

# titan<sup>TM</sup>

aspiration catheter system



balt  
inspiring innovation

# titan™

## Aspiration Catheter System

Indicated for injection of intravascular fluids, the introduction of interventional devices into the peripheral & neuro vasculature, and/or aspiration of soft emboli and thrombi from the arterial system, including the peripheral and neuro vasculature.

### unique design

to optimize the balance between lumen diameter, proximal support & distal trackability

**14 transition zones** to maximize the distal zone flexibility

**Hydrophilic coating** on 90 cm long

### available in kit

Together with .070 & .036 designed to

- ease navigation in the ophthalmic region
- better track distally

### compatible with



ballast  
80, 90  
& 100



catchview  
& catch+  
stentriever



## ordering information

Reference	Product code	Working length	Proximal outer Ø	Distal outer Ø	Distal inner Ø	Hydrophilic coating length
Titan™ 036 Catheter System	TC036-1	160cm	.056"	.049"	.036"	90cm
Titan™ 070 Catheter System	TC070-2	128cm	.083"	.0805"	.070"	90cm
Titan™ Catheter System Kit	TC070-036-2	160 / 128cm	.056 / .083"	.049 / .0805"	.036 / .070"	90cm

Introductory tool included in the package

The TITAN™ 036 Catheter is indicated for general intravascular use, including the neuro and peripheral vasculature. It can be used to facilitate introduction of diagnostic and therapeutic agents. It is not intended for use in coronary arteries. The TITAN 070 Catheter is indicated for injection of intravascular fluids, the introduction of interventional devices into the peripheral and neuro vasculature, and removal/aspiration of soft emboli and thrombi from the arterial system, including the peripheral and neurovasculature. The TITAN™ Catheter System Kit is indicated for injection of intravascular fluids, the introduction of interventional devices into the peripheral and neuro vasculature, and removal/aspiration of soft emboli and thrombi from the arterial system, including the peripheral and neuro vasculature. Class III CE0297 in compliance with Medical Device Directive (MDD 93/42/EEC amended by 2007/47/EC). Manufactured by BALT USA LLC. Carefully read the instructions for use before use. Not reimbursed. First CE marking:2019.

The content of this document, in particular data, information, trademarks and logos is BALT SAS and affiliate's sole property. ©2019 BALT SAS and affiliates, all rights reserved. All representation and/or reproduction, whether in part or in full, is forbidden and would be considered a violation of BALT SAS and affiliates' copyrights and other intellectual proprietary rights. This document with associated pictures is non-contractual and is solely dedicated to healthcare professionals and BALT's distributors (BALT's supplier's distributors). It cannot be distributed or given to patients. The products commercialized by BALT shall exclusively be used in accordance with the instructions for use included in the boxes. DCO47GB (05/19)

### Balt

10, rue de la Croix Vigneron, 95160 Montmorency, France

Tél. : +33 (0)1 39 89 46 41

Fax : +33 (0)1 34 17 03 46

www.balt.fr

