MAGNA PERFORMANCE.
MITRAL EASE.

CARPENTIER-EDWARDS PERIMOUNT
MAGNA MITRAL EASE
PERICARDIAL BIOPROSTHESIS
Built on Proven Performance

- Outperforming mitral pericardial design in challenging and standard age groups\(^1\)

- Study after study, consistently demonstrates long-term endurance of Edwards mitral pericardial design\(^1\)-\(^7\)

\([\text{Graph showing actuarial freedom from structural valve deterioration, with details on study results below.}]\)

\(^{**}\)Patients and results are a subset of each study. Reporting methods vary among literature sources. See references for definitions.
No clinical data are available which evaluate the long-term impact of the Carpentier-Edwards ThermaFix process in patients.

Long, multi-bend handle enhances reachability enhanced Mitral implantability where MagNa performance meets MiTral ease of implant.

The Magna Mitral Ease valve offers outstanding performance and durability. With the addition of significant design enhancements for better access, placement, and suturing, Magna Mitral Ease provides performance you can count on to ease the challenges of mitral valve replacement.

**Enhanced Mitral Implantability**

- Color-coded valve holder confirms implant readiness
- Asymmetric design to position the posterior strut away from LV wall
- Supra-annular position to reduce the risk of LVOTO and ventricular injury
- Ultra-low profile valve optimizes outflow hemodynamics and physiologic flow
Pre-attached barrel and replica sizers facilitate valve sizing

Model 7300TFX
Nominal Specifications (mm)

<table>
<thead>
<tr>
<th>Size</th>
<th>25</th>
<th>27</th>
<th>29</th>
<th>31</th>
<th>33</th>
</tr>
</thead>
<tbody>
<tr>
<td>A. Stent Diameter (Wireform)</td>
<td>25</td>
<td>27</td>
<td>29</td>
<td>31</td>
<td>31</td>
</tr>
<tr>
<td>B. Tissue Annulus Diameter</td>
<td>28</td>
<td>29.5</td>
<td>31.5</td>
<td>33.5</td>
<td>33.5</td>
</tr>
<tr>
<td>C. External Sewing Ring Diameter</td>
<td>36</td>
<td>38</td>
<td>40</td>
<td>42</td>
<td>44</td>
</tr>
<tr>
<td>D. Anterior Effective Profile</td>
<td>7</td>
<td>7.5</td>
<td>8</td>
<td>8.5</td>
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</tr>
</tbody>
</table>

References

Brief Summary—Mitral Bioprosthesis
Indications: For use in patients whose mitral valvular disease warrants replacement of their natural or previously placed prosthetic valve and when the valve cannot be repaired. Contraindications: Do not use if surgeon believes it would be contrary to the patient's best interests. Complications and Side Effects: stenosis, regurgitation, endocarditis, hemolysis, thromboembolism, valve thrombosis, nonstructural dysfunction, structural valve deterioration, anemia, arrhythmia, hemorrhage, transient ischemic attack/stroke, congestive heart failure, myocardial infarction, angina, ventricular perforation by stent posts, any of which could lead to reoperation, explantation, permanent disability and death. Warnings: Alternative therapies should be considered in the presence of conditions affecting calcium metabolism or when calcium containing chronic drug therapies are used, including children, adolescents, young adults and patients on a high calcium diet or maintenance hemodialysis. Should be used with caution in the presence of severe systemic hypertension or when anticipated patient longevity is longer than the known longevity of the prosthesis.

For professional use. CAUTION: Federal (United States) law restricts this device to sale by or on the order of a physician. See instructions for use for full prescribing information, including indications, contraindications, warnings, precautions and adverse events.

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