

EuroValve

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Early and mid-term outcomes after aortic valve replacement with a sutureless prosthesis: single center experience.

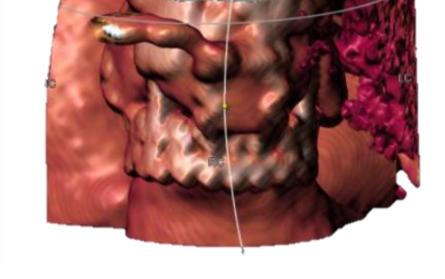
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AIM

We evaluated the outcomes of 68 consecutive patients undergone AVR with the sutureless Perceval aortic valve bioprosthesis (Sorin Group, Saluggia, Italy) from April 2011 to June 2014.





Perceval valve

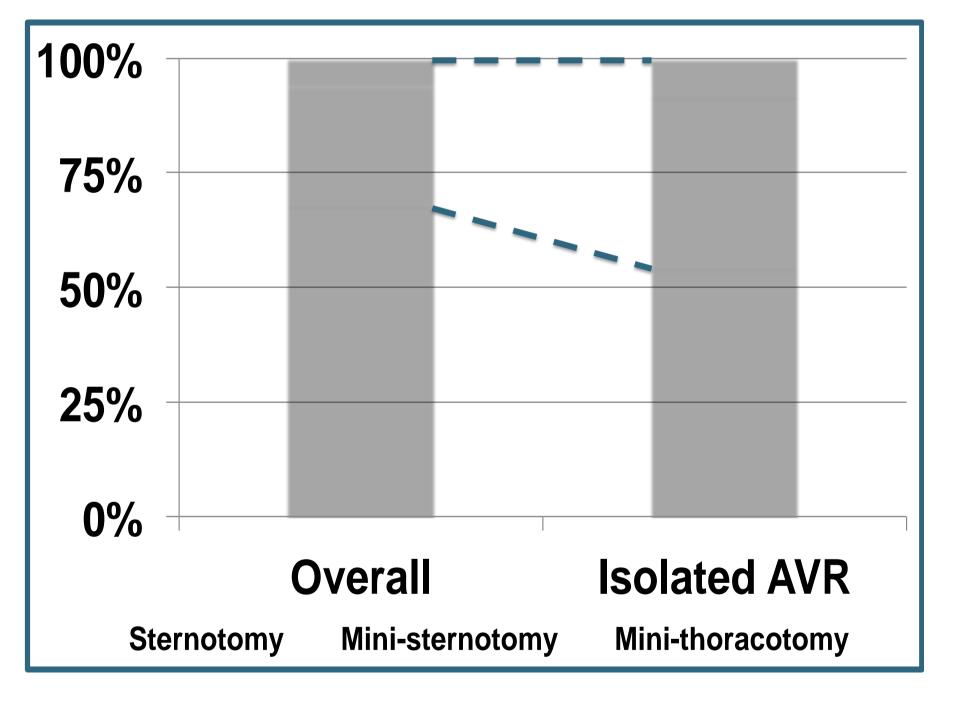
3D renderiing of the valve in the aortic box

RESULTS

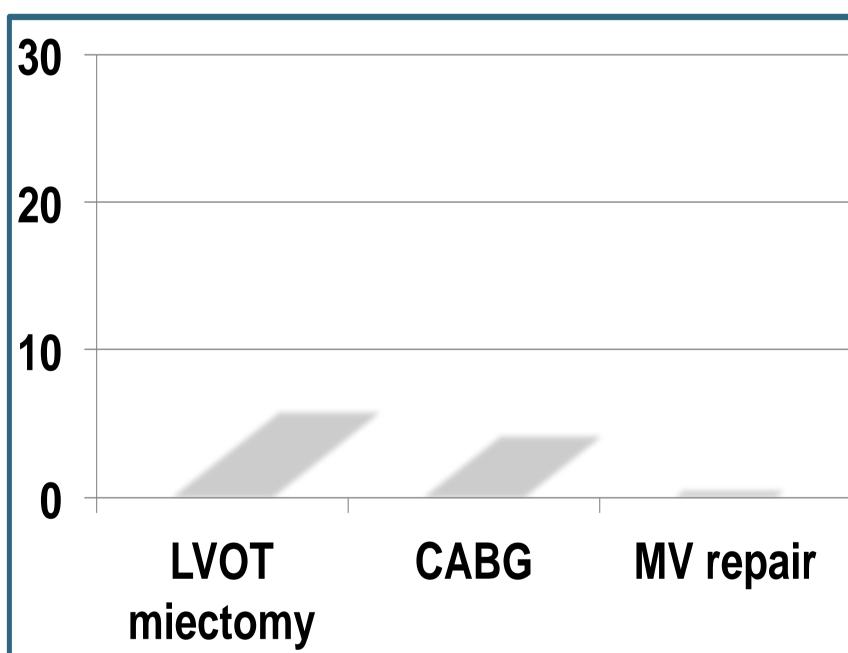
Preoperative variables

Variables	n=68	Variables	n=68
Age	74.8±5.4 yrs (58-92 yrs)	COPD	16 (23.5%)
Female	45 (66.2%)	EuroSCORE II	5.5±5.3% (1.1%-45.1%)
NYHA(3/4)	58 (85.3%)	EF	56.1±9.1% (25%-76%)
IDDM	19 (27.9%)	Annulus	19.5±6.8 mm (17-26 mm)

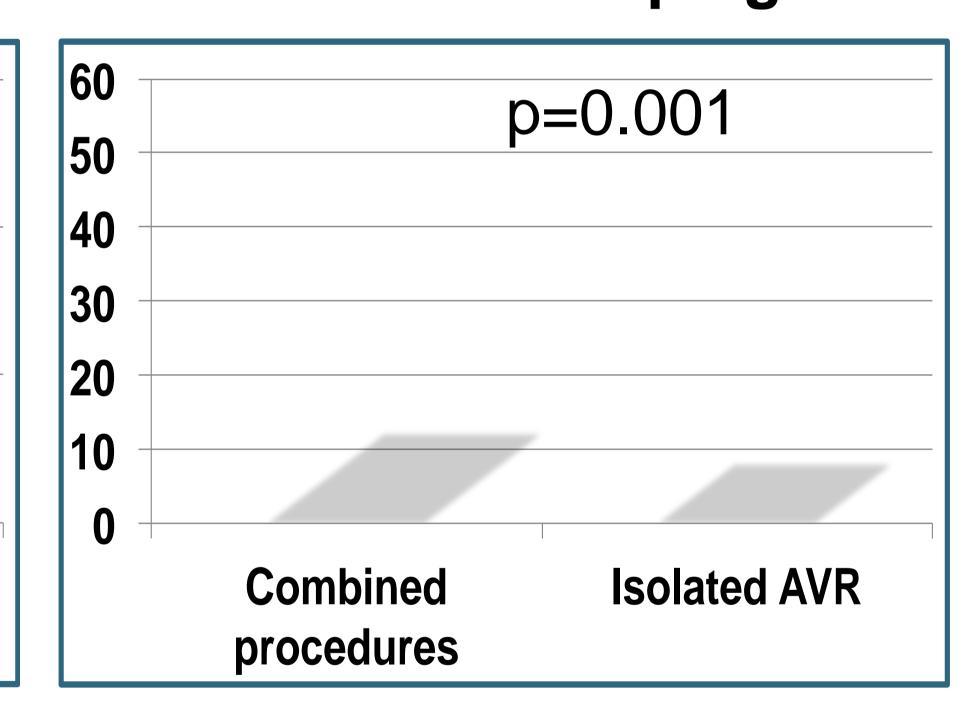
Surgical approach



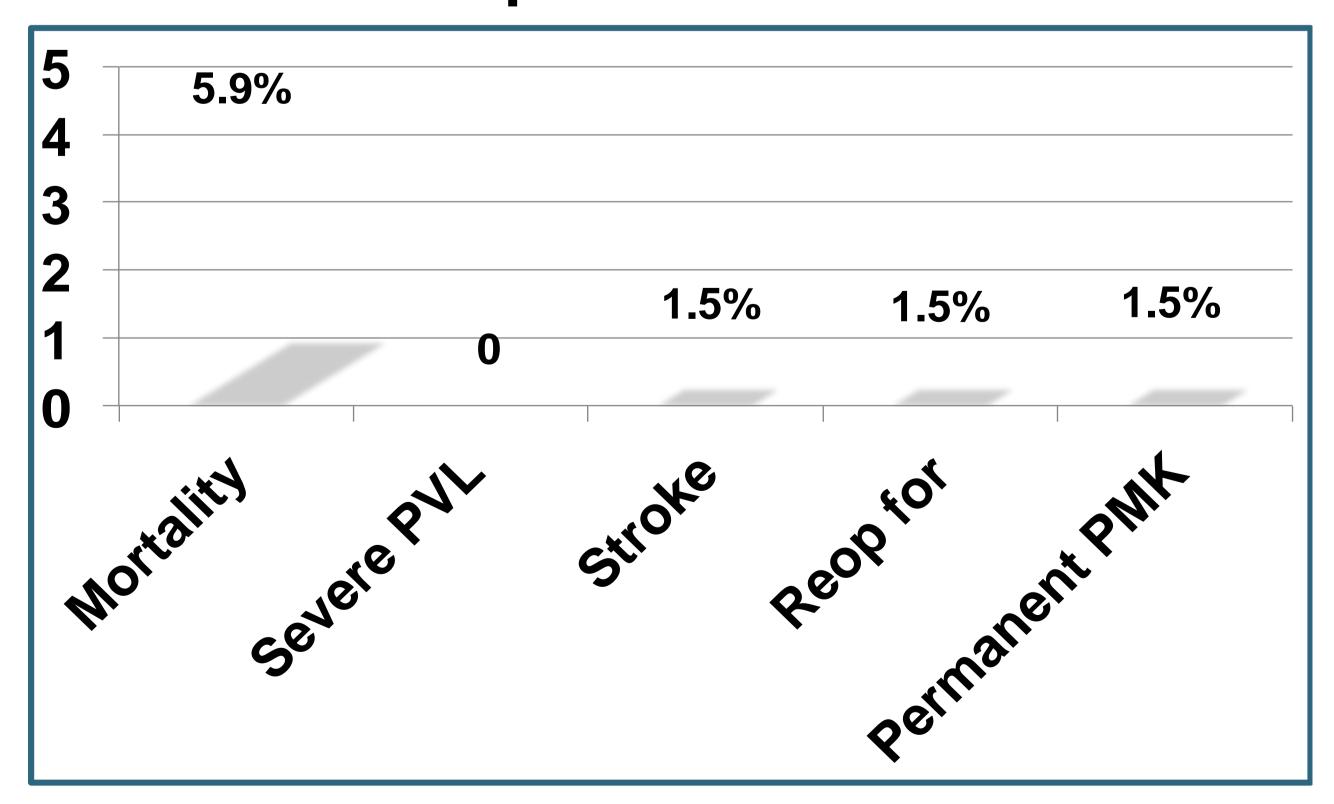
Associated procedures



Aortic cross-clamping time

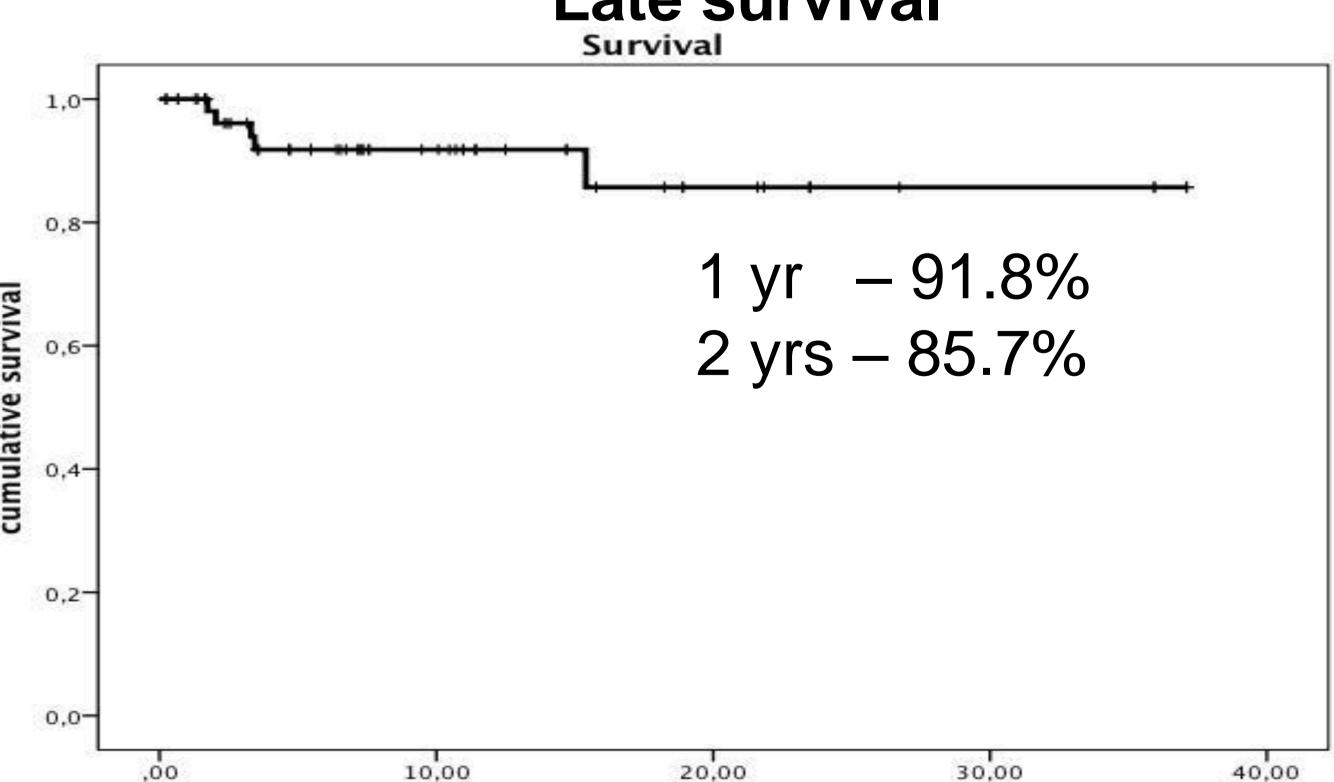


Hospital outcomes



100% device success

Late survival



100% freedom from valve-related mortality, stroke, redo, endocarditis

Months of follow-up

CONCLUSIONS

Perceval allows to minimize cross-clamping time and delivers excellent clinical outcome. Furthermore, this valve offers the unique advantage to implant a stentless valve with a sutureless technology, shifting the concept of minimally invasive surgery from small incisions to the minimal manipulation of the heart.