

Ordering information

150 cm Catheter Length					
Order Code	Balloon diameter (mm)	Balloon length (mm)	Nominal pressure (atm)	Rated burst pressure (atm)	Sheath size (F)
U41501Q1HRX		15	6	16	4
U41501Q2RX	1.25	20	6	16	4
U41501Q4RX		40	6	16	4
U41501H2RX		20	6	16	4
U41501H4RX		40	6	16	4
U41501H8RX	1.5	80	6	16	4
U41501H12RX		120	6	16	4
U415022RX		20	6	16	4
U415024RX		40	6	16	4
U415028RX		80	6	15	4
U4150212RX	2	120	6	15	4
U4150215RX		150	6	15	4
U4150220RX		200	6	15	4
U4150230RX		300	6	15	4
U41502H2RX		20	6	16	4
U41502H4RX		40	6	16	4
U41502H8RX		80	6	15	4
U41502H12RX	2.5	120	6	15	4
U41502H20RX		200	6	15	4
U41502H30RX		300	6	15	4
U415032RX		20	6	16	4
U415034RX		40	6	15	4
U415038RX		80	6	15	4
U4150312RX	3	120	6	15	4
U4150315RX		150	6	15	4
U4150320RX		200	6	15	4
U4150330RX		300	6	15	4
U41503H4RX		40	6	16	4
U41503H8RX		80	6	16	4
U41503H12RX	3.5	120	6	15	4
U41503H20RX		200	6	15	4
U41503H30RX		300	6	15	4

150 cm Catheter Length					
Order Code	Balloon diameter (mm)	Balloon length (mm)	Nominal pressure (atm)	Rated burst pressure (atm)	Sheath size (F)
U415042RX		20	6	16	4
U415044RX		40	6	16	4
U415048RX		80	6	15	4
U4150412RX		120	6	15	4
U4150512RX	4	120	6	15	4
U4150415RX		150	6	15	4
U4150420RX		200	6	15	4
U4150430RX		300	6	15	4
U415054RX		40	6	14	5
U415058RX		80	6	13	5
U4150515RX	5	150	6	13	5
U4150520RX		200	6	13	5
U4150530RX		300	6	13	5
U415064RX	6	40	6	14	5
U415074RX	7	40	6	12	5

200 cm Catheter Length					
Order Code	Balloon diameter (mm)	Balloon length (mm)	Nominal pressure (atm)	Rated burst pressure (atm)	Sheath size (F)
U420024RX		40	6	16	4
U4200210RX	2	100	6	15	4
U42002H10RX		100	6	15	4
U42002H4RX		40	6	16	4
U4200310RX	2.5	100	6	15	4
U420034RX		40	6	16	4
U4200310RX	3	100	6	15	4
U4200320RX		200	6	15	4
U42003H4RX		40	6	16	4
U42003H10RX	3.5	100	6	15	4

Ultraverse™ RX PTA Dilatation Catheter

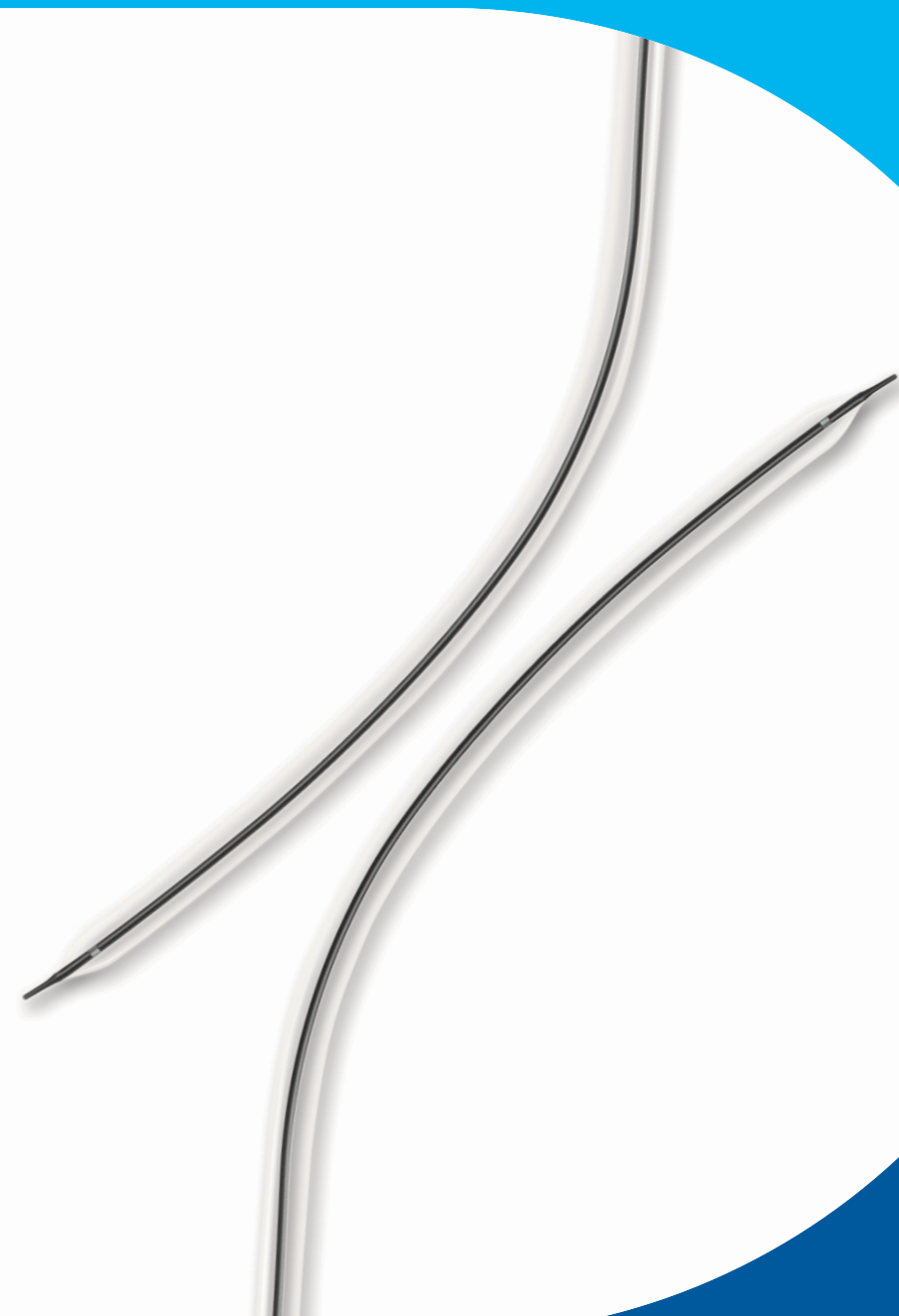
Indications for Use: ULTRVERSE™ RX PTA Dilatation Catheter is indicated for use in Percutaneous Transluminal Angioplasty of the renal, femoral, popliteal and infra-popliteal arteries. This catheter is not for use in coronary arteries.

Contraindications: None known.

Warnings: 1. Contents supplied STERILE using ethylene oxide (EO). Non-Pyrogenic. Do not use if sterile barrier is opened or damaged. Do not reuse, reprocess or resterilize. 2. This device has been designed for single use only. Reusing this medical device bears the risk of cross-patient contamination as medical devices – particularly those with long and small lumina, joints, and/or crevices between components – are difficult or impossible to clean once body fluids or tissues with potential pyrogenic or microbial contamination have had contact with the medical device for an indeterminate period of time. The residue of biological material can promote the contamination of the device with pyrogens or microorganisms which may lead to infectious complications. 3. Do not resterilize. After resterilization, the sterility of the product is not guaranteed because of an indeterminate degree of potential pyrogenic or microbial contamination which may lead to infectious complications. Cleaning, reprocessing, and/or resterilization of the present medical device increases

the probability that the device will malfunction due to potential adverse effects on components that are influenced by thermal and/or mechanical changes. 4. To reduce the potential for vessel damage, the inflated diameter and length of the balloon should approximate the diameter and length of the vessel just proximal and distal to the stenosis. 5. When the catheter is exposed to the vascular system, it should be manipulated while under high-quality fluoroscopic observation. Do not advance or retract the catheter unless the balloon is fully deflated. If resistance is met during manipulation, determine the cause of the resistance before proceeding. Applying excessive force to the catheter can result in tip or catheter breakage, catheter kink, or balloon separation. 6. Do not exceed the RBP recommended for this device. Balloon rupture may occur if the RBP rating is exceeded. To prevent over pressurization, use of a pressure monitoring device is recommended. 7. After use, this product may be a potential biohazard. Handle and dispose of in accordance with acceptable medical practices and applicable local, state, and federal laws and regulations.

Please consult product labels and instruction for use for indications, contraindications, hazards, warnings, and precautions.



Going the distance

Ultraverse™ RX
PTA Dilatation Catheter



BD Switzerland Sarl, Terre Bonne Park – A4, Route De Crassier, 17, 1262 Eysins, Vaud, Switzerland.

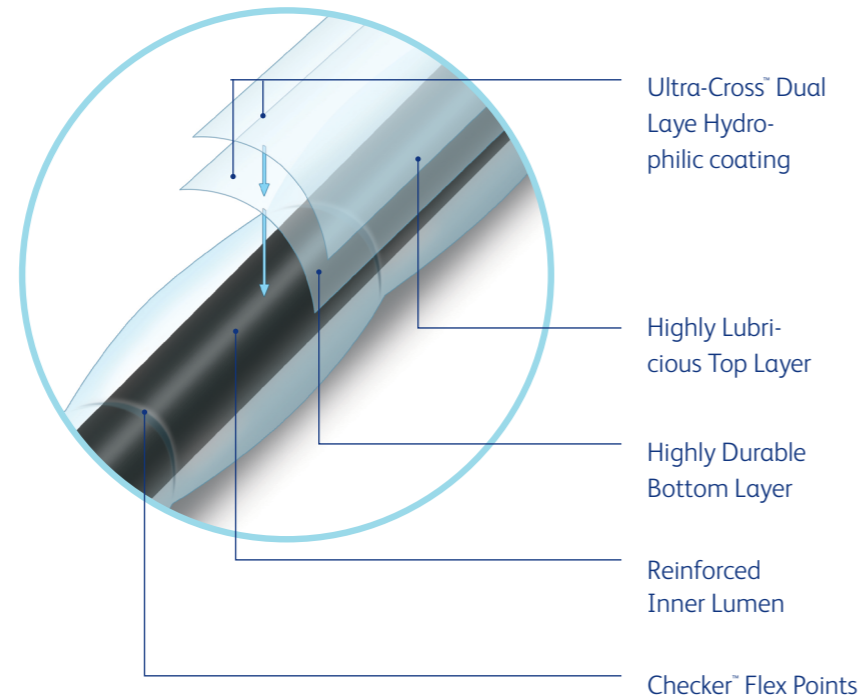
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Navigating the extremes

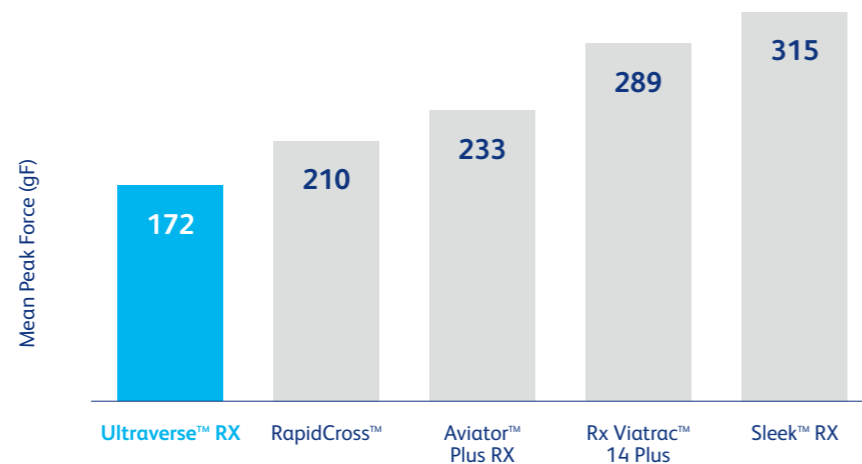
- Exceptional trackability
- Balloon lengths up to 300 mm
- Ultra-Cross™ Dual Layer Hydrophilic coating designed to reduce friction
- Checker™ Flex Points engineered to allow the balloon to flex in tortuous anatomy
- Reinforced inner lumen provides improved axial strength constructed to cross tight lesions



Excellent Trackability**

Requires up to 45% less force to track

(lower is better)



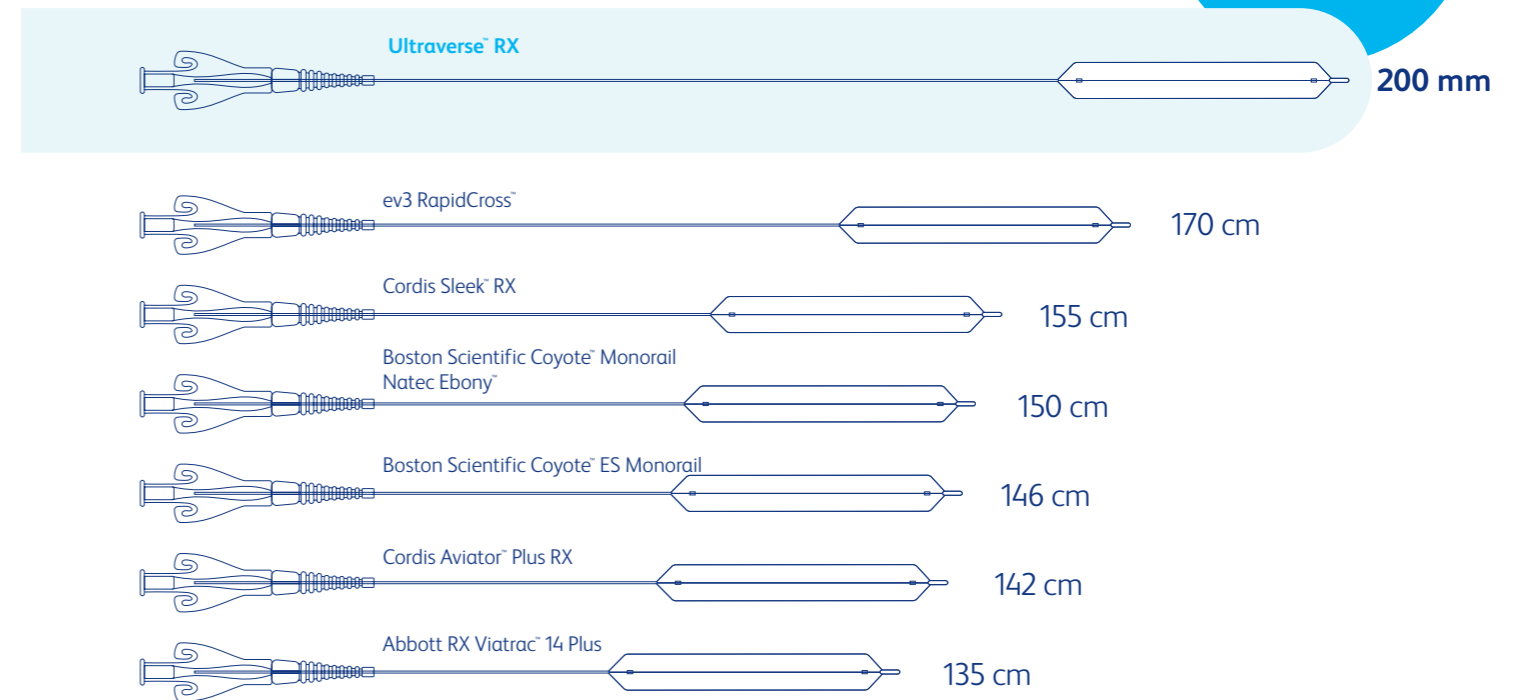
**4 mm x 40 mm x 150 cm Ultraverse™ RX - N=20; 4 mm x 40 mm x 170 cm RapidCross™ - N=20; 4 mm x 40 mm x 140 cm Aviator™ Plus RX - N=20; 4 mm x 40 mm x 135 cm Rx Viatrac™ 14 Plus - N=40; 4 mm x 40 mm x 150 cm Sleek™ RX - N=40. p<.05.
Bench data on file. Results may not necessarily be indicative of clinical performance. Different tests may yield different results.

Maneuver past the boundaries

- 200 cm shaft – longest on the market*
- Longest balloon available* – 300 mm

Enables a radial approach

Long Shaft*



Long Balloons*



Recommended Guidewire .014"		
Balloon Diameters	1.25 - 7 mm	
Balloon Lengths	15 - 300 mm	
Shaft Lengths	150, 200 cm	
Rated Burst Pressure up to	16 atm	
Nominal Pressure	6 atm	
Material	Nylon	