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Commentary: Is there life after cardiac death? Considering the challenges of heart donation after circulatory death

Alexander Raskin, MD,^a Farhan Zafar, MD,^b and David L. S. Morales, MD^b

Rajab and colleagues¹ provide an excellent synopsis on the current state of cardiac donation after circulatory death (DCD). The authors highlight a key issue of more patients being listed than are being transplanted each year. As a result, managing and minimizing waitlist mortality has become paramount. Increased use of DCD has been proposed as a means to increase the donor pool by as much as 15% to 30%, but this may be an overestimation. In the modern era, DCD continues to be minimally employed, with just more than 100 adult cases reported, almost all outside the United States.³ Several barriers may stand in the way of wider adoption of DCD, which is mired with legal and ethical considerations. The legal pronouncement of cardiac death revolves around the irreversible cessation of cardiopulmonary function. However, the required resuscitation and restoration of cardiac activity needed to carry out DCD transplantation may violate the tenant of irreversibility. Another consideration is that the legal definition of death may vary between jurisdictions,⁴ and this can make it harder to standardize widely adaptable DCD protocols. For example, some approaches call for ante-mortem placement of extracorporeal membrane oxygenation cannulas, which has been deemed unacceptable in certain institutions and jurisdictions. Variability in mandated standoff time, the time between circulatory arrest, and declaration of death may further alter warm ischemic time and graft quality. These

From the Divisions of ^aPediatric Cardiology, and ^bPediatric Cardiothoracic Surgery, Cincinnati Children's Hospital Medical Center, Cincinnati, Ohio.

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Alexander Raskin, MD, Farhan Zafar, MD, and David L. S. Morales, MD

CENTRAL MESSAGE

Cardiac donation after circulatory death may not be the solution to the currently evergrowing waitlist.

considerations are only amplified when trying to employ the DCD strategy internationally.

Data from adult DCD recipients have demonstrated greater rates of immediate graft dysfunction and increased extracorporeal membrane oxygenation use. Intermediate survival seems to be equivocal for adult DCD recipients, but the overall reported numbers are limited.⁵ Pediatric DCD outcomes, based on 21 reported cases to the International Society for Heart and Lung Transplantation Registry, are less encouraging. According to that data, DCD recipients had increased graft dysfunction and decreased 1-year survival of 61% versus 91% compared with donation after brain death.6 Recipient selection factors could have contributed to the survival outcomes. Unfortunately, donor characteristics were not reported in that series. These results may explain why DCD accounted for 0.5% of pediatric and less than 1% of adult transplants over the last 2 decades.^{6,7}

Perhaps we should maximize our current pool of donors, since up to 44% of pediatric and 48% of adult available allografts are discarded. Registry data from the Organ Procurement and Transplantation Network demonstrate that donor use has declined with time. Despite seemingly more stringent donor selection, recipient survival has not improved significantly. Many of the discarded hearts would have likely resulted in favorable clinical outcomes. In fact, adults who received adolescent allografts that were initially rejected by all pediatric centers went on to have excellent results. At this time, it is unclear how many of the unselected allografts would be appropriate for donation, but clearly some would be. The provincial protocols and beliefs about accepting organs that vary at each program have led to our inability to maximize the donor

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pool. More work using machine learning needs to be done to better understand and standardize the donor selection process while minimizing existing variability. We should be able to improve donor selection and use without adversely impacting recipient survival, in fact, improving it for the community as a whole. Perhaps there is life after cardiac death via donation, but we do not believe it will flourish in its current form or in the present system, as the past decade has revealed. Perhaps, the use of technology like the TransMedics Organ Care System (TransMedics, Andover, Mass) to resuscitate and evaluate the organs after cardiac death will bring life to DCD. Results from the EXPAND trial will be informative in this regard.

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