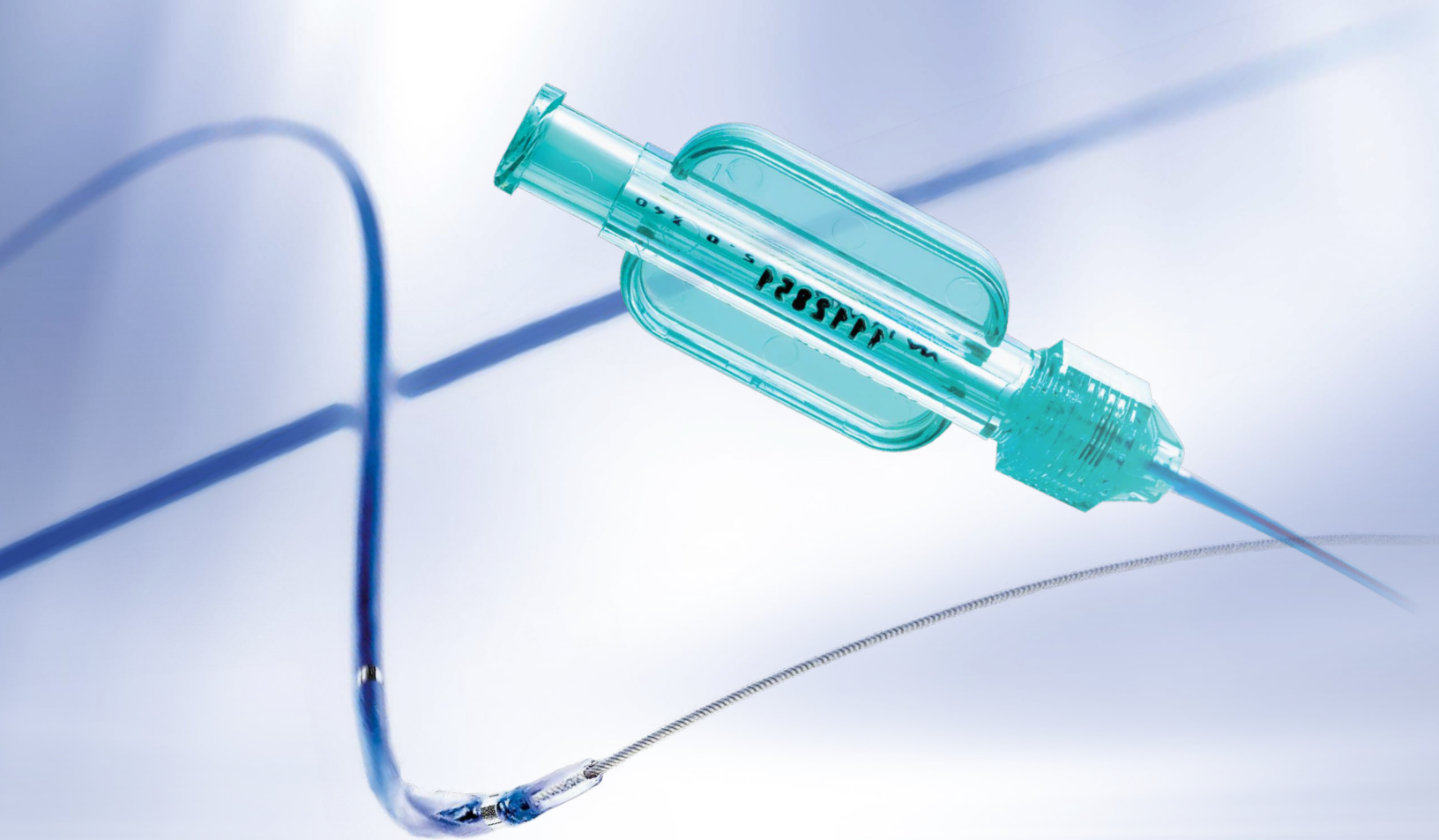


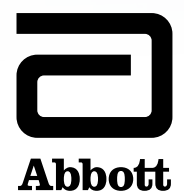
RX VIATRAC 14 PLUS

Peripheral Dilatation Catheter



CONTROL AND DURABILITY

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Check the regulatory status of the device in areas where CE marking is not the regulation in force.
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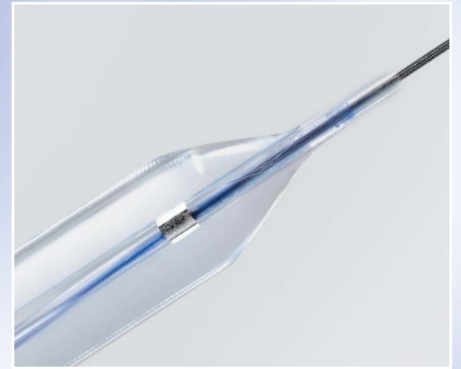


CONTROL AND DURABILITY

RAPID EXCHANGE SYSTEM DESIGNED FOR PERIPHERAL ARTERIES AND ARTERIOVENOUS DIALYSIS FISULAE

RAPID EXCHANGE

- Rapid exchange delivery system for single operator use and fast balloon exchanges
- Balloon lengths of 15, 20, 30 and 40 mm
- Catheter lengths of 80 and 135 cm
- 6F guide compatible across all balloon sizes and with shorter 130 cm guide wires



HIGH-PRESSURE STRENGTH

- Durable XCELON balloon material with high RBPs of 14 atm for sizing and flexibility
- Advanced balloon processing designed for outstanding rewrap and easy withdrawal

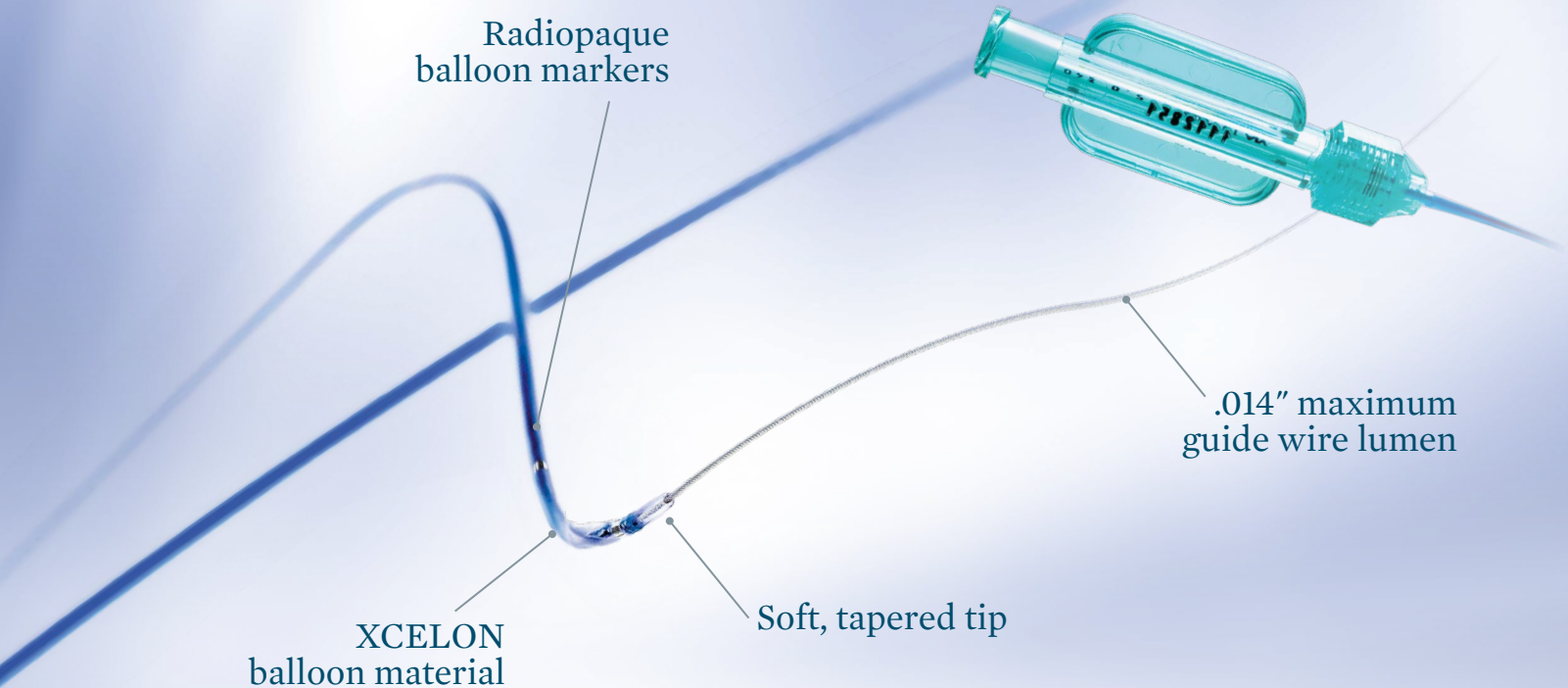


DELIVERABLE PERFORMANCE

- Flexible distal catheter with tapered support mandrel inside catheter to improve pushability and kink resistance
- Low profile design to cross tight lesions
- Hydrophobic coated shaft and balloon to reduce friction



DELIVERING PERFORMANCE THROUGH LOW PROFILE ACCESS



- .014" guide wire for easy lesion cross
- 8 atm Nominal, 14 atm RBP for all sizes
- 6F guide catheter on all sizes
- Easy contrast injections in a 6F guiding catheter
- 2-year shelf life

RX VIATRAC 14 PLUS

ORDERING INFORMATION

80 cm Catheter Length Art. No. (Prefix)	Balloon Length [mm] (Suffix)	Balloon Diameter [mm]	Crossing Profile [inch]	Nominal Pressure [bar]	Rated Burst Pressure [bar]	Prox./Dist. Shaft Ø [F]	Min. Guiding Catheter [inch]	Minimum Sheath Size [F]	Maximum Guide wire [inch]
1008190	-15, -20, -30, -40	4.0	0.036	8	14	3.3/3.3-2.9	6F (.067)	4	0.014
1008192	-15, -20, -30, -40	4.5	0.037	8	14	3.3/3.3-2.9	6F (.067)	4	0.014
1008194	-15, -20, -30, -40	5.0	0.038	8	14	3.3/3.3-2.9	6F (.067)	5	0.014
1008196	-15, -20, -30, -40	5.5	0.041	8	14	3.3/3.3	6F (.067)	5	0.014
1008198	-15, -20, -30, -40	6.0	0.043	8	14	3.6/3.3	6F (.067)	5	0.014
1008200	-15, -20, -30, -40	6.5	0.048	8	14	3.6/3.5	6F (.067)	5	0.014
1008202	-15, -20, -30, -40	7.0	0.049	8	14	3.6/3.5	6F (.067)	5	0.014

135 cm Catheter Length Art. No. (Prefix)	Balloon Length [mm] (Suffix)	Balloon Diameter [mm]	Crossing Profile [inch]	Nominal Pressure [bar]	Rated Burst Pressure [bar]	Prox./Dist. Shaft Ø [F]	Min. Guiding Catheter [inch]	Minimum Sheath Size [F]	Maximum Guide wire [inch]
1008189	-15, -20, -30, -40	4.0	0.036	8	14	3.3/3.3-2.9	6F (.067)	4	0.014
1008191	-15, -20, -30, -40	4.5	0.037	8	14	3.3/3.3-2.9	6F (.067)	4	0.014
1008193	-15, -20, -30, -40	5.0	0.038	8	14	3.3/3.3-2.9	6F (.067)	5	0.014
1008195	-15, -20, -30, -40	5.5	0.041	8	14	3.3/3.3	6F (.067)	5	0.014
1008197	-15, -20, -30, -40	6.0	0.043	8	14	3.6/3.3	6F (.067)	5	0.014
1008199	-15, -20, -30, -40	6.5	0.048	8	14	3.6/3.5	6F (.067)	5	0.014
1008201	-15, -20, -30, -40	7.0	0.049	8	14	3.6/3.5	6F (.067)	5	0.014

COMPLIANCE

Balloon Ø [mm]	Inflation Pressure [bar]																	
	2	3	4	5	6	7	8*	9	10	11	12	13	14**	15	16	17	18	
4.0	3.58	3.68	3.76	3.84	3.90	3.96	4.00*	4.04	4.08	4.11	4.14	4.16	4.19**	4.21	4.23	4.26	4.28	
4.5	3.88	4.00	4.11	4.20	4.36	4.43	4.50*	4.56	4.61	4.66	4.70	4.75	4.79**	4.83	4.87	4.92	4.97	
5.0	4.34	4.47	4.59	4.69	4.83	4.92	5.00*	5.07	5.14	5.21	5.27	5.34	5.40**	5.46	5.53	5.60	5.68	
5.5	4.94	5.06	5.17	5.26	5.35	5.43	5.50*	5.57	5.63	5.68	5.74	5.79	5.84**	5.90	5.95	6.01	6.08	
6.0	5.35	5.49	5.62	5.73	5.83	5.92	6.00*	6.07	6.13	6.19	6.24	6.29	6.34**	6.39	6.45	6.50	6.56	
6.5	5.76	5.93	6.08	6.21	6.32	6.42	6.50*	6.57	6.63	6.68	6.73	6.77	6.81**	6.85	6.88	6.92	6.97	
7.0	6.23	6.39	6.54	6.67	6.79	6.90	7.00*	7.09	7.18	7.26	7.34	7.41	7.49**	7.58	7.66	7.76	7.86	

*Nominal **Rated Burst Pressure [RBP]

Caution: This product is intended for use by or under the direction of a physician. Prior to use, reference the Instructions for Use provided inside the product carton (when available), at eifu.abbottvascular.com or at Manuals.sjm.com for more detailed information on Indications, Contraindications, Warnings, Precautions and Adverse Events. **Information contained herein for distribution outside the U.S. only.** Check the regulatory status of the device before distribution in areas where CE marking is not the regulation in force.

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