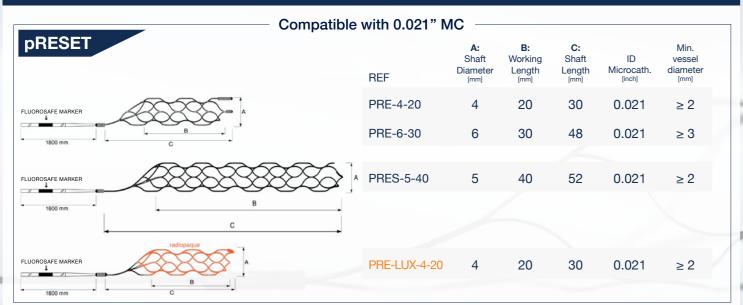
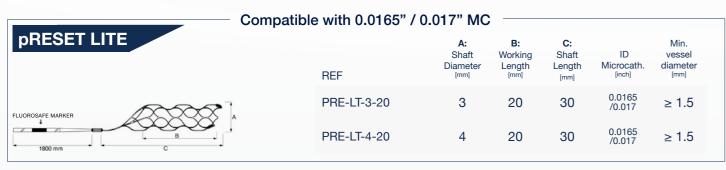




Specifications

When crossing profile is critical, the choice is yours







Continuous commitment to patient care - the extended **pRESET** family

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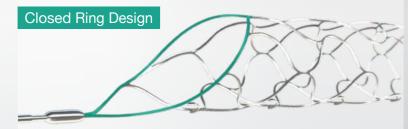
PRESETThrombectomy Device

Proven scientific excellence

Backed by clinical evidence





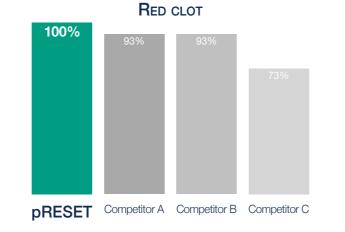


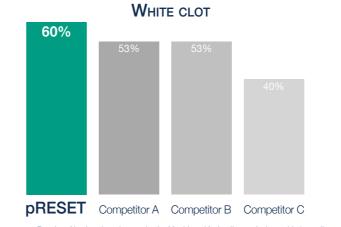
Unique design elements

- Helical slit maintains cell shape integrity independent of expansion diameter
- Closed Ring Design ensures stable opening and constant wall apposition during retrieval
- Dual Type Cell Design for deep clot integration combined with flexibility in tortuous anatomies



Best-in-class clot retention and removal of red and white clot





Results of in vitro thrombectomies by Machi et al.2; Applies to devices with 6mm diameter

ARTESp¹ study conclusion

- safety and efficacy of mechanical thrombectomy with pRESET
- excellent recanalization rate
- excellent long-term neurological outcome regardless of patient's age



TICI 2b/3 **84,4%**

	ARTESp	MR CLEAN	SWIFT-PRIME	EXTEND-IA	ESCAPE	
nRS 0-2 90 days	62,5%	32,6%	60%	71%	53%	
TCI 2b/3	84,4%	58,7%	88%	86%	73,4%	

- 1 Prothmann S et al.; Acute Recanalization of Thrombo-Embolic Ischemic Stroke with pRESET (ARTESp): the impact of occlusion time on clinical outcome of directly admitted and transferred patients; J NeuroIntervent Surg 2016; doi:10.1136/neurintsurg-2016-012556.
- 2 Machi P et al.; Experimental Evaluation of Stent Retrievers' Mechanical Properties and Effectiveness; J NeurovIntervent Surg 2016; doi: 10.1136/neurintsurg-2015-012213. Applies for pRESET 6-30.
- 3 Lamprecht S et al.; Penetration Depth of Stent Retrievers Into Clots is Highly Dependent on Stent Design; ASNR 2017. Submitted for publication. Applies for pRESET LITE 4-20.

The pRESET, pRESET LITE and pRESET LUX Thrombectomy Devices have received the CE Mark (CE 0297)
They are not approved for sale nor are they available for sale or use in the United States