Preoperative voiding detrusor pressures do not predict stress incontinence surgery outcomes

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Abstract

Introduction and hypothesis The aim of this study was to determine whether preoperative voiding detrusor pressures were associated with postoperative outcomes after stress incontinence surgery.

Methods Opening detrusor pressure, detrusor pressure at maximum flow ($p_{det \, Q_{max}}$), and closing detrusor pressure were assessed from 280 valid preoperative urodynamic studies in subjects without advanced prolapse from a multicenter randomized trial comparing Burch and autologous fascia sling procedures. These pressures were compared between subjects with and without overall success, stress-specific success, postoperative detrusor overactivity, and postoperative urge incontinence using independent sample $t$ tests.

Results There were no clinically or statistically significant differences in mean preoperative voiding detrusor pressures in any comparison of postoperative outcomes.

Conclusions We found no evidence that preoperative voiding detrusor pressures predict outcomes in women with stress predominant urinary incontinence undergoing Burch or autologous fascial sling procedures.

Keywords Closing detrusor pressure · Opening detrusor pressure · Urinary stress incontinence · Urinary urge incontinence · Urodynamics · Voiding detrusor pressure
Introduction

Preoperative urodynamic studies (UDS) are commonly performed prior to stress incontinence (SUI) surgery. If urodynamic measures could predict overall success, stress-specific success, postoperative detrusor activity (DO), or the need for postoperative urge urinary incontinence (UUI) treatment, then these measures might be useful for preoperative counseling.

Recent studies suggest that voiding detrusor pressure values during preoperative UDS predict postoperative outcomes. Digesu et al. found that postoperative incontinent women after Burch colposuspension had lower preoperative opening detrusor pressures (ODP) and closing detrusor pressures (CDP) than continent women [1]. Similarly, in a study of 35 women with urodynamic mixed incontinence, defined as urodynamic stress incontinence (USI) and DO, investigators found that higher ODP during preoperative urodynamic testing predicted persistent DO after tension-free vaginal tape (TVT) surgery. ODP was also higher in subjects with postoperative overactive bladder (OAB), but this difference was not statistically significant [2].

We hypothesized that preoperative voiding detrusor pressures would be associated with postoperative outcomes after SUI surgery in women with stress predominant SUI. Specifically, we sought to determine whether ODP, detrusor pressures at maximum flow ($p_{detQ_{max}}$), or CDP were associated with, or predictive of, overall success, stress-specific success, postoperative DO, and postoperative treatment for UUI. We also evaluated whether preoperative detrusor after-contractions were more common in women who were treated for postoperative UUI.

Materials and methods

SISTEr trial and subjects

This study is a secondary analysis of data from the Stress Incontinence Surgical Treatment Efficacy Trial (SISTEr). In the original trial, 655 women were enrolled from February 2002 to June 2004. Details of the SISTEr study methods have been published previously [3] and are briefly outlined here. Inclusion criteria for enrollment included: (1) predominant SUI: stress>urge score on the Medical, Epidemiological and Social Aspects of Aging Questionnaire (MESA) [4], (2) positive stress test (observed leakage from the external urethral meatus coincident with a cough or Valsalva maneuver with a bladder volume $\leq$ 300 ml), (3) urethral hypermobility as evidenced by resting or maximum straining Q-tip angle $>30^\circ$ [5], (4) maximum cystometric capacity (MCC) $\geq$ 200 ml, and (5) postvoid residual (PVR) $<150$ ml. All study procedures were approved by the institutional review board of each participating clinical center and the Biostatistical Coordinating Center with written informed consent obtained from all women prior to enrollment.

Urodynamic studies and quality control

Baseline urodynamic testing (free uroflowmetry, filling cystometry, and pressure-flow studies) were performed on all participants prior to surgery based on a standardized protocol implemented by all 20 urodynamic testers at the nine continence treatment centers. The SISTEr UDS protocol complied with terminology from the Standardization Committee of the International Continence Society [6] and technical recommendations from the Good Urodynamic Practice guidelines [7]. Standardized interpretation guidelines were used by on-site (local) physician reviewers. The details of our specific UDS Protocol and Interpretation Guidelines are available on the Urinary Incontinence
Treatment Network website at http://www.uitn.net/resources forphysicians.htm. A Urodynamics Working Group supervised all aspects of urodynamic protocol development, procedural performance, and interpretation reliability [8, 9].

Pressure-flow studies

After standing filling cystometry, study participants sat to void for pressure-flow studies (PFS). For PFS pressure data to be considered valid, they had to satisfy the following criteria: (1) legible signals, (2) subject voided, (3) bladder pressure (p-ves) and abdominal pressure (p-abd) measuring systems properly functioning at PFS baseline and Q-max, (4) p-det at PFS baseline greater than −5 cm H2O, (5) p-det at PFS baseline no more than 15 cm H2O greater than p-det at MCC (insuring that change of position and transducer adjustment did not cause artifact in pressure measurement), (6) prevoid cough spikes demonstrated at least 70% agreement between the p ves and p-abd signals (smaller spike at least 70% of the larger spike) [10] to demonstrate that the bladder and abdominal pressure transducers were functioning well, and (7) the local reviewer found no other reason to invalidate the study. For this analysis to be consistent with our other plausibility criteria, we also excluded all PFS with voiding detrusor pressure measures (ODP, p-detQ-max, or CDP) less than −5 cm H2O.

Preoperative voiding detrusor pressure measurements

We followed the methods used by Panayi et al. 2009 [2] to determine ODP and CDP, defining them as the detrusor pressure 1 s prior to the onset of flow and 1 s prior to cessation of flow, respectively, as recorded by the uroflowmeter. For ODP, to evaluate the appropriateness of the 1-s delay for the urine to flow through the urethra, emerge from the patient, and register on the uroflowmeter, we also recorded the highest detrusor pressure in the 1 to 5 s preceding the onset of flow on the uroflowmeter and how many seconds prior to flow it occurred. In more than 90% of the signals, the highest detrusor pressure in the 1 to 5 s preceding the onset of flow was equal to ODP at 1 s prior to onset of flow, and therefore, we only report ODP at 1 s for this analysis. Detrusor pressure at maximum flow (p-detQ-max) was recorded from the Q-max auto-annotation on each UDS signal. The reviewers were blinded to all outcome measures when the urodynamic data were extracted.

After-contractions are sometimes found as a pressure increase after flow ceases at the end of micturition [11]. The significance of this event is not understood, and there is no universally accepted definition. For the purpose of this review, we defined after-contractions as any detrusor pressure within 5 s of the end of flow greater than any detrusor pressure during flow.

Preoperative voiding detrusor pressure quality control

Fifty arbitrarily selected urodynamic signals were jointly reviewed by the first and second authors for education, training, and clarification of definitions. Subsequently, a statistician (HL) randomly selected 22 signals for independent review to evaluate inter-rater reliability. A priori, 80% agreement was determined to be acceptable inter-rater reliability, and pressure measures with a numerical difference of 10% or less were considered the same since ≤10% differences in pressure are seldom clinically important. It was determined a priori that, if acceptable reliability could be established, a single interpreter would perform the remainder of the data extraction.

Overall treatment success and stress-specific success

Success was evaluated at 24 months. Overall treatment success was defined as no self-reported SUI symptoms, a negative provocative standardized 300-ml stress test, no re-treatment for SUI, less than 15 g of urine on a 24-h pad test, and no leakage on a 3-day diary. Stress-specific success was defined as no SUI symptoms, a negative stress test, and no SUI re-treatment [12].

Detrusor overactivity

DO was defined as a urodynamic observation of involuntary detrusor contractions during the filling phase [6]. Preoperative DO was DO during the baseline UDS and postoperative DO was DO during UDS at the 24-month urodynamic visit. Since preoperative DO is likely to be associated with postoperative DO, in our analysis of postoperative DO, we performed two analyses, one excluding and one including subjects with preoperative DO.

Treatment of postoperative urge incontinence

Treatment of postoperative UUI was defined as treatment of clinically diagnosed new-onset or persistent UUI after the 6-week follow-up visit with any clinically acceptable treatment for OAB. This parameter was assessed at 3, 6, 12, 18, and 24 months.

Preoperative pelvic organ prolapse

Pelvic organ prolapse was assessed during preoperative physical examination using POPQ definitions [12, 13]. For this secondary analysis, we excluded subjects with pelvic organ prolapse past the introitus (stage 3 or 4) because prolapse could artificially increase pressures by causing urethral obstruction, and exclusion of prolapse subjects is consistent with previous literature evaluating the role of preoperative voiding detrusor pressures [1, 2].
Statistics

The distributions of preoperative voiding detrusor pressure measures were assessed for normality. There was no strong evidence of deviation from normality, so independent sample *t* tests were used to test for differences in detrusor pressures by overall and stress success status, DO status, and UUI treatment status. Based on the number of subjects with and without postoperative UUI, there was at least 80% power to detect a difference in mean ODP values of 5 cm H$_2$O assuming that mean ODP of subjects with postoperative UUI was 13 with a standard deviation of 10 [14]. Pearson chi-square tests were used to test for relationships between categorical variables (e.g., after-contractions). Analyses were carried out using SAS statistical software Version 9.2 (SAS Institute, Inc., Cary, NC).

Results

SISTEr trial and subjects

The 655 study subjects had a mean age of 52 years (standard deviation (SD) 10, range 28 to 81) and a mean BMI of 30 kg/m$^2$ (SD, 6 kg/m$^2$). The ethnic distribution was: 73% non-Hispanic White, 11% Hispanic, 7% non-Hispanic Black, and 9% non-Hispanic Other.

Pressure-flow studies and detrusor pressure quality control

The 655 subjects all underwent preoperative UDS. Seventy-eight UDS had invalid or implausible cystometrogram (CMG) studies, and 191 had invalid or implausible PFS as described previously [9]. We excluded an additional 13 studies that were not captured on standardized electronic equipment, one study from which data were not properly extracted, nine studies in which the pressure catheter fell out before $Q_{\text{max}}$, and 46 signals with implausible pressures (less than $-5$cm H$_2$O) at ODP ($n=14$), $p_{\text{det}}Q_{\text{max}}$ ($n=8$), or CDP ($n=23$). We subsequently excluded 46 of these 326 high-quality PFS because the subjects had stage 3 or 4 prolapse. This left 280 PFS studies that met our predetermined quality control measures for this analysis.

Evaluation for potential selection bias

An analysis was performed to determine if there was possible selection bias introduced by excluding subjects with invalid or implausible UDS subjects. Table 1 compares the postoperative outcome rates in the 280 subjects with valid UDS who were included in this analysis to the postoperative outcome rates in the 269 subjects that were excluded (549 subjects in SISTEr had stage 2 prolapse or less, and not all subjects had all outcome measures recorded). No differences between those included and excluded were detected.

Inter-rater reliability

Inter-rater agreement was 95% (21/22) for ODP, 82% (18/22) for $p_{\text{det}}Q_{\text{max}}$, 91% (20/22) for CDP, and 91% (20/22) for the presence or absence of after-contractions. This inter-rater agreement met our reliability criteria, and most of the data extraction was then obtained by a single interpreter (AK).

Preoperative voiding detrusor pressures

Mean and standard deviation (SD) preoperative voiding detrusor pressures for all 280 studies were ODP 13.5 (9.7), $p_{\text{det}}Q_{\text{max}}$ 19.8 (11.3), and CDP 18.3 (15.8) cm H$_2$O.

Overall treatment success and stress-specific success

Table 2 shows the mean preoperative voiding detrusor pressures in subjects with overall success and failure and stress-specific success and failure. There were no significant differences in mean ODP, $p_{\text{det}}Q_{\text{max}}$, or CDP between the success and failure groups.

Postoperative detrusor overactivity

In SISTEr, 9.3% (60/645) of subjects had preoperative DO and 8.9% (45/506) had postoperative DO. Table 3 presents mean preoperative voiding detrusor pressures grouped by postoperative DO status in subjects stratified by the presence or absence of preoperative DO. In subjects with preoperative USI or preoperative urodynamic mixed incontinence (USI and DO), there was no significant difference

| Table 1 Postoperative outcome rates for included and excluded subjects |
|-----------------------------------------------|
|                  | Included subjects | Excluded subjects | *p* value |
|------------------|-------------------|------------------|-----------|
| Overall success  | 38.5% (85/221)    | 35.1% (74/211)   | 0.47      |
| Stress-specific success | 52.9% (120/227) | 57.3% (129/225) | 0.34      |
| Postoperative DO | 12% (25/207)      | 7.5% (16/214)    | 0.11      |
| Treatment for postoperative UUI | 19.6% (55/280)  | 22.7% (61/269)   | 0.38      |

DO detrusor overactivity, UUI urinary urge incontinence
in voiding detrusor pressures in subjects with and without postoperative DO.

Treatment for postoperative urge incontinence

Of the 55 subjects treated for postoperative UUI in this study, all 55 were treated with medications, 11 were also treated with biofeedback or behavioral training, and four were treated with other therapies. Table 4 shows the mean preoperative voiding detrusor pressures for subjects who were and were not treated for postoperative UUI. No preoperative voiding detrusor pressures were statistically different.

After-contractions

Twenty-six percent (8/31) of subjects with after-contractions received treatment for postoperative UUI compared to 19% (47/249) of those without after-contractions. This rate difference was not statistically different ($p=0.36$).

Discussion

In this analysis of preoperative urodynaminc data from a large stress incontinence surgery trial, we found no voiding detrusor pressure parameter that was associated with overall success, stress-specific success, DO, or treatment for UUI after surgery. This implies that none of these measures could be used clinically to predict which patients would have these postoperative outcomes.

There was no difference in the prevalence of after-contractions when we compared those who received treatment for postoperative UUI to those who did not, leaving the significance of this finding still not understood [11].

We acknowledge that lack of significance of superiority tests does not necessarily imply lack of difference. However, this study had an adequate sample size ($n=280$) to detect large, clinically relevant differences, so it is unlikely that low power was the reason for not detecting differences in preoperative voiding detrusor pressures by success, DO, or UUI treatment. For instance, based on the number of subjects with and without postoperative UUI ($n=225$ vs $n=55$), there was at least 80% power to detect a difference in mean ODP values of 5 cm H$_2$O assuming that mean ODP of subjects with postoperative UUI was 13 with a standard deviation of 10 [14].

Unlike previous investigators who found that subjects with postoperative continence had ODP values 9 cm H$_2$O higher than those with postoperative incontinence, [1] we found that our subjects with postoperative continence had ODP values 1 to 2 cm H$_2$O lower than those with postoperative incontinence. Our studies were similar, and both had over 200 subjects, but they differ in the following ways: (1) our study included Burch and sling and their study had Burch procedures only, (2) our incontinence outcome measures were more clearly defined, and (3) we included the 9% of subjects with preoperative DO, and they excluded preoperative DO. Given these conflicting results, in different directions, we think it is highly unlikely that ODP can predict postoperative incontinence.

### Table 2 Preoperative voiding detrusor pressures in success and failure groups

|                        | Overall success ($n=85$) | Overall failure ($n=136$) | $p$ value | Stress-specific success ($n=120$) | Stress-specific failure ($n=107$) | $p$ value |
|------------------------|--------------------------|---------------------------|-----------|---------------------------------|---------------------------------|-----------|
| ODP                    | 11.3 (9.5)               | 13.6 (8.6)                | 0.07      | 12.3 (9.6)                      | 13.6 (8.9)                      | 0.31      |
| $p_{detQ_{max}}$       | 19.1 (11.8)              | 19.7 (11.2)               | 0.70      | 19.8 (11.5)                     | 19.4 (11.4)                     | 0.79      |
| CDP                    | 16.0 (12.9)              | 19.0 (15.7)               | 0.14      | 16.8 (13.0)                     | 19.1 (16.4)                     | 0.24      |

Values are reported as mean in centimeter H$_2$O (SD)

$ODP$ opening detrusor pressure, $p_{detQ_{max}}$ detrusor pressure at maximum flow, $CDP$ closing detrusor pressure

### Table 3 Preoperative voiding detrusor pressures in urodynamic stress incontinent subjects and postoperative DO

|                        | No preoperative DO ($n=167$) | Preoperative DO ($n=20$) |
|------------------------|------------------------------|--------------------------|
|                        | No postoperative DO ($n=152$) | Postoperative DO ($n=15$) | $p$ value |
|                        | No postoperative DO ($n=11$)  | Postoperative DO ($n=9$)  | $p$ value |
| ODP                    | 12.4 (9.3)                   | 11.4 (7.4)                | 0.68      | 17.0 (11.3)                    | 15.8 (8.3)                | 0.79      |
| $p_{detQ_{max}}$       | 19.6 (12.0)                  | 19.3 (8.1)                | 0.91      | 21.5 (12.7)                    | 22.4 (11.7)               | 0.86      |
| CDP                    | 16.8 (13.6)                  | 19.9 (13.9)               | 0.41      | 16.3 (12.5)                    | 21.4 (16.3)               | 0.43      |

Values are reported as mean in centimeters H$_2$O (SD)

$ODP$ opening detrusor pressure, $p_{detQ_{max}}$ detrusor pressure at maximum flow, $CDP$ closing detrusor pressure, $DO$ detrusor overactivity
Also, unlike previous investigators who reported that subjects with persistent postoperative DO had ODP values 17 cm H2O higher than subjects without persistent postoperative DO, [2] we found that subjects with persistent postoperative DO had ODP values 1 cm H2O lower than subjects without persistent postoperative DO. Our studies were similar in that both were small (35 and 20 subjects, respectively), but they differed in that our study was Burch and autologous fascial sling surgeries, and their study was TVTs. However, if these preoperative urodynamic variables could reliably predict surgical outcomes, we think they should predict outcomes for both traditional incontinence surgeries and the newer midurethral slings.

In SISTEr, only 9% of subjects had preoperative DO, but 93% had a positive response to at least one urge question on MESA [15], therefore, we consider our subjects to have anywhere between a 9% and 93% mixed incontinence rate. Because of the ambiguity of the mixed incontinence definition, we decided to concentrate our study on the broader, more generalizable question of preoperative voiding detrusor pressure predictors in all patients undergoing SUI surgery rather than only a small subgroup of subjects with preoperative urodynamic mixed incontinence. Therefore, we also studied 187 subjects without preoperative DO and found similar preoperative voiding detrusor pressures for those with and without postoperative DO.

We chose to evaluate postoperative UUI using the two most unambiguous measures: DO on UDS and treatment for UUI. DO on UDS was the outcome chosen by previous investigators and most likely reflects the most severe UUI patients. Treatment for UUI has a clear definition in our study and arguably is the most clinically relevant overactive bladder condition. We recognize that both of these definitions have limitations. DO on UDS represents events during just a single filling cycle and is not sensitive for the detection of UUI. Treatment for UUI can be influenced by unmeasured variability in clinician and patient preferences with respect to treatment of any UUI that is present. We found no difference in preoperative voiding detrusor pressures for postoperative DO or postoperative treatment for UUI. Given all these results, we think it is highly unlikely that preoperative voiding detrusor pressures can predict postoperative DO and urge incontinence.

The strengths of this study include the large subject population, the multiple sites which improve generalizability, and our careful, standardized urodynamic quality control process. This quality control process excludes poor quality signals. Most urodynamic publications do not perform quality control on their signals and do not evaluate and exclude signals with poor pressure transmission concordance on prevoid cough spikes. A quality control process, especially for pressure-flow studies, which are the most prone to quality challenges, may result in the exclusion of a large number of studies and has the potential to introduce selection bias, but it is unlikely in this study because our postoperative outcome rates were the same in our included and excluded subjects. Therefore, we think insisting on the highest quality, valid and plausible studies strengthens our results and conclusions rather than weakens them by limiting sample size.

In this same study population, other urodynamic predictors were not found to be good predictors of success or failure. [16] It is likely that during preoperative counseling, clinicians will continue to rely on simple clinical factors to predict postoperative outcomes, such as a patient’s age, as a predictor of failure and postoperative UUI. [17, 18] Our results suggest no change in this practice.

In conclusion, we did not find that voiding detrusor pressures during preoperative urodynamic studies predict postoperative success, stress-specific success, DO, or treatment of UUI in a large sample of women undergoing Burch or fascial sling procedures for SUI. We do not recommend routine measuring of these urodynamic measures for predictive or prognostic purposes in women undergoing preoperative urodynamics before stress incontinence surgery.

Conflicts of interest None.

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