

# GORE® PRECLUDE® PERICARDIAL MEMBRANE CAN FACILITATE REOPERATIONS IN LEFT VENTRICULAR ASSIST DEVICE (LVAD) PROCEDURES<sup>1-4</sup>



Clinical studies/reports utilizing GORE® PRECLUDE® Pericardial Membrane in LVAD and total artificial heart (TAH) surgeries found

**100%** of patients were free of severe adhesion at reoperation.<sup>1-5</sup>

**Reoperation patients implanted with an LVAD or TAH  
and GORE® PRECLUDE® Pericardial Membrane<sup>1-5</sup>** **N = 65**

Patients with severe adhesions	0 (0.0%)
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The quote was modified to include the current trademark of the GORE® PRECLUDE® Pericardial Membrane

“At reoperation, the polytetrafluoroethylene membrane [GORE® PRECLUDE® Pericardial Membrane] was . . . clearly separate from the sternum and resembled a sheet of plastic covering both the heart and the pump conduit, made of knitted polyester material; no problematic adhesions were observed between the membrane, the chest wall, the extracardiac conduit, and the underlying epicardium. The [GORE® PRECLUDE® Pericardial Membrane] sheet wrapping the outflow cannula is very useful to accomplish a safe dissection of this conduit from the right atrium to quickly perform bypass.”<sup>1</sup>

\*Reprinted from *Operative Techniques in Thoracic and Cardiovascular Surgery*, Vol:19, Osami Honjo, Vivek Rao, Implantation of HeartWare Left Ventricular Assist Device in Pediatric Population, Figure 8a, page 93, 2014, with permission from Elsevier. Image courtesy Osami Honjo, M.D., Ph.D. Used with permission.

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# References

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Consult Instructions  
for Use

[eifu.goremedical.com](http://eifu.goremedical.com)

Refer to *Instructions for Use* at [eifu.goremedical.com](http://eifu.goremedical.com) for a complete description of all applicable indications, warnings, precautions and contraindications for the markets where this product is available. REGD

Products listed may not be available in all markets.

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