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To: [HRSA Paperwork](#)
Subject: [EXTERNAL] National Marrow Donor Program Patient Support Center Survey
Date: Wednesday, July 1, 2026 5:33:11 PM

This email is in regards to the notice on [federalregister.gov](https://www.federalregister.gov) titled:

Agency Information Collection Activities: Proposed Collection: Public Comment Request; Information Collection Request Title: National Marrow Donor Program Patient Support Center Survey

I hope you find these comments helpful. I have a family member that donated a kidney, and my comments are based on an interview I did with him.

Ways to enhance the quality, utility, and clarity of the information to be collected

This commentary applies if the data collection is similar for any organ donors. The data being collected was intuitive, however, the format used for my family member was a paper questionnaire that was sent every three months for multiple years. The frequency wasn't a big problem, however, the questionnaire needed to be completed in its entirety every time. It would be better if the information was pre-loaded based on the prior submission, and then the donor could just update the items that have changed.

Thanks for your time.

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