

SAPPHIRE[®] II PRO 1.0 mm - 1.5 mm

Coronary Dilatation Catheter

Cross with Confidence Every Time

Redefining the standard for crossing chronic total occlusions

Single-component **Sub-Zero tapered** tip, tailored for the smaller catheter size

Durable balloon material and **higher RBP** for resistant CTO lesions

1.0 mm
for Complex
CTO lesions

Increased inner body wall thickness to **reduce guidewire friction**

Single marker band and improved balloon folding for a **low crossing profile**

Spiral-cut hypotube transition zone to reduce kinking and maintain **excellent force transmission**

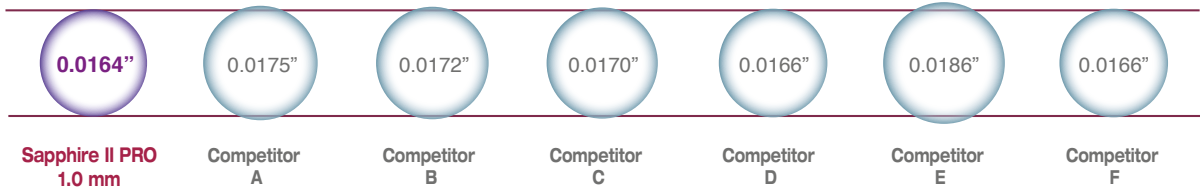
Technical Specifications

Catheter Type	Rapid Exchange	Compliance	Semi-compliant
Proximal Shaft Diameter	1.9F	Crossing Profile	0.0216"
Distal Shaft Diameter	2.36F	Marker Bands	1
Catheter Working Length	140 cm	Coating	Hydrophilic (distal tip to guidewire exit marker); Hydrophobic (guidewire lumen)
Leading Tip Length	1.5 mm	Nominal Pressure	6 atm
Balloon Material	Nylon	Rated Burst Pressure	16 atm
Balloon Folds	2 (Ø 1.0 mm); 3 (Ø 1.25 mm - 1.5 mm);	Guiding Catheter Compatibility	5F

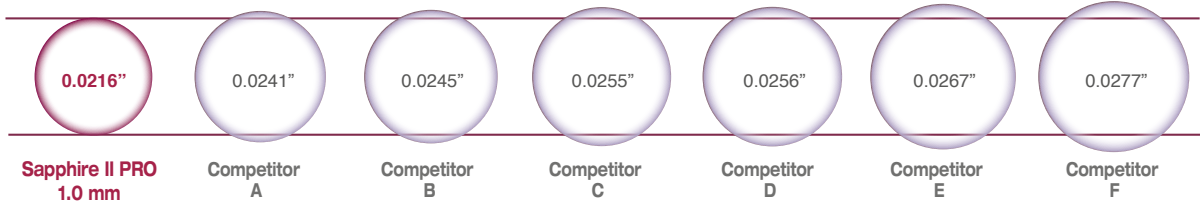
SAPPHIRE[®] II PRO 1.0 mm - 1.5 mm

Coronary Dilatation Catheter

Tip Entry Comparison*



Crossing Profile Comparison*



*Data on file, scale depictions

Ordering Information

Non-EU Manufacturing Catalogue Numbers

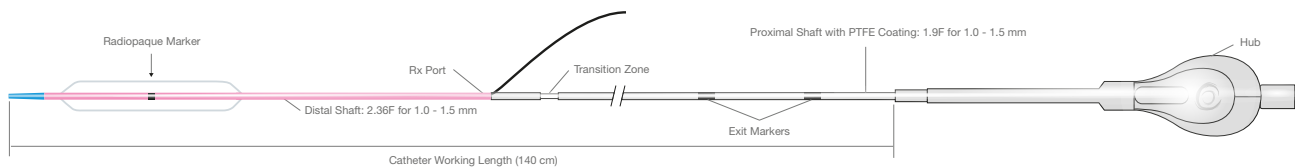
Balloon Diameter (mm)	Balloon Working Length (mm)					
	5	8	10	12	15	20
1.0	210-053-5UJ	210-083-5UJ	210-103-5UJ	—	210-153-5UJ	—
1.25	212-053-5UJ	212-083-5UJ	212-103-5UJ	—	212-153-5UJ	—
1.5	—	—	215-103-5UJ	215-123-5UJ	215-153-5UJ	215-203-5UJ

Compliance Chart

Pressure (atm)	Balloon Diameter (mm)		
	1.0	1.25	1.5
2	0.94	1.18	1.42
4	0.97	1.22	1.46
6 NOM*	1.00	1.25	1.50
8	1.03	1.28	1.54
10	1.06	1.32	1.58
12	1.09	1.35	1.62
14	1.12	1.38	1.66
16 RBP**	1.15	1.42	1.70
18	1.18	1.45	1.74
20	1.21	1.48	1.78
22	1.24	1.52	1.82

* Nominal Pressure. The nominal in-vitro device specifications do not take into account any lesion resistance.

** Rated Burst Pressure. Do not exceed RBP.



For more information please visit our website at www.OrbusNeich.com or contact us:

Corporate Headquarters

Unit 303 & 305, 3/F, Building 20E
 Hong Kong Science Park
 Shatin, N.T., Hong Kong, China
 Phone +852.2802.2288
 Fax +852.2507.3532

EMEA Regional Headquarters

Drs. W. van Royenstraat 5
 3871 AN Hoevelaken
 The Netherlands
 Phone +31.33.254.1150 Toll Free Phone 00800.0254.1150*

* Only for Belgium, Denmark, France, Germany, Ireland, Netherlands, Norway, Sweden and UK

©2024 OrbusNeich Medical Group Holdings Limited or its affiliates. All rights reserved.

The Sapphire II PRO product is now approved in the USA and distributed with USA specific catalog numbers.

G-70-0410 Rev06

Sapphire and OrbusNeich are registered trademarks of OrbusNeich Medical Group Holdings Limited or its affiliates. All data and photos on file. Illustrations are not to be considered as engineering drawings or photos.

