

ANNUAL BURDEN ESTIMATES

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Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ACF-196	51	4	10	2,040

Estimated Total Annual Burden Hours: 2,040.

In compliance with the requirements of the Paperwork Reduction Act of 1995 (Pub. L. 104-13, 44 U.S.C. Chap 35) Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington DC 20201. Attn: ACF Reports Clearance Officer. Email address: infocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2016-14945 Filed 6-23-16; 8:45 am]

BILLING CODE 4184-01-P

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 ICR should be received no later than August 23, 2016.

ADDRESSES: Submit your comments to paperwork@hrsa.gov or mail the HRSA Information Collection Clearance Officer, Room 14N-39, 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 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: Ryan White HIV/AIDS Program Outcomes within the Context of the Affordable Care Act, OMB No. 0906-xxxx—New.

Abstract: The Health Resources and Services Administration's HIV/AIDS Bureau (HRSA/HAB) implements the Ryan White HIV/AIDS Program (RWHAP). This program provides HIV-related services in the United States for those who do not have sufficient health care coverage or financial resources for coping with HIV disease. Enacted in 2010, the Affordable Care Act has had profound impacts on health care financing and delivery that are continuing to unfold. The expansion of health care coverage impacted many of RWHAP's traditional clients who are now eligible to receive health care coverage through Medicaid coverage

and qualified health plans available on Health Insurance Marketplaces. These changes have required RWHAP sites to adapt in order to fill different gaps in care experienced by clients across the varying health care coverage options. The purpose of this evaluation study is to determine the effect that Affordable Care Act related health care coverage has had on overall health outcomes, service utilization, and gaps in care for people living with HIV. This evaluation seeks to understand how RWHAP provider sites meet the needs of clients under the variety of health care coverage options clients are encountering across the country.

Need and Proposed Use of the Information: The expansion of health care coverage now offers new options of obtaining health care services for many individuals with HIV. Due to these changes, additional information concerning overall client health outcomes, pharmaceutical and core medical processes and outcomes, and client access to and utilization of support services is needed. Data from this evaluation study will be used to provide HRSA/HAB with the necessary information to understand the changes in primary health care outcomes of RWHAP clients' pre- and post-implementation of the Affordable Care Act. This will inform how the RWHAP can best serve clients in the environment of the health care reform.

Likely Respondents: RWHAP administrators, RWHAP care providers, and RWHAP clients are the likely respondents.

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
Site Survey	305	1	305	0.5	152.5
Medical Records Sample Selection Guide	25	1	25	1	25
Site Interview Guide	50	1	50	2	100
Focus Groups Guide	60	1	60	1.5	90
Total	440	440	367.5

HRSA specifically requests comments on (1) the necessity and practical utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the agency's estimate of the burden of the proposed collection of information; (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 Executive Secretariat.

[FR Doc. 2016-14951 Filed 6-23-16; 8:45 am]

BILLING CODE 4165-15-P

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.

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Officer, Room 14N-39, 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 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: Healthy Start Evaluation and Quality Improvement OMB No. 0915-0338—Revision.

Abstract: The National Healthy Start Program, funded through HRSA's Maternal and Child Health Bureau (MCHB), has the goal of reducing disparities in infant mortality and adverse perinatal outcomes. The program began as a demonstration project with 15 grantees in 1991 and has expanded over the past 2 decades to 100 grantees across 37 states and Washington, DC. Healthy Start grantees operate in communities with rates of infant mortality at least 1.5 times the U.S. national average and high rates for other adverse perinatal outcomes. These communities are geographically, racially, ethnically, and linguistically diverse low-income areas. Healthy Start covers services during the perinatal period (before, during, after pregnancy) and follows the woman and infant through 2 years after the end of the pregnancy. The Healthy Start program has five approaches, including: (1) Improving women's health; (2) promoting quality services; (3) strengthening family resilience; (4) achieving collective impact; and (5) increasing accountability through quality assurance, performance monitoring, and evaluation.

MCHB seeks to implement a uniform set of data elements for monitoring and conducting a mixed-methods evaluation to assess the effectiveness of the

program on individual, organizational, and community-level outcomes. Data collection instruments will include a National Healthy Start Program Survey; Community Action Network Survey; Healthy Start Site Visit Protocol; Healthy Start Participant Focus Group Protocol—these instruments have not been changed. The Preconception, Pregnancy and Parenting (3Ps) Information Form will also be used as a data collection instrument; however the 3Ps Information form has been redesigned from one form into six forms. The six forms include: (1) Demographic Intake Form; (2) Pregnancy Status/History; (3) Preconception; (4) Prenatal; (5) Postpartum; and (6) Interconception/Parenting. The purpose of this redesign is to enhance the 3Ps Information Form to ensure collected data is meaningful for monitoring and evaluation, as well as screening and care coordination, and streamline previously separate data systems. The 3Ps Information Form was also redesigned to allow questions to be administered in accordance with the participant's enrollment/service delivery status and perinatal period. In addition to redesigning the 3Ps Information Form, HRSA deleted questions that are neither critical for evaluation nor programmatic purposes. HRSA also added questions to the 3Ps Information Form to allow the Form to be used as an all-inclusive data collection instrument for MCHB and Healthy Start grantees. The additional questions extend and refine previously approved content, allowing for the collection of more granular and/or in-depth information on existing topics. Adding these questions allows Healthy Start grantees to better assess risk, identify needed services, provide appropriate follow-up activities to program participants, and improve overall service delivery and quality.

Need and Proposed Use of the Information: The purpose of the data collection instruments is to obtain consistent information across all grantees about Healthy Start and its