Capio™ SLIM
Suture Capturing Device

Building on the history, reducing the profile
The Capio SLIM Suture Capturing device features a reduced profile to minimize the space required within the surgical field, easy suture loading and a funnel shaped catch, designed for consistent ease of use.

Since 1995, the Capio Device has provided consistent suture placement in difficult to access pelvic floor locations.

The Capio SLIM Suture Capturing Device

New “funnel” suture dart catch design

Ergonomic Handle Providing 3 Finger Grip

36% Reduction in Head Width

70% Reduction in Shaft Diameter

Easy Suture Dart Loading

The evolution continues...

1995
The Capio Suture Capturing Device debuts, revolutionizing the field of urogynecology.

1999
The Capio Device is used with Repliform™ Graft, offering another treatment option.

2006
The Capio Device is used with Xenform™ Graft, offering more material options for prolapse repair.
## Comparison of Device Heads

<table>
<thead>
<tr>
<th></th>
<th>Capio™ SLIM Suture Capturing Device</th>
<th>Digitex™ Suture Delivery System</th>
<th>FiXT™ Suturing Device</th>
</tr>
</thead>
</table>

Building on the history and reducing the profile, the Capio SLIM Suture Capturing Device features one of the smallest device profiles and suture dart carrier diameter while keeping the same bite depth as the current Capio Open Access Suturing Device.

### Fixation Device Comparison

<table>
<thead>
<tr>
<th></th>
<th>Shaft Diameter (mm)</th>
<th>Head Width (mm)</th>
<th>Device Weight (g)</th>
<th>Carrier Diameter (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Capio OPEN ACCESS Suture Capturing Device**</td>
<td>10.1</td>
<td>9.8</td>
<td>45.4</td>
<td>1.2</td>
</tr>
<tr>
<td>*<em>Capio SLIM Suture Capturing Device</em></td>
<td>3.0</td>
<td>6.3</td>
<td>38.2</td>
<td>1.2</td>
</tr>
<tr>
<td>Digitex Suture Delivery System*</td>
<td>9.5</td>
<td>9.6</td>
<td>133.3</td>
<td>2.1</td>
</tr>
<tr>
<td>FiXT Suturing Device*</td>
<td>4.8</td>
<td>6.9</td>
<td>121.1</td>
<td>2.2</td>
</tr>
</tbody>
</table>

*Measurements were recorded using one (1) of each device. Data on file at Boston Scientific.

**Head width and device weight values are an average of five (5) device measurement recordings. Shaft diameter values are an average of fifteen (15) device measurement recordings. Data on file at Boston Scientific.

---

**Figure 1:** Comparison of Device Heads illustrates the difference in carrier diameter amongst these fixation devices. The Capio SLIM Suture Capturing Device carrier diameter is 45% less than the FIXT Suturing Device and 43% less than the Digitex Suture Delivery System.

**Figure 2:** Fixation Device Comparison illustrates the device profile differences amongst these fixation devices. The Capio SLIM Suture Capturing Device has the smallest device profile based on the following measurements: shaft diameter, head width and device weight.

---

**2008**
The Capio Device is used with the Uphold™ System, one of the smallest mesh footprints for pelvic floor reconstruction.

**2011**
The Capio Device remains “the foundation” of Pelvic Floor Reconstruction.

**2012**
Capio SLIM: the evolution continues.
Capio™ SLIM Suture Capturing Device

<table>
<thead>
<tr>
<th>Order Number</th>
<th>Description</th>
<th>Unit</th>
</tr>
</thead>
<tbody>
<tr>
<td>M0068318250</td>
<td>Capio SLIM Suture Capturing Device</td>
<td>bx 1</td>
</tr>
<tr>
<td>M0068318261</td>
<td>Capio SLIM Suture Capturing Device</td>
<td>bx 5</td>
</tr>
</tbody>
</table>

Capio Sutures

<table>
<thead>
<tr>
<th>Order Number</th>
<th>Description</th>
<th>Size</th>
<th>Unit</th>
</tr>
</thead>
<tbody>
<tr>
<td>M0068331131</td>
<td>Non-absorbable, coated braided polyester, double armed with TC tapercut needle (dart), 48&quot;</td>
<td>0</td>
<td>bx 12</td>
</tr>
<tr>
<td>M0068331231</td>
<td>Non-absorbable, polypropylene monofilament, double armed with TC tapercut needle (dart), 48&quot;</td>
<td>0</td>
<td>bx 12</td>
</tr>
<tr>
<td>M0068332131</td>
<td>Absorbable, coated braided PGA, double armed with TC tapercut needle (dart), 48&quot;</td>
<td>0</td>
<td>bx 12</td>
</tr>
<tr>
<td>M0068331141</td>
<td>Non-absorbable, coated braided polyester, double armed with TC tapercut needle (dart) and a T 26mm 1/2 circle taper needle, 36&quot;</td>
<td>0</td>
<td>bx 12</td>
</tr>
<tr>
<td>M0068331241</td>
<td>Non-absorbable, polypropylene monofilament, double armed with TC tapercut needle (dart) and a T 26mm 1/2 circle taper needle, 36&quot;</td>
<td>0</td>
<td>bx 12</td>
</tr>
<tr>
<td>M0068331371</td>
<td>Monodek™ Absorbable, monofilament PDO, double armed with TC tapercut needle (dart) and a T 26mm 1/2 circle taper needle, 48&quot;</td>
<td>0</td>
<td>bx 12</td>
</tr>
</tbody>
</table>

Your Pelvic Floor Reconstructive Tool to Meet a Variety of Your Algorithm Needs:
- Native Tissue Repair
- Biologic Graft Augmentation
- Synthetic Mesh And More...

All images are owned by Boston Scientific. All trademarks are the property of their respective owners.

For Xenform Soft Tissue Repair Matrix: CAUTION: Federal (US) law restricts this device to sale by or on the order of a physician trained in the use of surgical mesh for repair of pelvic organ prolapse.

Repliform Tissue Regeneration Matrix complies with U.S. Regulations 21 CFR part 1271 Human Tissue Intended for Transplantation. Repliform Matrix is not CE marked and therefore is not available for distribution in European markets.

Products shown for informational purposes only - not meant as a promotion or offer for sale - certain components pending CE Mark, not available for sale in the European Economic Area (EEA)

Data on file. Bench test results may not necessarily be indicative of clinical performance. Results from case studies are not predictive of results in other cases. Results in other cases may vary.

CAUTION: Federal Law (USA) restricts these devices to sale by or on the order of a physician.

Refer to package insert provided with these products for complete Indications for Use, Contraindications, Warnings, Precautions, Adverse Events, and Instructions prior to using these products.

CAUTION: The law restricts these devices for sale by or on the order of a physician. Information for use only in countries with applicable health authority registrations. Material not intended for use in France.

Products shown for INFORMATION purposes only and may not be approved for sale in certain countries.

Please check availability with your local sales representative or customer service.