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Dynamic Maximum Urethral Closure Pressures Measured by High Resolution Manometry Increase Markedly after Sling Surgery

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Abstract

Introduction and Hypothesis: The premise of midurethral sling (MUS) surgery is to apply a tension-free vaginal tape in the midurethra that does not constrict the urethra at rest but stabilizes the urethra and prevents downward descent and opening of the urethra during stress maneuvers, but current technology has limitations in measuring urethral pressures during dynamic conditions. Our objective was to describe the change in maximum urethral closure pressures (MUCPs) after MUS surgery using an 8-French high resolution manometry system (HRM) that can measure urethral pressures during cough and strain maneuvers (ManoScan® ESO, Given Imaging) without migration or withdrawal limitations.

Methods: We measured rest, cough, and strain MUCPs in 26 women before and after retropubic or transobturator MUS for stress urinary incontinence using the HRM system.

Results: The objective success rate after MUS was 92.3% based on postoperative cough stress testing. Mean resting MUCPs measured by HRM did not change after surgery (59.3 preop vs 59.7 cm H₂O postop) ($p = 1.0$). Mean cough MUCPs measured by HRM increased from 36.9 to 100.7 cm H₂O ($p < 0.001$), and strain MUCPs increased from 35.0 to 92.7 cm H₂O ($p < 0.001$).

Conclusions: Advanced HRM technology to measure MUCPs under cough and strain conditions without withdrawal techniques provides new insights into the continence mechanism

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Authors' contributions

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after tension-free MUS: MUCPs do not change at rest but do increase significantly during cough and strain maneuvers.

Brief summary

Mean cough and strain maximum urethral closure pressures in women with stress urinary incontinence increase markedly after midurethral sling surgery

Keywords

High resolution manometry; maximum urethral closure pressures; stress urinary incontinence; midurethral sling surgery

Introduction

Urinary incontinence affects 16% of adult women [1]. Stress urinary incontinence (SUI), the involuntary loss of urine on effort or physical exertion such as cough and strain [2], can be treated with pelvic floor muscle exercises, pessaries, and surgeries, with surgeries being the most successful treatment [3]. Approximately 260,000 stress incontinence surgeries are performed each year in the U.S. [4]. The most frequently performed surgical treatments for SUI are mesh midurethral slings (MUS), which have a success rate of approximately 80% [5].

The premise of MUS surgery is to apply a tension-free vaginal tape in the midurethra that does not constrict the urethra at rest but stabilizes the urethra and prevents downward descent and opening of the urethra during stress maneuvers. Indeed, most studies report that resting maximum urethral closure pressures (MUCPs) do not change significantly after MUS surgery [6–8]. It is more difficult to assess and understand the function of the urethra during typical stress maneuvers like cough and strain. Current technologies are hindered by slow response times or the need for withdrawal techniques. Instruments that measure a single point of pressure in the urethra can be displaced from the maximum pressure zone with a cough or strain and therefore changes may just be artifact. Pressure transmission ratios measure the ratio of urethral pressure to bladder pressure with repetitive coughs while withdrawing a microtip catheter through the urethra. These ratios have generally been shown to increase after SUI surgery [9], but in several decades of existence, these ratios have never become a widespread or clinically useful tool. The search for a simple tool that can measure urethral pressures during dynamic conditions and help us understand successful or failed treatments remains elusive.

A high resolution manometry (HRM) system (ManoScan® ESO, Given Imaging) has recently been shown to be well tolerated in the bladder and urethra and to correlate well with water-perfusion urethral pressure profilometry at rest [10]. The potential advantage of this technology is that it has a fast 25-millisecond response time and can measure circumferential pressures at many points along the urethra at once without withdrawal techniques. Because of its ability to capture MUCPs during cough and strain, it may be clinically useful in evaluating outcomes after MUS for SUI. Although resting MUCP's may not change after

surgery, the more important question is whether cough and strain MUCPs increase after MUS, and HRM technology can measure this.

The purpose of this investigation is to determine if this new HRM technology allows us to better understand how MUS affect urethral pressures under dynamic conditions. Specifically, we sought to measure changes in rest, cough, and strain MUCPs after MUS. We hypothesized that the MUS functions by increasing urethral pressures during cough and strain maneuvers but not at rest.

Materials and Methods

The HRM system we used is a pediatric esophageal catheter and pressure-sensing software produced by Given Imaging for use in gastroenterology (ManoScan® ESO, Given Imaging) [11]. This catheter is 2.75mm in diameter (approximately 8-French) and has circumferential pressure sensors along 26.5cm of its length. The miniature solid state sensors are 4mm long with 2mm of active sensing area separated by 3mm of flexible molding giving a 7.5mm on center spacing between each of the 36 sensors. Each individual sensor has 16 pressure sensitive segments circumferentially distributed around it (Figure 1). Computer processing of the signal comes from the pressure sensing elements and calculates average circumferential pressures. Pressures between sensing elements are interpolated, and the data are presented on a color graphic display (Figure 2).

Women with primary or recurrent urinary incontinence who were scheduled to undergo full length MUS at our institution were recruited to have HRM bladder and urethral pressure measurements before and 6 weeks after surgery. The study was approved by our Institutional Review Board, and written consent was obtained. All subjects were at least 18 years of age, not pregnant, without evidence of a UTI, and without pelvic organ prolapse beyond the hymen. Women with prior prolapse or incontinence surgeries were not excluded. All subjects had positive standing cough stress tests prior to surgery. Each subject completed demographic information including age, height, weight, parity, medical history, and surgical history as well as the Incontinence Severity Index (ISI, scale 0 to 12) [12], Incontinence Impact Questionnaire Short Form (IIQ-7, scale 0 to 100), and Urogenital Distress Inventory Short Form (UDI-6, scale 0 to 100) questionnaires [13]. Question 3 of the UDI-6 was specifically used to capture subjective SUI: "Do you experience and, if so, how much are you bothered by urine leakage related to activity, coughing, or sneezing?" Questionnaires were repeated 6 weeks after surgery.

Prior to obtaining pressure measurements, subjects' bladders were drained with a sterile catheter and then filled with 250ml sterile water or with the maximum volume she could comfortably hold for 15 minutes if this was less than 250ml. This catheter was removed, and the HRM catheter was placed such that the tip was approximately 5cm inside the bladder. Bladder and urethral pressures were obtained after 10 seconds of rest, during 3 coughs, and during 3 strain maneuvers. For strain maneuvers, subjects were instructed to bear down or push as though trying to have a baby or a bowel movement. MUCPs were calculated by subtracting the maximum bladder pressures 2cm proximal to the urethrovesical junction from the maximum urethral pressure anywhere along the urethra during each maneuver.

Because each 4mm sensor has a 2mm sensing area and there are 3mm spaces between sensors, there are 5mm sections along the urethra that are not measured with this system. Our previous experiments, in which the catheter was moved several millimeters and pressure recordings repeated, demonstrated that the location of this non-measured area does not appear to affect maximum pressure measures [10].

The primary endpoint was change in MUCP during dynamic maneuvers (cough and strain) after compared to before MUS surgery. We pooled retropubic and transobturator MUS because these surgeries, with only small differences in placement and outcomes, are thought to function similarly [5]. Pre- and postoperative MUCPs were compared using paired t-tests, questionnaire scores were compared using paired Wilcoxon tests, and the change in the percent of patients with positive cough stress tests before and after surgery was tested with chi-square. Tests were adjusted for multiple comparisons using the Bonferroni method, and $p < 0.01$ was considered significant. We recruited 26 subjects in order to have 80% power to detect a mean change of 14cm H₂O (standard deviation 20 cm H₂O) during cough and strain after MUS based on previous studies [14] with an 85% success rate after MUS surgery [15] and 20% drop-out rate. Analyses were performed using R (<http://r-project.org>). In a secondary analysis, we defined composite surgical success as objective negative postoperative cough stress test (CST) and subjective improvement in ISI score, and we compared composite success rates in subjects with increased postoperative cough and strain MUCPs to subjects with decreased postoperative MUCPs using a Fisher's exact test.

Results

26 women were included in this study, 20 of whom had retropubic and 6 of whom had transobturator MUS. The median age was 53.5 years (range 38–77), median BMI 27.9 kg/m² (range 20.8–37.2), and median vaginal parity 2 (range 0–5). 7 subjects (26.9%) had at least one prior continence surgery. Pre- and postoperative continence severity, quality of life measures, cough stress tests, and MUCPs are shown in Table 1. All continence severity and quality of life measures improved significantly after surgery. All subjects had positive cough stress tests before surgery, and 2 had positive stress tests 6 weeks after surgery (objective success rate of 92.3%). ISI scores improved in 96.2% (25/26) of subjects. Composite success (negative CST and improvement in ISI) was 88.5% (23/26).

Figure 3 shows pre- and postoperative MUCPs after MUS surgery for rest, cough, and strain for each subject. Resting MUCPs generally did not change after surgery while cough and strain MUCPs increased in the vast majority of subjects. Cough and/or strain MUCPs decreased after surgery in 3 subjects; the composite success rate for these subjects was 66.7% (2/3), compared to 91.3% (21/23) subjects in whom cough and strain MUCPs increased. The difference in success rates in the two groups was not statistically different ($p = 0.3$), but this study was not powered for this secondary analysis.

Mean resting MUCPs were not different after surgery (59.3 preop vs 59.7 cm H₂O postop) ($p = 1.0$). Mean cough MUCPs increased from 36.9 to 100.7 cm H₂O ($p < 0.001$) and strain MUCPs increased from 35.0 to 92.7 cm H₂O ($p < 0.001$) (Table 1, Figure 4).

When we analyzed retropubic and transobturator MUS separately, the trends remained similar to what were seen for all subjects. Retropubic MUS subjects had enough subjects (20) to meet our power calculation because no subjects dropped out of the trial, and, similar to our primary endpoint for all subjects, cough and strain MUCPs increased significantly with p-values < 0.001 while rest MUCPs did not change significantly (p = 0.2). With only 6 subjects who underwent transobturator MUS, the same trends were seen, but we were not powered to show statistical significance in this sub-group: change in cough and strain nearly met significance (p = 0.05 and p = 0.02, respectively), while change in rest MUCP was not significant (p = 0.16).

We included women with prior continence surgeries in this study because they were surgical failures and candidates for MUS surgery regardless of what surgeries were done in the past. When we analyzed the subjects with and without prior continence surgeries separately, we saw similar results for subjects with and without prior surgeries with only slightly stronger results in those without prior surgeries (data not shown).

Discussion

The major finding of our study is that the mean MUCPs during cough and strain measured by a high resolution manometry system increase markedly (60 cm H₂O) and significantly after MUS surgery. Postoperative cough and strain MUCPs were similar to MUCPs seen in continent women [10]. We found this high mean increase even including those subjects who did not improve after surgery. In contrast to the change after surgery in MUCP with cough or strain, the resting MUCP did not change. These results support our hypothesis that MUS does not appreciably affect the urethra at rest but has a profound effect on the urethra with cough and strain.

No change in resting pressure after MUS is consistent with most previous studies using other technologies [6–8,16]. The increase we observed in MUCPs after surgery during dynamic conditions is also consistent with studies that have found increased pressure transmission ratios after surgery [6,9], but the magnitude of the effect can only be appreciated with this non-withdrawal technique.

Most cough and strain MUCPs increased after surgery, and the few subjects in whom MUCPs did not increase had somewhat worse though not statistically different postoperative outcomes. This suggests that this technology may be clinically useful for measuring success after MUS and may even help clinicians determine whether a sling revision, sling tightening, or an additional sling is likely to improve continence in patients who remain incontinent after MUS surgery. Since this is a preliminary pilot study using this new technology before and after surgery, and we had few failures, additional studies with sufficient power would be needed to determine its clinical utility in differentiating success from failures.

HRM technology is an improvement over other urodynamic measurements because of its fast response rate and multiple pressure-sensing segments that assure it can register millisecond pressure changes during coughs and capture maximum pressures along the length of the urethra even if the catheter changes position. There remain some limitations in

its accuracy because its inherent rigidity and circumference can produce artifact. Nonetheless, it has allowed us to dramatically demonstrate the mechanism of action of MUS to improve continence and may prove to be clinically useful in diagnosing and assessing treatment success in women with SUI.

Advanced HRM technology to measure MUCPs under cough and strain conditions without withdrawal techniques provides new insights into the continence mechanism after tension-free MUS. MUCPs do not change at rest, however marked increases occur during cough and strain maneuvers. Future studies are planned to evaluate this technology as an intraoperative tool to adjust sling tensioning or understand surgical failures

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Figure 1. High resolution manometry (HRM) catheter. 2.75mm diameter with 4mm copper pressure sensors spaced 3mm apart (left). Each sensor has 16 circumferential pressure sensing foci (right).

Figure 2. High resolution manometry (HRM) system output display. Y-axis is distance along the catheter and x-axis is time. Color display pressures are shown during 3 coughs (left) and 3 strains (right) measured in a continent subject.

Figure 3. Pre- and postoperative maximum urethral closure pressures (MUCPs) after midurethral sling (MUS) surgery during rest, cough, and strain for 26 individual subjects.

Figure 4. Mean change in maximum urethral closure pressures (MUCPs) after midurethral sling (MUS) surgery during rest, cough, and strain with standard error bars (n = 26).

Table 1.

Demographics, maximum urethral closure pressures, continence severity, and quality of life measures (n = 26)

	Preoperative	Postoperative	delta	p-value
Rest MUCP (cm H ₂ O) **	59.3 (4.3)	59.7 (3.8)	+ 0.4	p = 1.0 ^t
Cough MUCP (cm H ₂ O) **	36.9 (4.1)	100.7 (12)	+ 63.8	p < 0.001 ^t
Strain MUCP (cm H ₂ O) **	35 (4.3)	92.7 (10)	+ 57.7	p < 0.001 ^t
ISI *	8 (4–12)	0 (0–8)	– 8	p < 0.001 ^w
UDI-6 *	50 (22.2–72.2)	11.1 (0–55.6)	– 38.9	p < 0.001 ^w
UDI-6 Q3 *	3 (1–3)	0 (0–2)	– 3	p < 0.001 ^w
IIQ-7 *	33.3 (4.8–100)	0 (0–47.6)	– 33.3	p < 0.001 ^w
Positive cough stress test (n,%)	26 (100%)	2 (8%)	– 92%	p < 0.001 ^c

ISI: Incontinence Severity Index. IIQ-7: Incontinence Impact Questionnaire Short Form. UDI-6: Urogenital Distress Inventory Short Form. UDI-6 Q3: question #3 of UDI-6.

* median (range).

** mean, SE.

^t: paired t-tests.

^w: wilcoxon tests.

^c: chi square.