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Toledo L G¹, Cabral B H², Casella C L², Politi D E², Cardoso S N², Mello F F², Glina S² **1.** Santa Casa of São Paulo Medical School and Ipiranga Hospital, **2.** Ipiranga Hospital, São Paulo, Brazil

PROGNOSTIC VALUE OF URETHRAL MOBILITY AND VALSALVA LEAK POINT PRESSURE FOR FEMALE TRANSOBTURATOR SLING PROCEDURE

Hypothesis / aims of study

The transobturator sling (TOT) procedure is a well-established technique for the surgical treatment of stress urinary incontinence. Nonetheless, the mechanisms through which continence is obtained, and thereafter the prognostics determinants, are incompletely understood. The prognostic significance of urethral mobility and Valsalva leak point pressure (VLPP), which is a marker of intrinsic sphincter deficiency, is still debated.^(1,2) We analyzed the influence of urethral mobility and Valsalva leak point pressure on postoperative outcomes.

Study design, materials and methods

A prospective cohort was conducted including 66 patients submitted to TOT from March 2006 to May 2009. Urethral hypermobility was defined as mobility \geq 30° on Q-tip test, and Valsalva leak point pressure (VLPP) was classified as grater than 60 cmH₂O or 60 and less on preoperative urodynamics. Postoperative objective success was defined as absence of any urinary loss during full bladder standing Valsalva maneuver and no need of pads, while subjective success was achieved when the patients considered themselves much better or cured, the level of satisfaction was \geq 8 (according to a visual analogic scale from 0 to 10) and there was no report of stress incontinence. Quality of life was analysed by ICIQ-SF. Statistical analysis was accomplished and the results rendered significant if p< 0.05.

Results

Among the 73 patients undergoing the TOT procedure, seven were excluded due to incomplete preoperative data (VLPP or urethral mobility) and 66 patients were included for analysis. Overall, 94% and 79% of our patients were cured according to objective and subjective success criteria, respectively. There were two cases of mesh extrusion to the vaginal wall and one case of erosion to the bladder neck. None of them presented local severe infection and were submitted to the correcting surgery electively. They were all healthy and continent on completion of follow-up.

Baseline characteristics (age, BMI, gestational status, preoperative ICIQ, type of continence and previous incontinence surgery) had similar distribution when different groups were compared. These possible prognostic factors were individually analysed and had no influence on postoperative outcomes. Mean preoperative ICIQ was 15.5 and 16.7 in patients with low and high mobility (p= 0.2 - Mann-Whitney's test), respectively.

Urethral hypermobility was an important determinant of objective success (p= 0.04), as just one patient with mobility \geq 30° failed therapy. Subjective success rate was also higher in this group, but the difference did not reach statistical significance (p= 0.07). Conversely, VLPP had no role as prognostic factor, either when analysed as categorical variable or when numeric VLPP values were compared. In the objective success analysis, mean VLPP was 77 cmH₂0 and 95 cmH₂0 in successful and unsuccessful cases, respectively (p= 0.21 – Mann-Whitney's test). In the subjective success analysis, patients who failed therapy had mean VLPP similar to those who succeed (79.3 cmH₂0 *versus* 73.6 cmH₂0, respectively) (p= 0.31 – Mann-Whitney's test).

There was no statistical difference between mean postoperative ICIQ score when there was high and low urethral mobility (mean 3.7 and 2.8, respectively) (p= 0.16).

when compared to those with hypermobility and low VLPP (table I). The former also presented lower rates of objective (p= 0.01)

In the subgroup analysis, patients with concomitant low mobility and high VLPP had worse objective success rates

and subjective (p= 0.04) success in relation to the remaining of the study population. Table I. Comparison between specific subgroups.							
Subgroups	Objective success – n(%)		р*	Subjective success – n(%)		р*	
	Yes	No		Yes	No		
Mobility < 30° and VLPP>60 cmH ₂ 0	8 (72.7)	3 (27.3)		6 (54.5)	5 (45.5)	0.00	
Mobility ≥30° and VLPP≤ 60 cmH ₂ 0	18 (100)	0 (0)	0.04	13 (72.2)	5 (27.8)	0.28	
TOTAL	26 (89.7)	3 (10.3)		19 (65.5)	10 (34.5)		

Interpretation of results

To more adequately analyze the interaction between urethral mobility and sphincter intrinsic dysfunction (indicated by VLPP \leq 60 cmH₂0), we compared subgroups and demonstrated that high urethral mobility predicts cure, even when VLPP indicates a theoretically more dysfunctional sphincter. Low mobility was a predictor of failure even in women without intrinsic sphincter deficiency (high VLPP), although cure rates remain acceptable. This type of analysis seeks to circumvent selection bias. A contemporary Turkish study had a similar conclusion ⁽³⁾.

Theoretically, a successful sling procedure restores continence not by increasing resting urethral pressure but by providing a support the holds the mid-urethra in place while the proximal urethra descends under stress, allowing better pressure transmission and, more importantly, a kinking of urethra during straining. When urethra doesn't move well, this kinking does not occur. That's the advocated mechanism for urethral mobility as a prognostic factor and not a cause of incontinence.

We propose a graphic to illustrate how VLPP and urethral mobility interact to determine prognosis following sling procedure (figure 1).

Figure 1. Prognostic interaction between urethral mobility and sphincter dysfunction in women with SUI after sling procedure. Note that some degree of sphincter dysfunction is required for incontinence to occur.



Concluding message

Urethral mobility is an important independent prognostic factor for TOT surgery. Low urethral mobility predicts higher failure rates, but it does not preclude surgery as most of these patients are cured following the procedure. No association was found between postoperative outcomes and VLPP.

References

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Specify source of funding or grant	None				
Is this a clinical trial?	Yes				
Is this study registered in a public clinical trials registry?	Yes				
Specify Name of Public Registry, Registration Number	This study was approved by Insitutional Review Board (Ethics				
	Committee) and registered in the national (Brazilian) research				
	database called SISNEP (www.saude.gov.br/sisnep) by the				
	number (CAAE): 0001.0.148.000-10. This database has public				
	access.				
Is this a Randomised Controlled Trial (RCT)?	No				
What were the subjects in the study?	HUMAN				
Was this study approved by an ethics committee?	Yes				
Specify Name of Ethics Committee	Ethics Commitee of Santa Casa de São Paulo Medical School.				
Was the Declaration of Helsinki followed?	Yes				
Was informed consent obtained from the patients?	Yes				