

maintain a text message library, which addresses topics of HIV disease management, e.g. appointment keeping, retention in care, and medication adherence rates; and (2) to develop, implement, conduct, and evaluate a pilot study of delivering text messages to targeted youth receiving care at Ryan White grantee sites and other clinical sites.

The first phase of this project will include focus group interviews with the target audience to test the messages (Aim 1). Approximately 128 individuals will be screened to assess focus group eligibility. Four focus groups will be conducted with up to eight participants in each for a total sample size of 32.

The second phase of this project involves the evaluation of the pilot study (Aim 2). This will encompass data

collection with patients and providers. Patient participants for the pilot study will be recruited from ten clinical sites, some of which will be Ryan White grantees. Up to 1,000 individuals will be screened to determine eligibility for the pilot study to recruit a sample of 500 participants (50 from each clinical site). Patient participants will complete a baseline survey, 3-month survey, 6-month survey, and follow-up survey at 9 months. In addition, ten patient participants from each clinical site will be selected to participate in an in-depth, qualitative telephone interview for a total of 100 interviews. Finally, up to three clinic staff from the ten participating clinics will take part in in-depth, qualitative telephone interviews (N=30).

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
Patient Focus Group Screener	128	1	128	0.25	32
Patient Focus Group Interview	32	1	32	2.0	64
Patient Pilot Study Screener	1000	1	1000	0.25	250
Patient Pilot Study Surveys	500	4	2000	0.75	1500
Patient Pilot Study Qualitative Interviews	100	1	100	1.0	100
Clinic Staff Pilot Study Qualitative Interviews	30	1	30	0.75	22.5
Total	1790	3290	1968.5

Dated: August 28, 2013.

Bahar Niakan,

Director, Division of Policy and Information Coordination.

[FR Doc. 2013–21557 Filed 9–4–13; 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.

DATES: Comments on this Information Collection Request must be received within 60 days of this notice.

ADDRESSES: Submit your comments to paperwork@hrsa.gov or mail the HRSA Information Collection Clearance Officer, Room 10–29, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 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: HRSA Telehealth Outcome Measures.

OMB No.: 0915–0311—Extension.

Abstract: To help carry out its mission, the Office for the Advancement of Telehealth (OAT) created a set of

performance measures that grantees can use to evaluate the effectiveness of their services programs and monitor their progress through the use of performance reporting data.

Need and Proposed Use of the Information: As required by the Government Performance and Review Act of 1993 (GPRA), all federal agencies must develop strategic plans describing their overall goal and objectives. The Office for the Advancement of Telehealth (OAT) has worked with its grantees to develop performance measures to be used to evaluate and monitor the progress of the grantees. Grantee goals are to: improve access to needed services; reduce rural practitioner isolation; improve health system productivity and efficiency; and improve patient outcomes. In each of these categories, specific indicators were designed to be reported through a performance monitoring Web site.

Likely Respondents: Telehealth Network Grantees.

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
Performance Improvement Measurement System (PIMS) ..	700	2	1400	7	9,800

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: August 28, 2013.

Bahar Niakan,

Director, Division of Policy and Information Coordination.

[FR Doc. 2013-21567 Filed 9-4-13; 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.

DATES: Comments on this Information Collection Request must be received within 60 days of this notice.

ADDRESSES: Submit your comments to 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 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 National Health Service Corps Loan Repayment Program

OMB No. 0915-0127—Revision

Abstract: The National Health Service Corps (NHSC) Loan Repayment Program (LRP) was established to assure an adequate supply of trained primary care health professionals to provide services in the neediest Health Professional Shortage Areas (HPSAs) of the United States. Under this program, the Department of Health and Human Services agrees to repay the qualifying educational loans of selected primary

care health professionals. In return, the health professionals agree to serve for a specified period of time in a federally designated HPSA approved by the Secretary for LRP participants. The forms utilized by the LRP include the following: the NHSC LRP Application, the Authorization for Disclosure of Loan Information form, the Privacy Act Release Authorization form, the Verification of Disadvantaged Background form, and the Private Practice Option form. The first four of the aforementioned NHSC LRP forms collect information that is needed for selecting participants and repaying qualifying educational loans. The last referenced form, the Private Practice Option Form, is required by statute (42 U.S.C. 254n(a)) for all participants wishing to exercise that service option.

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	Responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
NHSC LRP Application	8,200	1	8,200	1.00	8,200
Authorization for Disclosure of Loan Information Form	150	1	150	.10	15
Privacy Act Release Authorization Form	100	1	100	.10	10