

Most heart transplant centers' websites get failing grade when it comes to clear, accessible information

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In a University of Michigan-led study, researchers have found that most websites for heart transplant centers in the United States are difficult to



understand, with more than 40% lacking information in languages other than English.

The majority of the 139 websites analyzed were at a college junior's reading level, much higher than the average American's 8th-grade reading level. The study is published in the *Journal of Cardiac Failure*.

"There is a huge problem and a disconnect between physicians and patients regarding what we know and do in the hospital versus what we bring to patients and evaluating what they actually need," said first author James W. Stewart II, M.D., MSc., a national clinician scholar with the University of Michigan Institute for Healthcare Policy & Innovation and a general surgery resident at the Yale School of Medicine. "A large part of the problem is patients not properly understanding the information—and an even bigger problem is that we're not communicating and giving patients the resources to properly understand."

Just over 56% of the heart transplant centers' websites offered resources for non-English speakers. The mid-Atlantic region performed the worst, with only six of 16 heart transplant centers in Pennsylvania, New Jersey, West Virginia, Maryland, Delaware and Washington, D.C., providing online information in a language other than English.

Since 1980, the number of people speaking a language other than English at home has increased by almost 200%. Spanish (including Spanish Creole) is the most common <u>language</u> spoken after English.

Stewart believes that professional transplant societies along with the United Network for Organ Sharing, the nonprofit that manages the American organ transplant system, should create a common source of information at a reading level that patients can understand, with professionally translated information about heart transplants.



Until then, he says, <u>heart</u> transplant centers need to take it upon themselves to make sure their information is accessible.

"During transplant meetings, evaluate what information on the website is important for patients to know and review the site every six months or so," he suggested. "Then ask patients questions like, 'Did you go to the website? Did you find it useful? What did you find useful? Is this something you understand?' Following through on whether this information is helpful and involving patients more is critical."

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More information: James W. Stewart et al, Readability and Non-English Language Resources of Heart Transplant Center Websites in the United States, *Journal of Cardiac Failure* (2022). DOI: 10.1016/j.cardfail.2022.09.016

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