hybrid

the ideal combination

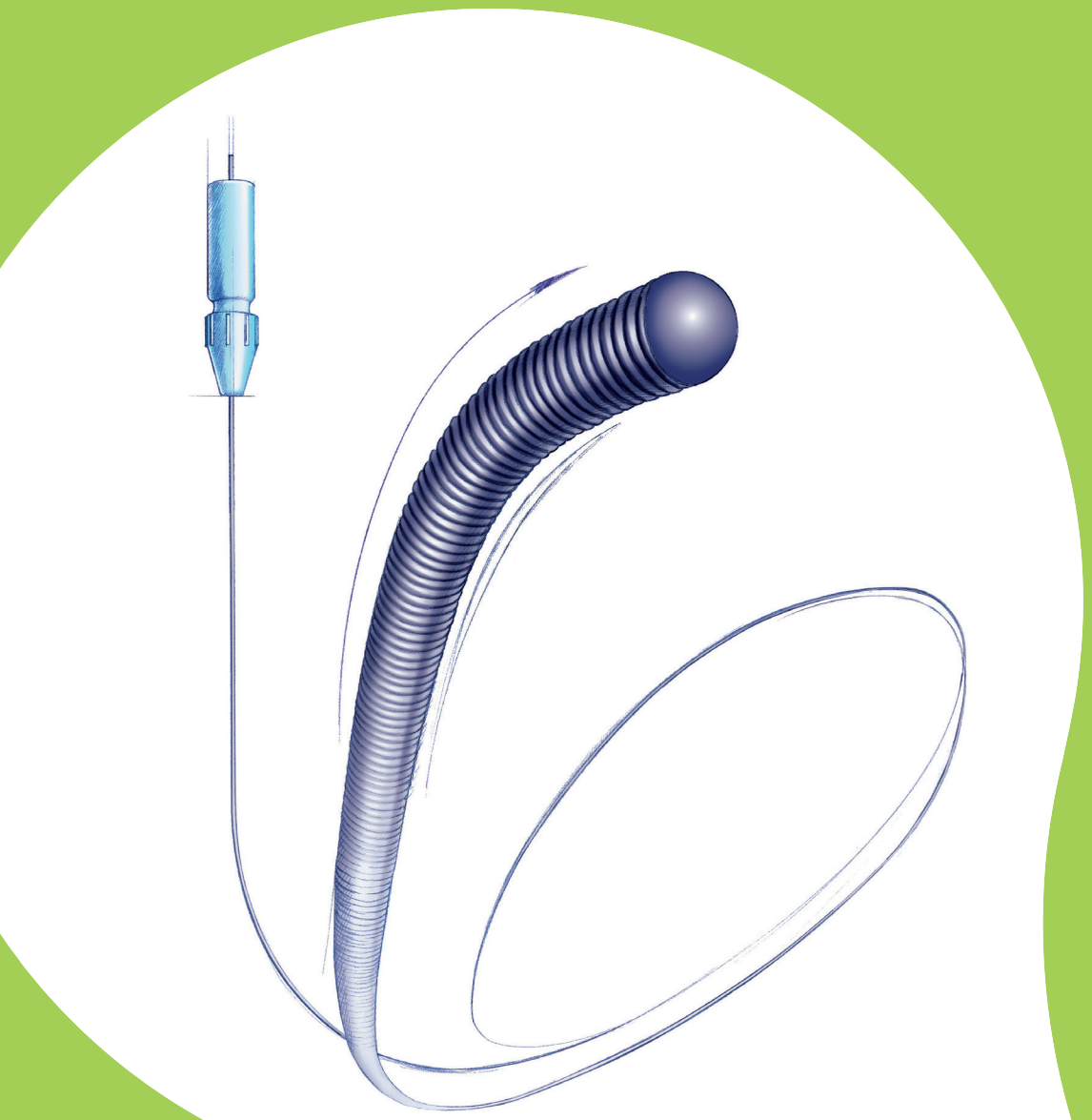
balt

inspiring innovation

hybrid

Hydrophilic guidewire

Designed to facilitate the insertion of catheters into intracranial vascular branches for diagnostic or therapeutic use.



O.D. as small as .007"

access treatment

Optimized navigation

Trackability and pushability
given by the stainless steel proximal part

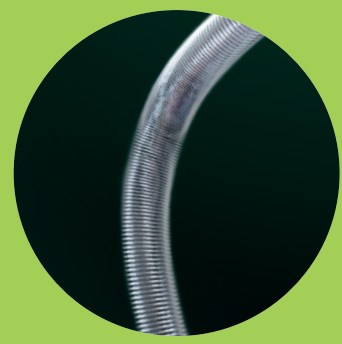
Shape retention and suppleness
thanks to the distal part in Nitinol

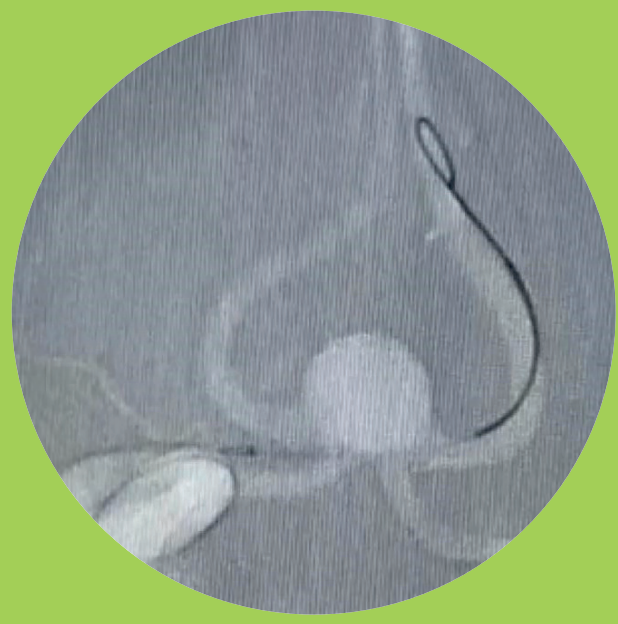
Smooth navigability
enhanced by the hydrophilic coating

Visibility
ensured by the distal radiopaque coil tip

Access in tortuous vasculature

thanks to the double angle shaped tip and broad options of O.D. as small as .007"

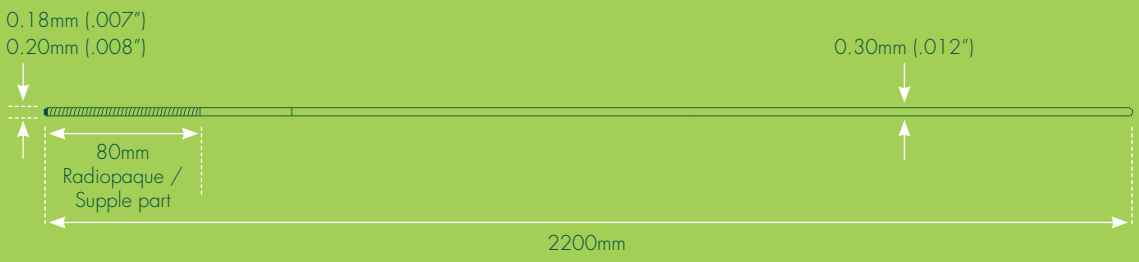




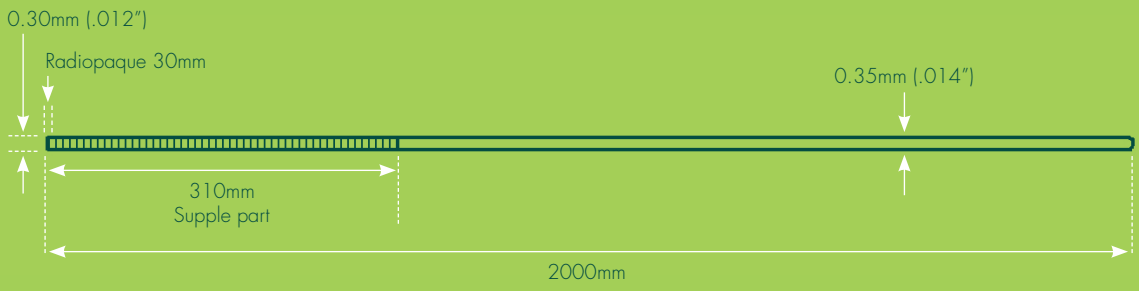
radiopaque coil tip

available in multiple lengths

HYBRID007 & HYBRID008D



HYBRID1214D



ordering information

Developed portfolio with unique references

Hybrid

Reference	Distal O.D.	Length (cm)	Compatible with	Curve
HYBRID007D	.007" (0,18mm)	220	SONIC all references MAGIC all references BALTACCI all references & others*	Straight (D)
HYBRID007J				Double Angle (J)
HYBRID008D	.008" (0,20mm)	220	SONIC all references except 1.2F MAGIC all references except 1.2F BALTACCI all references except 1.2F & others*	Straight (D)
HYBRID008J				Double Angle (J)
HYBRID007D.120	.007" (0,18mm)	120	Shorter length	Straight (D)
HYBRID1214D	.012" (0,30mm)	200	Any microcatheter compatible with .014" guidewire*	Straight (D)
HYBRID1214DA				Double Angle (DA)
HYBRID014D	.014" (0,35mm)	200	Any microcatheter compatible with .014" guidewire* COPERNIC RC	Straight (D)
HYBRID10D300	.010" (0,25mm)	310	Exchange microguidewires	Straight (D)
HYBRID12D300	.012" (0,30mm)			

*Check compatibility on products labelling

HYBRID are guidewires designed to facilitate the insertion of catheters into intracranial vascular branches for diagnostic or therapeutic use. Class III CE0297 in compliance with Medical Device Directive (MDD 93/42/EEC amended by 2007/47/EC). Manufactured by BALT Extrusion SAS. Carefully read the instructions for use before use. Not reimbursed. First CE marking:2010. SONIC are braided micro-catheters intended for selective and hyper selective vascular catheterization for diagnostic or therapeutic purposes. Class III CE0297 in compliance with Medical Device Directive (MDD 93/42/EEC amended by 2007/47/EC). Manufactured by BALT EXTRUSION SAS. Carefully read the instructions for use before use. Not reimbursed. First CE marking: 2005. BALTACCI are micro-catheters intended for selective and hyperselective vascular catheterization for diagnostic and therapeutic purposes. Class III CE0297 in compliance with Medical Device Directive (MDD 93/42/EEC amended by 2007/47/EC). Manufactured by BALT EXTRUSION SAS. Carefully read the instructions for use before use. Not reimbursed. First CE marking: 1998. MAGIC are micro-catheters intended for selective and hyperselective vascular catheterization for diagnostic and therapeutic purposes. Class III CE0297 in compliance with Medical Device Directive (MDD 93/42/EEC amended by 2007/47/EC). Manufactured by BALT EXTRUSION SAS. Carefully read the instructions for use before use. Not reimbursed. First CE marking: 1998. VASCO+ are reinforced micro-catheters intended: for injection of diagnostic or therapeutic products; to position pushable coils "SPIRALES" or detachable coils especially the ones of MDS « mechanical detachment system »; for the use of the self-expanding stent LEO+ or SILK+. Class III CE0297 in compliance with Medical Device Directive (MDD 93/42/EEC amended by 2007/47/EC). Manufactured by BALT EXTRUSION SAS. Carefully read the instructions for use before use. Not reimbursed. First CE marking: 2004. COPERNIC are occlusion catheters indicated for use in the neurovasculature and peripheral system to temporarily stop or control blood flow, to treat vasospasmes and embolization of aneurysms with balloons. Class III CE0297 in compliance with Medical Device Directive (MDD 93/42/EEC amended by 2007/47/EC). Manufactured by BALT EXTRUSION SAS. Carefully read the instructions for use before use. Not reimbursed. First CE marking: 1998. DC044GB (10/19)

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