OUTBACK® Elite
Re-Entry Catheter

True Precision.
True Control.
True Lumen.
The OUTBACK® Elite Re-Entry Catheter enables faster and more precise re-entry into the true lumen in the most challenging cases.¹

- NEW! 80 CM shaft
- NEW! Ergonomic handle for greater control
- Increased precision of target site re-entry¹
- Robust Nitinol Cannula

In a recent study by Gandini et al., the OUTBACK® LTD Re-Entry Catheter was shown to have a higher success rate of precision re-entry. Use of the device also saved the operator and the patient from additional fluoroscopy and procedure time.

OUTBACK® Elite Re-Entry Catheter

Clinical Performance

<table>
<thead>
<tr>
<th>Success Rate (entry w/in 5cm of target)¹</th>
<th>Percent</th>
</tr>
</thead>
<tbody>
<tr>
<td>OUTBACK™</td>
<td>100</td>
</tr>
<tr>
<td>Wire</td>
<td>42</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Mean Procedural and Fluoroscopy Times</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean Procedure Time (min)</td>
</tr>
<tr>
<td>N=26 OUTBACK™</td>
</tr>
<tr>
<td>Mean Fluoroscopy (min)</td>
</tr>
<tr>
<td>N=26 Wire</td>
</tr>
</tbody>
</table>


Image Courtesy of Ali Amin, MD, FACS, FACC, RVT
Quick Reference Procedural Guide

This is intended as a quick reference guide only. Prior to use, please read the full Instructions For Use for complete information.

Step 1 - Preparation

- REMOVE cannula tip cover
- FLUSH device at the flush and guidewire exit port, then repeat
- ACTUATE cannula 2-3 times; turn rotating knob to confirm functionality
- RETRACT cannula, and ENSURE the cannula is fully retracted and the deployment slide is locked in the most proximal position
- If cannula does not fully retract back into shaft after initial preparation, REPEAT flush sequence
- SELECT recommended 0.014” guidewire

Step 2 - Position Catheter

- ADVANCE the catheter over the wire to the desired site.
- RETRACT the 0.014” guidewire into the catheter approximately 5 cm.
- ROTATE the image intensifier so the catheter appears adjacent to the target vessel under fluoroscopy.
- ORIENT the “L” marker to point toward the target vessel by rotating the Rotator Knob.
- ROTATE the image intensifier so the catheter appears superimposed over the target vessel under fluoroscopy (90° orthogonal view).
- ADJUST the catheter by rotating the Rotator Knob such that the radiopaque marker appears as a “T”.
- RELEASE any stored torque in the catheter.

Step 3 - Re-Entry

- ROTATE the image intensifier back to the previous position to confirm “L” marker.
- DEPLOY cannula into the target vessel by advancing the Deployment Slide.
- ADVANCE the 0.014” guidewire into the target vessel.
- RETRACT the cannula and remove catheter over the guidewire.

Wire Compatibility List

Cordis Corporation
- ATW™ All Track Wire
- STABILIZER® Plus Guidewire
- STABILIZER® XS Guidewire

Abbott Vascular
- Hi-Torque All Star™ Guidewire
- Hi-Torque Balance Middleweight (BMW) ELITE™ Guidewire
- Hi-Torque Command™ Guidewire
- Hi-Torque Command™ ES Guidewire
- Hi-Torque Ironman™ Guidewire
- Hi-Torque SpartaCore™ Guidewire
- Hi-Torque Whisper™ Guidewire

Asahi Intec
- Confianza® Guidewire

Boston Scientific
- ChoICE™ Extra Support Guidewire
- Journey™ Guidewire
- Luge™ Guidewire
- Mailman™ Guidewire
- Platinum Plus™ Guidewire
- PT2™ Moderate Support Guidewire

Ordering Information

<table>
<thead>
<tr>
<th>Part Number</th>
<th>Description</th>
<th>Length</th>
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<tbody>
<tr>
<td>OTB59080A</td>
<td>OUTBACK® Elite Re-Entry Catheter</td>
<td>80 cm</td>
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<tr>
<td>OTB59120A</td>
<td>OUTBACK® Elite Re-Entry Catheter</td>
<td>120 cm</td>
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Caution: Federal (USA) law restricts these devices to sale by or on the order of a physician. This quick reference guide includes demonstration of the use of a medical device but is not intended to be used as a training guide. The steps demonstrated may not be the complete steps of the procedure. Individual physician preference and experience, as well as patient needs, may dictate variation in procedure steps. Before using any medical device, including those demonstrated or referenced in this quick reference guide, review all relevant package inserts and labeling, with particular attention to the indications, contraindications, warnings and precautions, and steps for use of the device.

Important Information: Prior to use, refer to the Instructions for Use supplied with this device for indications, contraindications, side effects, suggested procedure, warnings, and precautions.

As part of the Cordis policy of continuous product development, we reserve the right to change product specifications without prior notification.

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Contact your Cordis sales representative for product availability and ordering information. EU751 6/15 - 2E-800-2118-1