## AdVance™ Male Sling System

## Patient selection criteria<sup>1</sup>

## Six (6) patient selection criteria for AdVance Sling (post-prostatectomy incontinence):

- 1 <4 pads/day or <300g daily pad weight
- Intact appearing external urinary sphincter without segmental defects on cystoscopy; ability to volitionally contract their external urinary sphincter
- **3** No history of pelvic radiation or cryotherapy
- ▶ 4 No history of previous surgical procedures to treat incontinence
- **5** Volitional detrusor contraction when voiding
- 6 Post-void residual (PVR) urine volume <100 mL

	Patient meets all 6 criteria	Patient does not meet all 6 criteria	P value
N=95	72	23	
Mean daily preoperative <b>pad use</b> (range)	2.6 (1-4)	4.4 (1-9)	<.05
Mean daily preoperative <b>pad weight, g</b> (range)	131 (10-280)	520 (80-1200)	<.05
Mean daily postoperative <b>pad use</b> (range)	<b>0.6</b> (0-6)	2.4 (0-7)	<.05
Mean daily postoperative <b>pad weight, g</b> (range)	<b>16</b> (0-310)	201 (0-800)	<.05
Pad reduction percentage	77%	46%	
Pad weight reduction percentage	88%	61%	
Patient satisfaction	92%	30%	

Mean follow-up: 28 months

1. Sturm RM, Guralnick ML, Stone AR, et al. Comparison of clinical outcomes between "ideal" and "nonideal" transobturator male sling patients for treatment of postprostatectomy incontinence. *Urology*. 2014 May;83(5):1186-8.

Prior to use, please review the Instructions for Use for a complete listing of indications, contraindications, warnings, precautions and potential adverse events.

CAUTION: Federal Law restricts this device to sale by or on the order of a physician. This product is only intended for physicians trained in its use.

INDICATIONS: The AdVance™ male sling system is intended for the placement of a suburethral sling for the treatment of male stress urinary incontinence (SUI).

CONTRAINDICATIONS: Patients with urinary tract infections, blood coagulation disorders, a compromised immune system or any other condition that would compromise healing, with renal insufficiency, and upper urinary tract relative obstruction.

POTENTIAL ADVERSE EVENTS: Extrusion, erosion through the urethra or other surrounding tissue, migration of the device from the desired location, fistula formation and inflammation. Rx Only

