Boston Scientific

Advancing science for life™

Ultra

Retropubic Mid-Urethral Sling Family

Consistent.

Innovative.

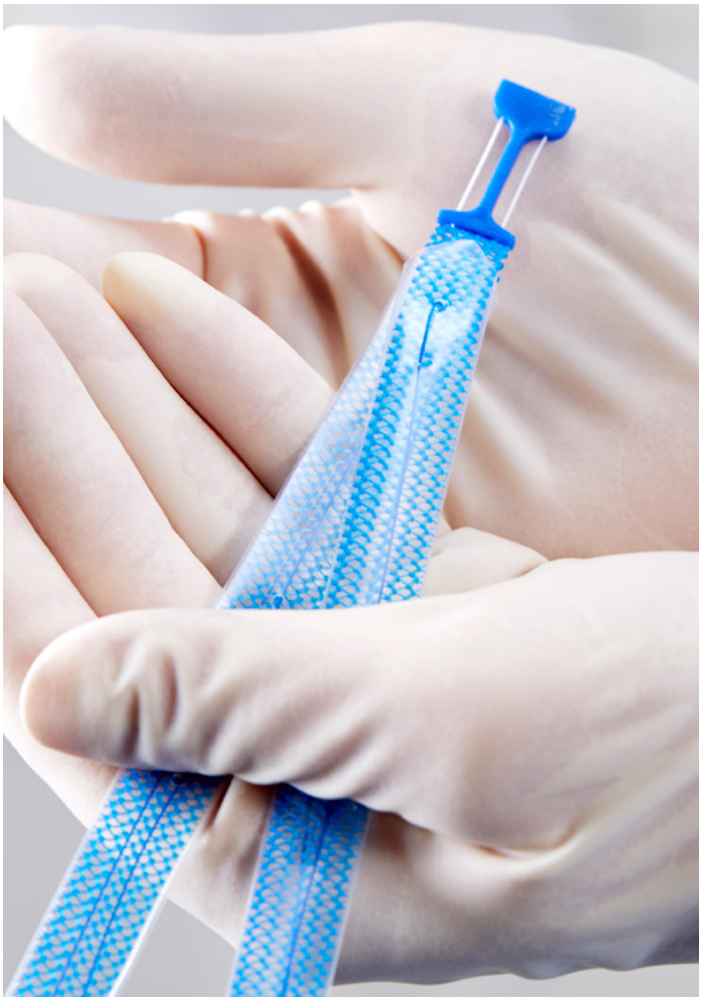
Precise.

The newest evolution in sling systems

Fueled by physician insights and feedback, the Ultra Sling Family is designed to improve provider experience through more precise sling placement.

Paired with Boston Scientific's clinically supported Advantage™ optical blue mesh, its features are designed to help drive:

- Procedural efficiency
- Mesh visualization
- Tensioning consistency

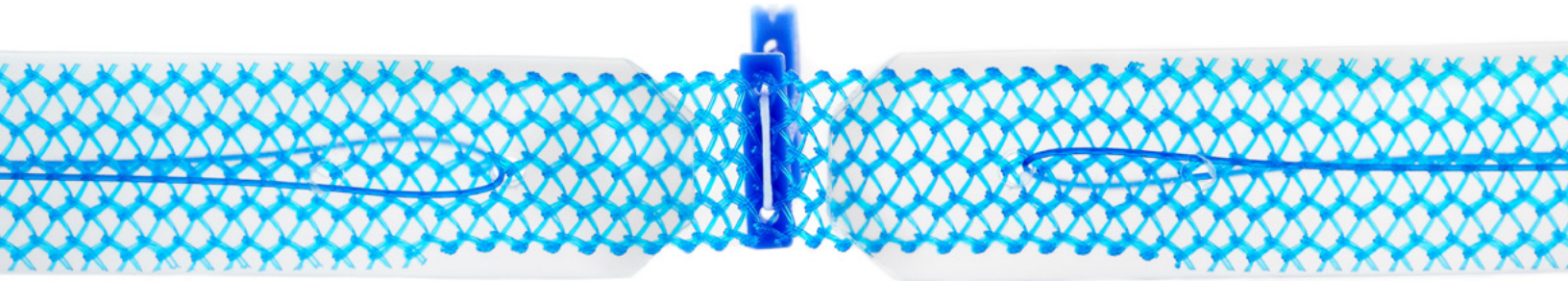


The Ultra Sling Family delivers physician-driven innovation across Boston Scientific's full-length mid-urethral sling portfolio.

Advantage™ Ultra Sling System

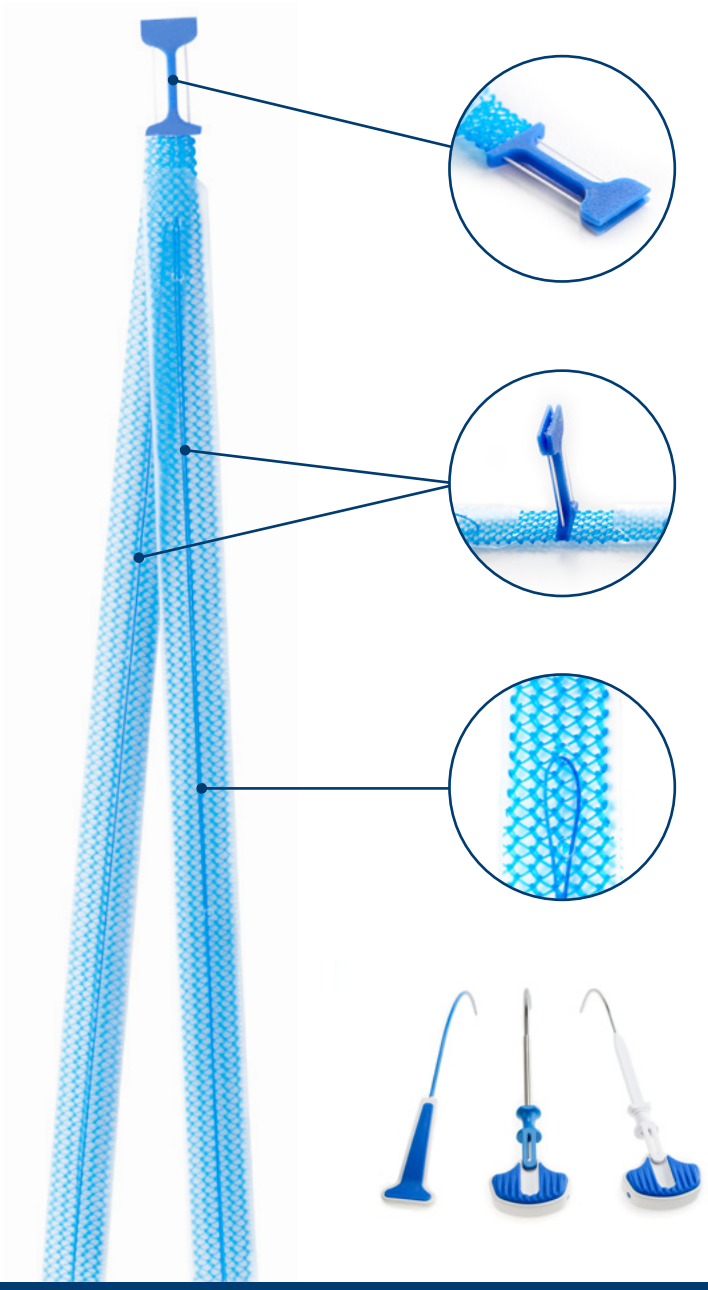
Advantage Fit™ Ultra Sling System

Lynx™ Ultra Sling System



You asked. We answered.

Boston Scientific's latest, physician-driven innovation is designed to drive efficiency.



Sleek, suture-released centering tab

- Designed with 1 cm exposed mesh for better visualization of the surgical field
- Updated removal process
- Ability to visualize tensioning and centering after sleeve removal

Lay-flat, two-sleeve design

- Designed to provide a slim, smooth surface during tissue interaction

Clinically supported Advantage blue mesh

- Offers the same material and features as the patented Advantage™ clear mesh in an optical blue color
- Enhances intra-operative visibility of mesh against the urethra

Enhancements are consistent across the Ultra Sling Family.

Ultra innovation includes Boston Scientific's clinically supported Advantage mesh, which is documented in more than **100 publications** to date and has been used in more than **1 million slings**.[1]

Dedicated to women

Since 1995, Boston Scientific has been committed to significantly and consistently investing in physician-driven, patient-centered pelvic floor research and innovation to deliver impactful solutions for women.

Ultra Retropubic Mid-Urethral Sling Family featuring Advantage™ Blue Mesh

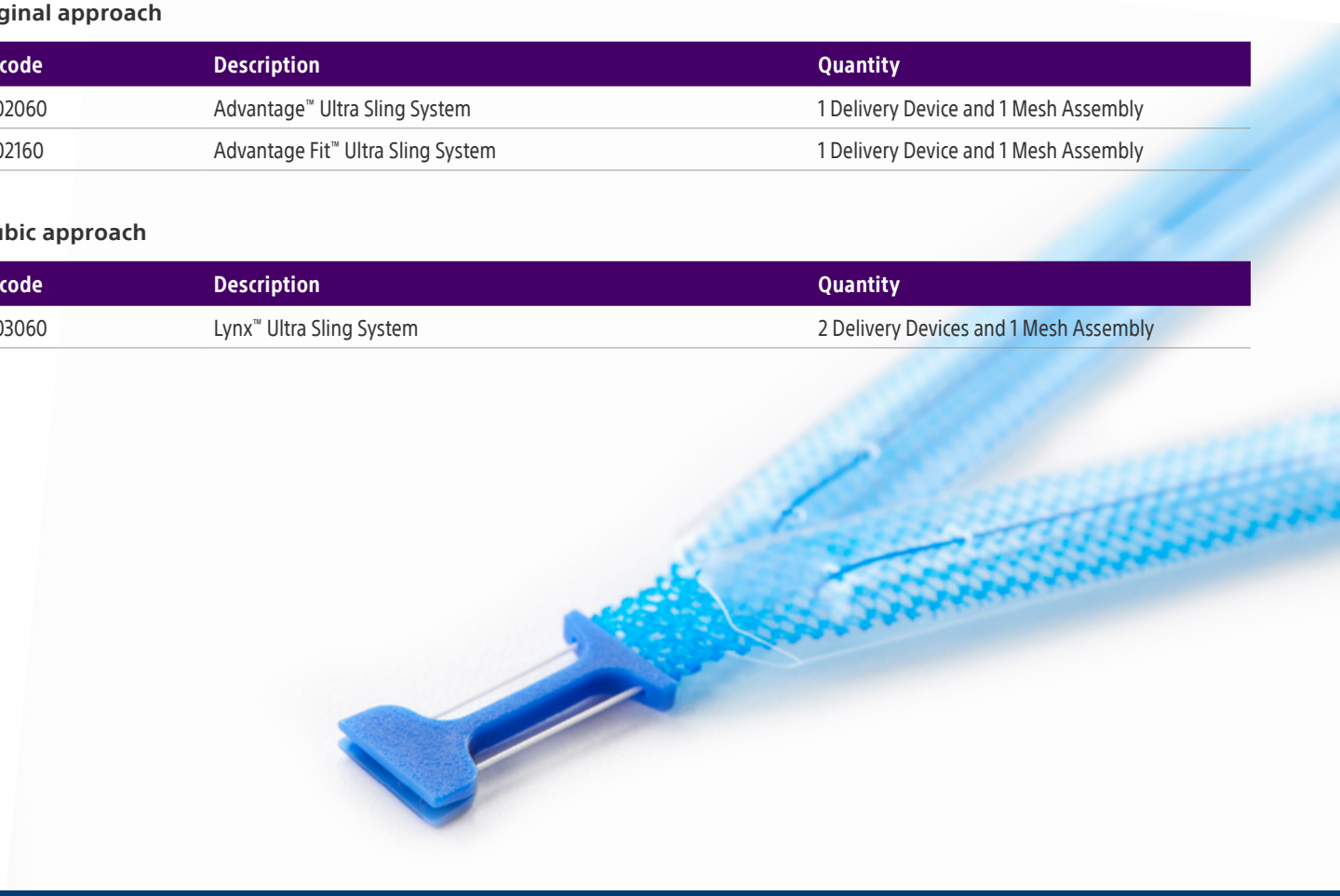
Ordering information

Transvaginal approach

Product code	Description	Quantity
M0068502060	Advantage™ Ultra Sling System	1 Delivery Device and 1 Mesh Assembly
M0068502160	Advantage Fit™ Ultra Sling System	1 Delivery Device and 1 Mesh Assembly

Suprapubic approach

Product code	Description	Quantity
M0068503060	Lynx™ Ultra Sling System	2 Delivery Devices and 1 Mesh Assembly



To learn more, contact your Boston Scientific representative or visit **bostonscientific.com/Ultra**

Reference:

1. Data on file with Boston Scientific

CAUTION: The following adverse events have been reported due to suburethral sling placement, any of which may be ongoing, but are not limited to: As with all implants, local irritation at the wound site and/or a foreign body response may occur, Foreign body reaction may be acute or chronic, Pain (pelvic, vaginal, groin/thigh, suprapubic, dyspareunia) (acute or chronic), Dyspareunia, Tissue responses to the mesh implant could include: erosion into organs (urethra, bladder or other surrounding tissues); exposure/extrusion into the vagina, Mesh contact with urine via erosion/exposure/extrusion may result in stone formation, scarring/scar contracture, Necrosis, fistula formation (acute or chronic), inflammation (acute or chronic), Mesh contracture, Tissue contracture, Vaginal shortening or stenosis that may result in dyspareunia and/or sexual dysfunction, Pain with intercourse that may not resolve, Exposed mesh may cause pain or discomfort to the patient's partner during intercourse, Sexual dysfunction, including the inability to have intercourse. Like all foreign bodies, the mesh may potentiate an existing infection. Allergic reaction has been reported. Known risks of surgical procedures for the treatment of incontinence include: pain, ongoing pain (pelvic, vaginal, groin/thigh, suprapubic, dyspareunia), Severe, chronic pain, Apareunia, Leg weakness, Infection, De novo detrusor instability, Complete failure of the procedure/failure to resolve a patient's stress urinary incontinence, Voiding dysfunction (incontinence, temporary or permanent lower urinary tract obstruction, difficulty urinating, pain with urination, overactive bladder, and retention), Bruising, bleeding (vaginal, hematoma formation), Abscess, Vaginal discharge, Dehiscence of vaginal incision, Edema and erythema at the wound site, Perforation or laceration of vessels, nerves, bladder, urethra or bowel may occur during placement. The following additional adverse events have been reported for the Solyx SIS System: Dysuria, Hematuria. The occurrence of these events may require surgical intervention and possible removal of the entire mesh. In some instances, these events may persist as a permanent condition after surgical intervention or other treatment. Removal of mesh or correction of mesh-related complications may involve multiple surgeries. Complete removal of mesh may not be possible and additional surgeries may not always fully correct the complications.

CAUTION: For Female Mid-Urethral Slings: Federal (US) law restricts this device to sale by or on the order of a physician trained in use of surgical mesh for repair of stress urinary incontinence.

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

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