RWHAP and how the waivers have prepared states for implementation of the Affordable Care Act.

**Likely Respondents:** Data will be collected through qualitative interviews, guided by discussion tools with questions tailored for four specific groups of individuals from: (1) State Medicaid agencies; (2) RWHAP Part B grantees and service providers; (3)

RWHAP Part A grantees and service providers; and (4) and RWHAP White Part C grantees and clinical providers.

**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 |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------|------------------------------------|-----------------|----------------------------------------|--------------------|
| Qualitative Interview Data Collection Tool for State Medicaid Agency Groups .....                                                                               | 40                    | 1                                  | 40              | 2                                      | 80                 |
| Qualitative Interview Data Collection Tool for Ryan White Part A Administrators and Members of Planning Councils .....                                          | 64                    | 1                                  | 64              | 2                                      | 128                |
| Qualitative Interview Data Collection Tool for Ryan White Part A Administrators and Members of Planning Councils .....                                          | 16                    | 1                                  | 16              | 2                                      | 32                 |
| Qualitative Interview Data Collection Tool for Ryan White Part B and ADAP (AIDS Directors, Part B Coordinators and ADAP Coordinators) .....                     | 80                    | 1                                  | 80              | 2                                      | 160                |
| Qualitative Interview Data Collection Tool for Ryan White Clinical Providers (RW Part C Grantees in clinical settings or Similar Clinical Care Providers) ..... | 80                    | 1                                  | 80              | 2                                      | 160                |
| Total .....                                                                                                                                                     | 280                   | .....                              | .....           | .....                                  | 560                |

Dated: July 3, 2013.

**Bahar Niakan,**

*Director, Division of Policy and Information Coordination.*

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#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

##### Health Resources and Services Administration

##### Agency Information Collection Activities; Proposed Collection; Public Comment Request

**AGENCY:** Health Resources and Services Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** In compliance with the requirement for opportunity for public comment on proposed data collection projects (Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995), the Health Resources and Services Administration (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 Information Collection Request must be received within 60 days of this notice.

**ADDRESSES:** Submit your comments to [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or mail the HRSA Information Collection Clearance

Officer, Room 10-29, Parklawn Building, 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:  
Scholarships for Disadvantaged  
Students Program OMB No. 0915–  
0149—Renewal**

The purpose of the Scholarships for Disadvantaged Students (SDS) Program is to provide funds to eligible schools to provide scholarships to full-time, financially needy students from disadvantaged backgrounds enrolled in health professions and nursing programs.

To qualify for participation in the SDS program, a school must be carrying out a program for recruiting and retaining students from disadvantaged backgrounds, including students who are members of racial and ethnic

minority groups (section 737(d)(1)(B) of the PHS Act). A school must meet the eligibility criteria to demonstrate that the program has achieved success based on the number and/or percentage of disadvantaged students who graduate from the school. In awarding SDS funds to eligible schools, funding points must be given to schools based on the proportion of graduating students going into primary care, the proportion of underrepresented minority students, and the proportion of graduates working in medically underserved communities (section 737(c) of the PHS Act).

**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              | Number of respondents | Number of responses per respondent | Total responses | Hours per response | Total hour burden |
|-------------------|-----------------------|------------------------------------|-----------------|--------------------|-------------------|
| Application ..... | 400                   | 1                                  | 400             | 13                 | 5,200             |
| Total .....       | 400                   | 1                                  | 400             | 13                 | 5,200             |

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.

Dated: July 3, 2013.

**Bahar Niakan,**

*Director, Division of Policy and Information Coordination.*

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**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 within 30 days of this notice.

**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](mailto: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](mailto:paperwork@hrsa.gov) or call (301) 443–1984.

**SUPPLEMENTARY INFORMATION:**

**Information Collection Request Title:  
Health Center Program Application  
Forms**

OMB No. 0915–0285—Revision  
**Abstract:** Health centers (section 330 grant funded and Federally Qualified Health Center Look-Alikes) deliver comprehensive, high quality, cost-effective primary health care to patients regardless of their ability to pay. Health centers have become an essential primary care provider for America's most vulnerable populations. Health centers advance the preventive and primary medical/health care home

model of coordinated, comprehensive, and patient-centered care, coordinating a wide range of medical, dental, behavioral, and social services. More than 1,200 health centers operate nearly 9,000 service delivery sites that provide care in every state, the District of Columbia, Puerto Rico, the U.S. Virgin Islands, and the Pacific Basin.

The Health Centers Program is administered by HRSA's Bureau of Primary Health Care (BPHC). HRSA/BPHC uses the following application forms to oversee the Health Center Program. These application forms are used by new and existing health centers to apply for various grant and non-grant opportunities, renew their grant or non-grant designation, and change their scope of project.

**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