

percent each year for the first 2 years and 25 percent for the third year. In exchange, the nurses agree to serve full-time as a registered nurse for 2 or 3 years at a health care facility with a critical shortage of nurses.

NELRP requires the following information:

1. Applicants must provide information on their nursing education, employment, and proposed service site;

2. Applicants must provide information on their outstanding nursing educational loans;

3. Applicants must provide banking information from their financial institution; and

4. Employers must provide information on the health care facility and on the employment status of applicants and participants.

#### ESTIMATES OF ANNUALIZED HOUR BURDEN ARE AS FOLLOWS FOR APPLICANTS

Form	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
NELRP Application .....	5,000	1	5,000	1.5	7,500
Loan Verification Form .....	5,000	3	15,000	1	15,000
Applicant Employment Verification Form .....	5,000	1	5,000	.5	2,500
Payment Information Form .....	5,000	1	5,000	1	5,000
Application Checklist .....	5,000	1	5,000	.5	2,500
Pre-Award Confirmation Checklist .....	600	1	600	.25	150
Total .....	5,000		35,600		32,650

#### ESTIMATES OF ANNUALIZED HOUR BURDEN ARE AS FOLLOWS FOR PARTICIPANTS

Form	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Participant semi-annual employment verification form .....	1,300	2	2,600	.5	1,300
Total .....	1,300	2	2,600	.5	1,300

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to [OIRA\\_submission@omb.eop.gov](mailto:OIRA_submission@omb.eop.gov) or by fax to 202-395-6974. Please direct all correspondence to the "attention of the desk officer for HRSA."

Dated: October 17, 2007.

**Alexandra Huttinger,**

*Acting Director, Division of Policy Review and Coordination.*

[FR Doc. E7-20940 Filed 10-23-07; 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: Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Public Law 104-13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed

for submission to the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, call the HRSA Reports Clearance Officer on (301) 443-1129.

Comments are invited on: (a) The proposed collection of information for the proper performance of the functions of the agency; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance 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.

#### Proposed Project: Ryan White HIV/AIDS Program Core Medical Services Waiver Application Requirements (OMB No. 0915-0307): Revision

Title XXVI of the Public Health Service (PHS) Act, as amended by the Ryan White HIV/AIDS Treatment Modernization Act of 2006 (Ryan White HIV/AIDS Program), requires that grantees expend 75 percent of Parts A, B, and C funds on core medical services, including antiretroviral drugs, for individuals with HIV/AIDS identified and eligible under the legislation,

effective fiscal year (FY) 2007. In order for grantees under Parts A, B, and C to be exempted from the 75 percent core medical services requirement, they must request and receive a waiver from HRSA, as required in the Act.

Beginning in FY 2008, HRSA will utilize new standards for granting waivers of the core medical services requirement for the Ryan White HIV/AIDS Program. These standards meet the intent of the Ryan White HIV/AIDS Treatment Modernization Act of 2006 to increase access to core medical services, including antiretroviral drugs, for persons with HIV/AIDS and to ensure that grantees receiving waivers demonstrate the availability of such services for individuals with HIV/AIDS identified and eligible under Title XXVI of the PHS Act. The core medical services waiver uniform standard and waiver request process will apply to Ryan White HIV/AIDS Program grant awards under Parts A, B, and C of Title XXVI of the PHS Act. Core medical services waivers will be effective for a one-year period consistent with the grant award period.

Grantees must submit a waiver request with the annual grant application containing the certifications and documentation which will be utilized by HRSA in making determinations regarding waiver requests.

Grantees must provide evidence that all of the core medical services listed in the statute, regardless of whether such services are funded by the Ryan White

HIV/AIDS Program, are available to all individuals with HIV/AIDS identified and eligible under Title XXVI of the

PHS Act in the service area within 30 days.

The estimated annual burden is as follows:

Application	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Waiver Request .....	20	1	20	6.5	130
Total .....	20	.....	20	.....	130

Send comments to Susan G. Queen, Ph.D., HRSA Reports Clearance Officer, Room 10-33, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: October 18, 2007.

**Alexandra Huttinger,**

Acting Director, Division of Policy Review and Coordination.

[FR Doc. E7-20945 Filed 10-23-07; 8:45 am]

BILLING CODE 4165-15-P

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Proposed Collection; Comment Request; NICHD Research Partner Satisfaction Surveys**

**SUMMARY:** Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute of Child Health and Human Development (NICHD), the National Institutes of Health (NIH), has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. The proposed information collection was previously published in the **Federal Register** on July 25, 2007, in Volume 72, No. 142, pages 40887-40888, and allowed 60 days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The NIH may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented after October 1, 1995, unless it displays a currently valid OMB control number.

*Proposed Collection: Title:* NICHD Research Partner Satisfaction Surveys.

*Type of Information Collection Request:* Extension without change. *Need and Use of Information Collection:* Executive Order 12862 directs agencies that provide significant services directly to the public to survey customers to determine the kind and quality of services they want and their level of satisfaction with existing services. With this submission, the NICHD seeks to obtain OMB's generic approval to conduct customer satisfaction surveys surrounding its research programs and activities.

The NICHD was founded in 1963. Its mission is to ensure, through research, the birth of healthy infants and the opportunity for each to reach full potential in adulthood, unimpaired by physical or mental disabilities. The NICHD conducts and supports research on the many factors that protect and enhance the process of human growth and development. The developmental focus of the NICHD means that its research portfolio is unusually broad. NICHD programs include research on infant mortality, birth defects, learning disorders, developmental disabilities, vaccine development, and demographic and behavioral sciences, among others. In addition to supporting basic research, clinical trials, and epidemiological studies that explore health processes, the NICHD forms partnerships with organizations or institutions to ensure effective use of scientific findings and research products.

The NICHD utilizes strategic assessments to support Institute planning and policy development, and to help determine programmatic and scientific objectives and priorities. Research partner surveys will augment NICHD's ongoing efforts to assess research-related activities. The two principal objectives are: (1) To measure the personal satisfaction of research

partners with NICHD programs or initiatives, including both responsiveness to scientific aims and convenience of operations to support research and its effective use; and (2) to learn from research partners the ways in which the NICHD can improve the overall planning and management of its programs and initiatives. Findings will be used to improve NICHD's research programs and initiatives in the following ways: (1) To assess the effectiveness and efficiency of operations; (2) to identify opportunities for improving program performance; (3) to develop plans to incorporate innovations in program management; (4) to measure partner satisfaction and document program outcomes for governmental accountability reporting; and (5) to identify the need for creating new programs or initiatives or restructuring existing ones to respond to emerging scientific opportunities.

*Frequency of Response:* Annual [As needed on an ongoing and concurrent basis]. *Affected Public:* Members of the public, researchers, practitioners, and other health professionals. *Type of Respondents:* Members of the public; eligible grant applicants and actual applicants (both successful and unsuccessful); clinicians and other health professionals; and actual or potential clinical trials participants. The annual reporting burden is as follows: *Estimated Number of Respondents:* 28,000; *Estimated Number of Responses per Respondent:* 1; *Average Burden Hours per Response:* Varies with survey type, see below; and *Estimated Total Annual Burden Hours Requested:* 5,883. The annualized cost to respondents is estimated at: \$109,541.46. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

Type of respondents	Estimated number of respondents	Estimated number of responses per respondent	Average burden hours per response	Estimated total annual burden hours requested
Web-based .....	24,000	1	0.167	4,008.00
Telephone .....	2,000	1	0.50	1,000.00