

12A—ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Total	1,340	1,340	1,510

* The medical records sample selection instrument has been previously submitted as part of the RWHAP Outcomes Study proposed data collection project.

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.

Amy McNulty,

Acting Director, Division of the Executive Secretariat.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Advisory Committee on Rural Health and Human Services; Notice of Correction

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

ACTION: Notice; correction.

SUMMARY: The Health Resources and Services Administration published a notice in the *Federal Register*, FR 2017-23562 (October 31, 2017), announcing the charter renewal of the National Advisory Committee on Rural Health and Human Services (NACRHHS).

FOR FURTHER INFORMATION CONTACT: Paul Moore, Designated Federal Officer, NACRHHS, HRSA, 5600 Fishers Lane, Room 17W41C, Rockville, Maryland 20857, telephone (301) 443-0835, fax (301) 443-2803 or by email at pmoore2@hrsa.gov.

Correction

In the *Federal Register*, FR 2017-23562 (October 31, 2017), please make the following correction:

In the Summary section, correct to read: The effective date of the renewed charter is October 29, 2017.

Amy McNulty,

Acting Director, Division of Executive Secretariat.

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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; Be The Match® Patient Services Survey, OMB No. 0906-0004—Revision

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

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, 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 no later than December 13, 2017.

ADDRESSES: Submit your comments, including the ICR Title, to the desk officer for HRSA, either by email to 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 Lisa Wright-Solomon, the HRSA Information Collection Clearance Officer at paperwork@hrsa.gov or call (301) 443-1984.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: Be The Match® Patient Services Survey.

OMB No.: 0906-0004—Revision.

Abstract: The National Marrow Donor Program®/Be The Match® is a HRSA contractor dedicated to helping patients and families get the support and information they need to learn about their disease and treatment options, prepare for a blood stem cell transplant, and thrive after a transplant procedure. The information and resources provided help individuals navigate the bone marrow or cord blood transplant process. Participant feedback is essential to understand the needs for transplant support services and educational information across a diverse population. This information is used to determine the helpfulness of existing services and resources. Feedback is also used to identify areas for improvement and develop future programs.

Need and Proposed Use of the Information: Barriers to access to bone marrow or cord blood transplant related care and educational information are multi-factorial. Feedback from participants is essential to understand the changing needs for services and information as well as to demonstrate the effectiveness of existing services. The primary use for information gathered through the survey is to determine helpfulness of participants' initial contact with Be The Match® Patient Services Coordinators (PSC) and to identify areas for improvement in the delivery of services. In addition, stakeholders use this evaluation data to make program and resource allocation decisions.

The survey includes the following items to measure: (1) Reason for contacting Be The Match®, (2) if the PSC was able to answer questions and easy to understand, (3) if the contact helped the participant to feel better prepared to discuss transplant with their care team, (4) increase in awareness of available resources, (5) timeliness of response, and (6) overall satisfaction.

Proposed changes to the survey instrument include updated references to the survey title and staff titles. Changes to the questions include minor changes to question one, changes to the instructions for questions three and four, and minor rewording of question