

PERCEVAL® PLUS THE OPTIMAL MIX



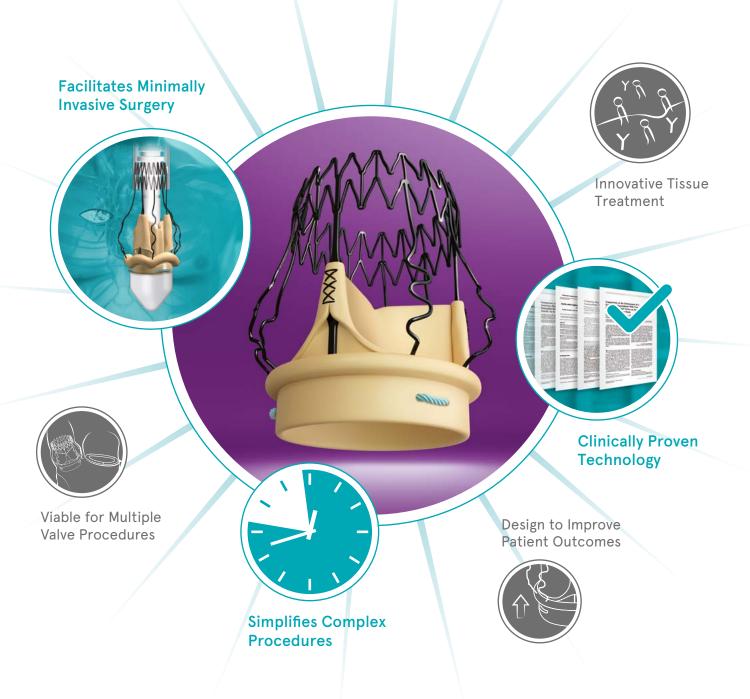
PERCEVAL PLUS: DESIGNED TO PERFORM.

PERCEVAL® PLATFORM

- First-in-human performed in 2007
- More than 10 years
 of successful clinical use
 - Solid clinical evidence
 - Truly global reach

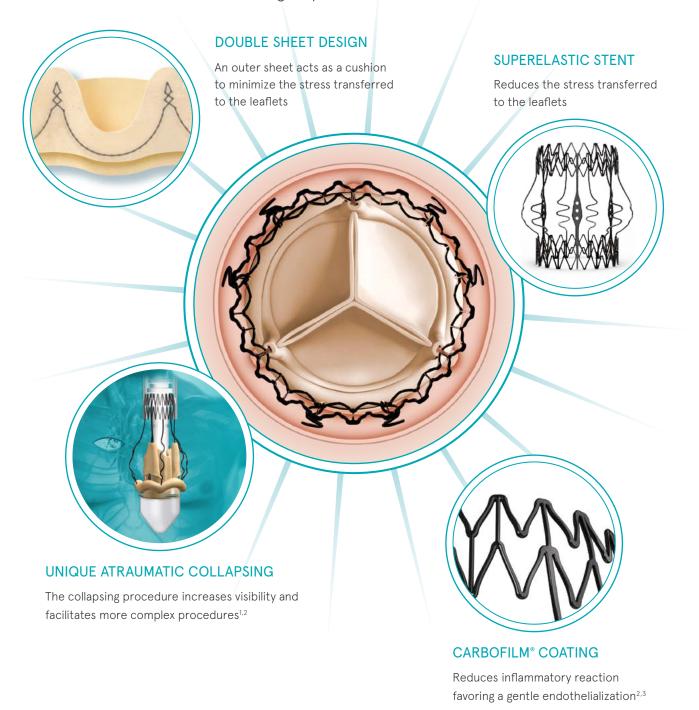
Designed to perform

Unique Valve Design



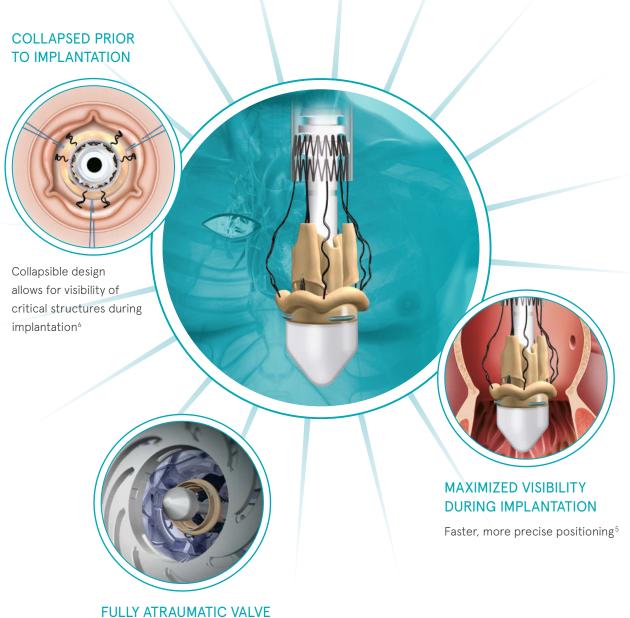
Unique Valve Design

LivaNova's tissue platform has a long clinical heritage, its proven double-sheet design principle has been used to manufacture biological prostheses since 1985.



Facilitates Minimally Invasive Surgery

The atraumatic collapsing does not impair leaflet functionality⁴, preventing possible damage to the tissue. The reduced profile allows the surgeon better visibility of the annulus and the anatomical structures during implantation and deployment for greater confidence and faster, more precise positioning at the implantation site.⁵

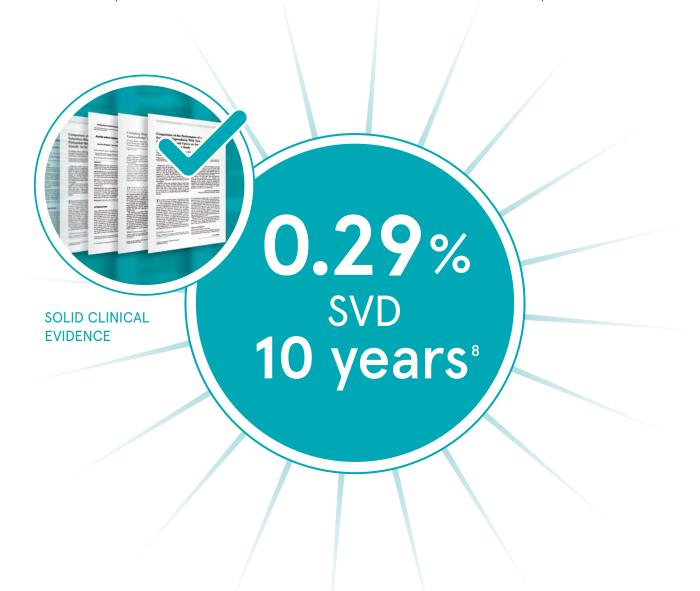


FULLY ATRAUMATIC VALVE COLLAPSING

Does not impair leaflet functionality⁷

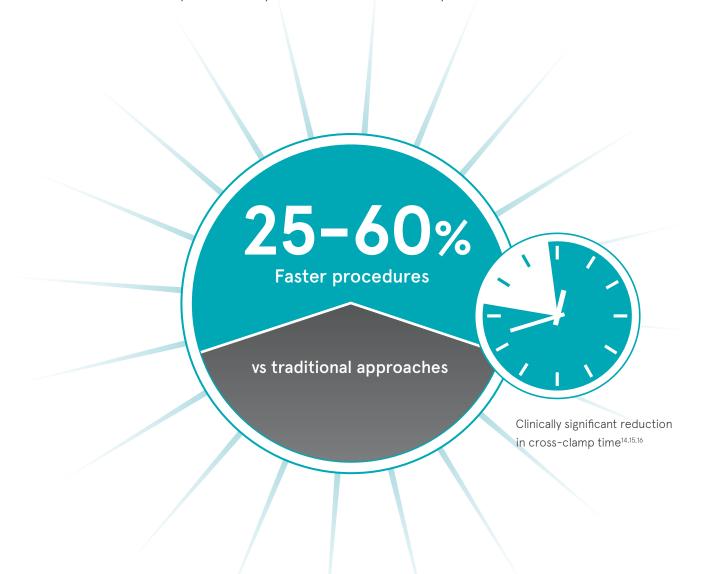
Clinically Proven Technology

PERCEVAL PLUS is based on the trusted PERCEVAL platform supported by more than 10 years of clinical experience that evidence excellent results in terms of durability.⁸



Simplifies Complex Procedures

Whether using a full sternotomy or a MICS approach, PERCEVAL PLUS enables a faster procedure^{9,10} and reduced cross-clamp¹¹ time compared to traditional valve prostheses.



PERCEVAL PLUS helps reduce complexity even in challenging and time consuming procedures, 12,15 while maintaining good hemodynamic outcomes. 11,13

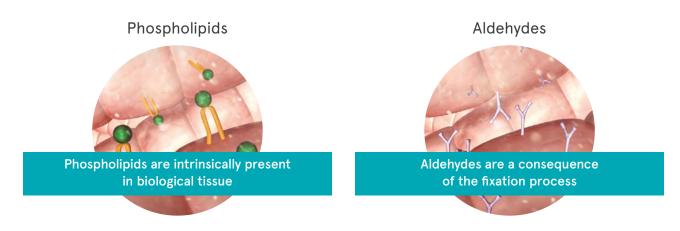
PERCEVAL PLUS is safe and effective, even in cases of concomitant cardiac procedures.¹⁰

Innovative Tissue Treatment Designed to Reduce Calcification

LivaNova's commitment to achieve the ideal tissue platform has led to constant improvements in tissue treatments, from the initial fixation process to the aldehyde neutralization treatment and now to the FREE tissue treatment that addresses both phospholipids and aldehydes.

The major causes of calcification

Phospholipids and aldehydes contribute to calcification as they are calcium binding sites.¹⁷⁻²¹



FREE addresses calcium binding sites



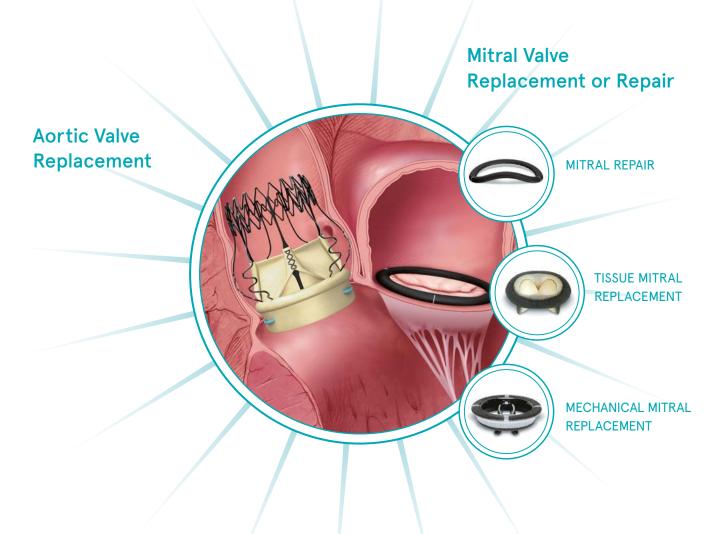
Supporting evidence of the FREE tissue treatment

-96% phospholipid content*

ZERO aldehyde toxicity

Viable for Multiple Valve Procedures When time really matters

The unique characteristics of PERCEVAL PLUS make it an ideal choice in complex implantations while extending the benefits to a wider patient population.*



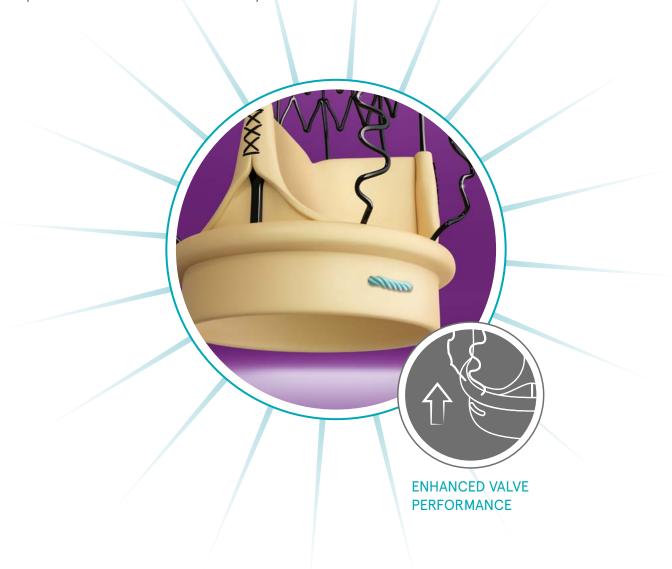
- Different clinical evidences already published to support the benefit of PERCEVAL PLUS in concomitant procedures. 23-30
- Concomitant PERCEVAL PLUS implantation and mitral valve repair/replacement is a feasible, safe and reproducible procedure.^{24,25,27,28,30}
- Minimally invasive surgery in combined procedures is a feasible approach that could be simplified with the use of PERCEVAL PLUS.²⁹

^{*} The decision of using PERCEVAL PLUS in the patients should be based on a careful individual assessment and limited to cases in which the benefits of using PERCEVAL PLUS justify the risks.

The available clinical data indicate that using PERCEVAL PLUS in patients with other prostheses may result in intraoperative valve misplacement or insufficient leaflet coaptation leading to valve replacement, due to possible interference with the other prostheses.

Designed to Improve Patient Outcomes

Implantation depth and therefore the stent prosthesis protrusion into the LVOT is a measure strongly associated with post-operative conduction disturbances.³¹ Therefore the reduction of the protrusion of the valve below the aortic annulus is expected to decrease the risk of impairment of the AV conduction system.







BECAUSE PATIENTS WISH TO RETURN TO A FULL AND ACTIVE LIFE



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INDICATIONS: The PERCEVAL PLUS prosthesis is indicated for the replacement of diseased native or a malfunctioning prosthetic aortic valve via open heart surgery. The prosthesis is indicated for use in adult patients who are diagnosed to have aortic valve stenosis or steno-insufficiency.

TOP POTENTIAL SIDE EFFECTS: The risks or potential adverse events associated with cardiac valve replacement with a bioprosthesis include, but may not be limited to: cardiac arrhythmias, death, endocarditis,

TOP POTENTIAL SIDE EFFECTs: The risks or potential adverse events associated with cardiac valve replacement with a bioprosthesis include, but may not be limited to: cardiac arrhythmias, death, endocarditis, heart failure, hemorrhage, intravalvular and/or paravalvular leak, stroke or any related neurologic disorders, structural valve deterioration, reoperation and explant. Beyond the previously mentioned adverse events, specific events related to the implant of the PERCEVAL PLUS prosthesis may include, but not be limited to dislodgment and/or migration of the prosthesis.

MRI conditional.

Liva Nova

Health innovation that matters

Manufactured by:

Sorin Group Italia Srl

A wholly-owned subsidiary of LivaNova PLC Via Crescentino - 13040 Saluggia (VC) Italy Tel: +39 0161 487472 - Fax: +39 0161 487316 info.cardiacsurgery@livanova.com

www.livanova.com





DIVISIÓN CIRUGÍA CARDÍACA

Can Sant Joan Business Park · Jesús Serra Santamans, 5 08174 Sant Cugat del Vallès (Barcelona) T: +34 934 006 552 F: +34 934 006 501 palexmedical@palex.es



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