

**React** with Speed.  
**React** with Agility.  
**React** with Confidence.

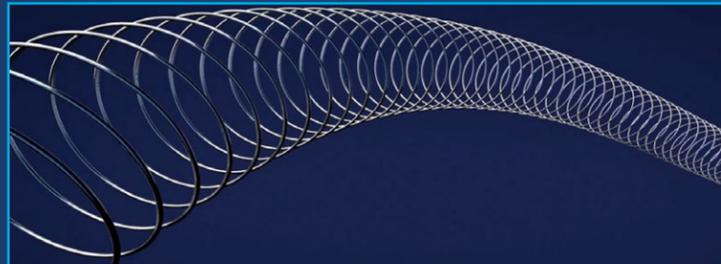


# React™ 68 Catheter

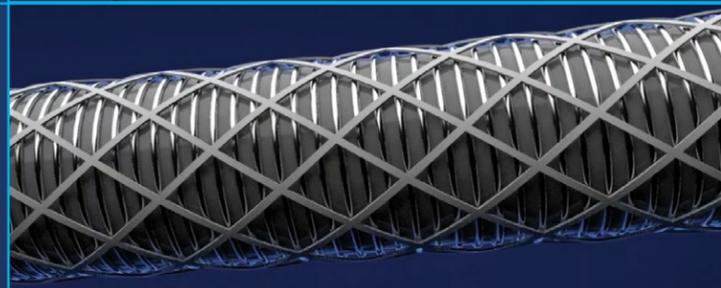
**Medtronic**  
Further.Together

STROKE IS NEVER EXPECTED,  
BUT KNOWING HOW TO **REACT**,  
HAS THE POWER TO CHANGE LIVES.

**NAVIGABILITY**  
OF A COIL<sup>1</sup>



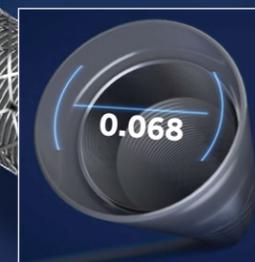
**PUSHABILITY**  
OF A BRAID<sup>1</sup>



**DURABILITY**  
OF NITINOL<sup>1</sup>



COMPATIBLE WITH  
**SOLITAIRE™ PLATINUM**<sup>1</sup>



- Large 0.068 Inner Diameter Lumen<sup>1</sup>
- Atraumatic Beveled Distal Tip<sup>1</sup>

FEATURING

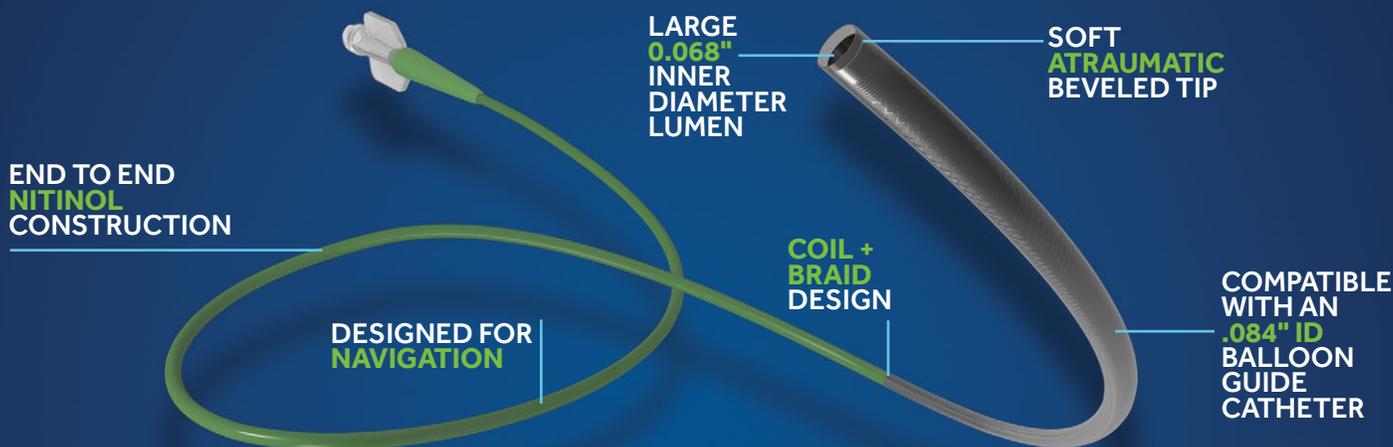
**COBRA**  
COIL + BRAID

TECHNOLOGY<sup>1</sup>

An overlapping coil +  
braid construction<sup>1</sup>



# PRODUCT FEATURES<sup>1</sup>



## REACT™ 68 CATHETER ORDERING INFORMATION<sup>2</sup>

Product Catalog Number	ID	MAX OD	Working Length		
<b>REACT-68</b>	<b>0.068"</b>	<b>0.083"</b>	<b>132cm</b>		

## BUNDLE ORDERING INFORMATION

Product Catalog Number	Riptide™ Aspiration Pump	React™ 68 Catheter	Riptide™ Aspiration Tubing	Riptide™ Collection Canister	Solitaire™ Platinum
<b>RBUNDLE-68</b>	<b>Qty. 1</b>	<b>Qty. 3</b>	<b>Qty. 3</b>	<b>Qty. 3</b>	
<b>RIPTIDE-68</b>		<b>Qty. 1</b>	<b>Qty. 1</b>	<b>Qty. 1</b>	
<b>STAT-6-40-68</b>		<b>Qty. 1</b>	<b>Qty. 1</b>	<b>Qty. 1</b>	<b>Qty. 1 6x40</b>
<b>STAT-4-40-68</b>		<b>Qty. 1</b>	<b>Qty. 1</b>	<b>Qty. 1</b>	<b>Qty. 1 4x40</b>
<b>REACT-6-40</b>		<b>Qty. 1</b>			<b>Qty. 1 6x40</b>
<b>REACT-4-40</b>		<b>Qty. 1</b>			<b>Qty. 1 4x40</b>

CAUTION: Federal (USA) law restricts this device to sale distribution and use by or on order of a physician. Indications, contraindications, warnings and instructions for use can be found in the product labeling supplied with each device.

The Riptide™ Aspiration System is intended for use in the revascularization of patients with acute ischemic stroke secondary to intracranial large vessel occlusive disease (within the internal carotid, middle cerebral M1 and M2 segments, basilar, and vertebral arteries) within 8 hours of symptom onset. Patients who are ineligible for intravenous tissue plasminogen activator (IV t-PA) or who fail IV t-PA therapy are candidates for treatment.

The React™ 68 Catheter is indicated for the introduction of interventional devices into the peripheral and neuro vasculature.

The Solitaire™ Revascularization Device is indicated for use to restore blood flow in the neuro-vasculature by removing thrombus for the treatment of acute ischemic stroke to reduce disability in patients with a persistent, proximal anterior circulation, large vessel occlusion, and smaller core infarcts who have first received intravenous tissue plasminogen activator (IV t-PA). Endovascular therapy with the device should be started within 6 hours of symptom onset.

The Solitaire™ Revascularization Device is indicated to restore blood flow by removing thrombus from a large intracranial vessel in patients experiencing ischemic stroke within 8 hours of symptom onset. Patients who are ineligible for IV t-PA or who fail IV t-PA therapy are candidates for treatment.

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1. All claims have been substantiated and data is on file TR-NV15399 Rev A.

2. DWGSREACT-68 Rev C Design Specification

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