Assessment of Correction of Female Stress Urinary Incontinence with Trans Obturator Tape; Traditional versus Tailored

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Abstract

Background: Female genuine stress urinary incontinence (SUI) affects a great number of females, with important social and economic consequences. SUI is more frequent after several deliveries, after menopause as a result of changes in the urethral support or the sphincter mechanism.

Aim of Study: To compare traditional tension free trans-obturator tape (TOT) and surgeon tailored mesh (STM) in treatment of female stress urinary incontinence as regard of safety, efficacy, side effects, cost, and follow-up.

Patients and Method: Prospective randomized study was done on sixty three female patients were suffered SUI with determining of eligibility of the study as shown in COSORT flow chart Fig. (1) at Al Zahraa University Hospital, Al Azhar University from December 2016 to April 2019. All patients subjected to full history, general, and local examination (cases of cystocele, rectocele, or previous vaginal surgeries were excluded). Cystometrogram (CMG) was done for exclusion of uninhibited detrusor contractions, small bladder capacity (<250 CC), impaired bladder sensation, or impaired detrusor compliance. Patients with Significant Post voiding residual urine (>100 CC) assessed by ultrasonography were excluded from this study. All patients requested surgical treatment and provided informed written consent, and the study was approved by the ethical committee. With follow-up for six month post-operatively.

Results: Depending on subjective criteria obtained from the Patric questionnaire, and objective criteria from physical examination. As regards traditional tape group, 13 patients (43.33%) were cured; two patients (6.67%) improved with no failed cases and STM group 14 patients (46.67%) were cured; one patient (3.33 %) improved with no failed cases (p value 0.542, Pearson Chi-Square test).

Conclusions: TOT (outside in technique) using both STM and traditional tape are minimally invasive techniques and are equally effective and safe in treatment of pure female SUI, with short hospital stay which reduces the cost of surgery. Both techniques could be used in treatment of pure SUI with good outcome. This technique is more economical due to the cheap ordinary polypropylene mesh. So, it should be considered as a low-cost alternative to the available commercial kits in the treatment of female SUI, mainly for public health systems with few financial resources.

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Key Words: Female stress urinary incontinence – Surgeon-tailored mesh – Polypropylene mesh – TVT-O – TOT – Cost.

Introduction

STRESS urinary incontinence (SUI) is the commonest subtype of urinary incontinence (UI) of female patients affecting their quality of life (QoL) and that of their families. Accounting for about 50% of the patients. Which is prevalent problem of all age, median prevalence of female UI is 27.6% [1].

The bothersome symptom of SUI may adversely affect social relationships and activities, restrict physical pursuits, impair personal hygiene and lead to avoidance of sexual relationship [2].

The AUA and Society of Urodynamics, Female Pelvic Medicine and Urogenital Reconstruction guidelines on the management of SUI recommend four types of surgical treatment: Bulking agents; midurethral sling; autologous fascia pubovaginal sling; and Burch colposuspension [3].

Midurethral sling method, standard midurethral slings (SMUSs) and single-incision mini-slings (SIMSs) were recommended for the surgical treatment of SUI in women. SMUSs have shown a high success rate [4].

Patients and Methods

Prospective randomized study was done on sixty three female patients were suffered SUI with determining of eligibility of the study using COSORT flow chart (Fig. 1). All operations were carried out at Al Zahraa University Hospital, Al Azhar University from December 2016 to April 2019. All patients subjected to full history, general, and local examination (cases of cystocele, rectocele, or previous vaginal surgeries were excluded). Cystometrogram (CMG) was done for exclusion...
of uninhibited detrusor contractions, small bladder capacity (<250 CC), impaired bladder sensation, or impaired detrusor compliance. Patients with Significant Post voiding residual urine (>100 CC) assessed by ultrasonography were excluded from this study. All patients requested surgical treatment and provided informed written consent, and the study was approved by the ethical committee. Follow-up for six month post-operatively. The peri-operative data of both groups were collected, summarized, and compared using chi-square, fisher exact, or Mann-Whitney tests.

All operations were done by the same surgeon and two assistant. The patient was placed in the lithotomy position, with the thighs flexed at an angle of 120. A 16 French bladder catheter was inserted and the balloon was inflated in the bladder. The labia are retracted laterally by fixation to the skin with 3/0 silk stay sutures in order to expose the vulvar vestibulum. The anterior vaginal wall is suspended with two Allis clamps on either side of the midline, 1 cm proximal to the urethral meatus. Submucosal saline injection to elevate the vaginal mucosa (hydrodissection) easing the dissection.

A median sagittal incision of the vaginal wall is started 1cm proximal to the Urethral meatus and is continued proximally over a 1 cm distance. Both vaginal mucosal and sub-mucosal tissues are incised and dissected off the underlying periurethral fascia, lateral dissection with Metzenbaum scissors toward the inferior pubic ramus bilaterally was created. The internal edge of obturator foramen was identified with the tip of the index finger. Skin incision was made in or just medial to the genito-femoral fold at the base of the adductor longus tendon, approximately at the level of the clitoris. The needle was passed from the obturator incision to the vaginal incision by supination movement until palpated with the finger placed in the vaginal incision which guided it to outside through the vaginal incision.

In tailored method Polypropylene strips of about 30cm length and 1 cm width are handly prepared by dividing the commercially available 30x30 prolene mesh (the same mesh used in hernial repair). A prolene mesh of 30 x 30cm will give about 30 tapes. Each strip can be packaged and resterilized by plasma machine.

The tape was fixed to the needle and drawn out through the thigh by pronation movement. Tension of the tape over the urethra or its twisting was avoided. Long Mayo scissors was placed between the sling and the urethra to provide a space and to avoid any tension. The tension was tested when the bladder was filled with 250ml of sterile fluid; the patient was asked to cough and continence assessed. Cystoscopy was performed when needed (suspected bladder or urethral injury).

The ends of the sling cut below the surface of the skin. The skin was closed with 3-0 vicryl in a sub-cuticular fashion. Finally, the anterior vaginal wall incision was closed with 3-0 vicryl in interrupted fashion. A vaginal pack soaked with povidone-iodine was placed in the vagina. The urethral catheter was re-placed into the bladder and connected to closed-bag drainage.

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**Results**

In this study we compared two groups of patients were suffered genuine stress urinary incontinence and underwent mid urethral sling Trans obturator tape (fifteen patients had traditional TOT as control group and fifteen had surgeon tailored tape as study group) to evaluate the efficacy and safety of STM in comparing to traditional TOT in relieving SUI.

The data was collected, summarized, and then statistically analyzed by using IBM SPSS v19 program.

The mean age of STM group was $41.60 \pm 6.957$ years (range 30-56 years old). Seven patients aged from 31 to 40 years, six patients aged from 41 to
50 years, and two patients aged from 51 to 56 years old. Whereas in the traditional tape group the mean age was 45.67 ± 8.482 years (range 30-57 years old). Three patients aged from 31 to 40 years, six patients aged from 41 to 50 years, and six patients aged from 51 to 56 years old. \( p \)-value, student’s \( t \)-test was 0.460 (statistically insignificant).

Table (1): Age categories of both groups.

<table>
<thead>
<tr>
<th>Age category</th>
<th>Surgeon tailored mesh</th>
<th>Traditional tape</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>30-39</td>
<td>6</td>
<td>3</td>
<td>9</td>
</tr>
<tr>
<td>40-49</td>
<td>7</td>
<td>6</td>
<td>13</td>
</tr>
<tr>
<td>50-57</td>
<td>2</td>
<td>6</td>
<td>8</td>
</tr>
<tr>
<td>Total</td>
<td>15</td>
<td>15</td>
<td>30</td>
</tr>
</tbody>
</table>

\( p \)-value 0.310 (Eta test).

None of the patients were nulliparous. The mean number of deliveries in STM group was 4.2 ± 1.56 (range 2-8 deliveries). The mean vaginal deliveries in the 30 patients were 4.1 with a range of 2-8 deliveries. Two patients only have 1 cesarean section in addition to their vaginal deliveries. The range of cesarean section (CS) in the 30 patients was 0 to 1.

Table (2): Distribution of parity in two group of study.

<table>
<thead>
<tr>
<th>Parity</th>
<th>Surgeon tailored mesh</th>
<th>Traditional tape</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>p2+0</td>
<td>5</td>
<td>1</td>
<td>6</td>
</tr>
<tr>
<td>p2+1</td>
<td>0</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>p2+2</td>
<td>1</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>p3+0</td>
<td>2</td>
<td>3</td>
<td>5</td>
</tr>
<tr>
<td>p3+1</td>
<td>3</td>
<td>2</td>
<td>5</td>
</tr>
<tr>
<td>p3+2</td>
<td>0</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>p4+0</td>
<td>0</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>p5+0</td>
<td>2</td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>p5+1</td>
<td>1</td>
<td>2</td>
<td>3</td>
</tr>
<tr>
<td>p7+1</td>
<td>1</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Total</td>
<td>15</td>
<td>15</td>
<td>30</td>
</tr>
</tbody>
</table>

\( p \)-value 0.156 (Eta test).

Seven (40%) out of fifteen cases of study group (STM) were had ALPP ranging 101-150 ML, while eight (53.33%) out of fifteen of control group (traditional tape) with ALPP 51-100 ML (\( p \)-value 0.150, Eta test).

Table (3): Abdominal leak point pressure (ALPP).

<table>
<thead>
<tr>
<th>Group</th>
<th>Surgeon tailored mesh</th>
<th>Traditional tape</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;50cc</td>
<td>1</td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td>51-100cc</td>
<td>6</td>
<td>8</td>
<td>14</td>
</tr>
<tr>
<td>101-150</td>
<td>7</td>
<td>6</td>
<td>13</td>
</tr>
<tr>
<td>151-200</td>
<td>1</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Total</td>
<td>15</td>
<td>15</td>
<td>30</td>
</tr>
</tbody>
</table>

\( p \)-value 0.150 (Eta test).

Table (4): Mean ± SD of operative time of both groups.

<table>
<thead>
<tr>
<th>Group</th>
<th>Number of cases</th>
<th>Mean</th>
<th>Std. Deviation</th>
<th>Std. Error Mean</th>
</tr>
</thead>
<tbody>
<tr>
<td>STM</td>
<td>15</td>
<td>20.47</td>
<td>3.603</td>
<td>0.930</td>
</tr>
<tr>
<td>Traditional tape</td>
<td>15</td>
<td>19.47</td>
<td>3.226</td>
<td>0.833</td>
</tr>
</tbody>
</table>

\( p \)-value 0.429 (student’s \( t \)-test).

Table (5): Operative time of both groups.

<table>
<thead>
<tr>
<th>Group</th>
<th>Surgeon Tailored Mesh</th>
<th>Traditional Tape</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>12-18 min</td>
<td>6</td>
<td>7</td>
<td>13</td>
</tr>
<tr>
<td>19-25 min</td>
<td>9</td>
<td>8</td>
<td>17</td>
</tr>
<tr>
<td>Total</td>
<td>15</td>
<td>15</td>
<td>30</td>
</tr>
</tbody>
</table>

\( p \)-value 0.50 (Pearson Chi-Square test).

Intra-operative complications:

In both groups there are minimal intra Operative bleeding with no need for use of diathermy coagulation. And one case expected bladder injury which need to cystoscopy and exclusion of perforation. There is no urethral injury of both groups.

Post-operative complications:

There are no post operative cases with early complications as obturator hematoma (dissecting hematoma), urine retention, and high post voiding residual urine estimated by ultrasound, and De novo urgency; or late complications as vaginal erosions, vaginal discharge (infection), and wound infection.

Transient thigh pain occurs in 2 cases (6.7%) of STM and 3 cases (10%) of traditional tape group which respond well to NSAIDs and completely resolved within one month after operation.
Post-operative follow-up:

Post-operative follow-up carried out at 1 week, 1 month and 3 months postoperatively. At each visit all patients were subjected to Clinical assessment in the form of any distressing problem. She was asked the same questions in the pre-operative questionnaire. She was asked specifically about stress incontinence, frequency of micturition, urgency, or urge incontinence (whether persisted or developed postoperatively), voided volume, any vaginal spotting, or bleeding especially in the early postoperative period. She was asked about any vaginal pain, dyspareunia, vaginal discharge, suprapubic pain or thigh pain. She was asked about the quality of her micturition regarding strength, intermittency, deviation of stream and post voiding dribbling. Examination for the presence of discharge, vaginal erosion, chronic retention, and cough stress test.

Outcome of the procedures was evaluated depending on subjective criteria obtained from the Patric questionnaire, and objective criteria from physical examination.

As regards traditional tape group, 13 patients (43.33%) were cured; two patients (6.67%) improved with no failed cases and STM group 14 patients (46.67%) were cured; one patient (3.33%) improved with no failed cases (p-value 0.542, Pearson Chi-Square test).

Discussion

Our two groups showed no statistically significant difference concerning age, number of deliveries, number of postmenopausal women, type, and grade of SUI. There was no statistically significant difference between the two groups in the mean abdominal leak point pressure or preoperative mean Q max.

There was no statistically significant difference in the operative time among the two groups (20.47±3.603min; 19.47±3.226min. for STM and traditional tape respectively) (p-value 0.429, student’s t-test).

Compared with study by Ogah et al. [5]. There is a significant statistically difference regarding the operative time when comparing our technique with the original TVT-O technique (in-out technique). Mean operative time of the original TVT-O was 14min (range: 11-20) in case of isolated SUI treatment [6].

Oliveira et al. [5] reported that no bladder or urethral injury in the original TVT-O technique and thus there is no need for routine cystoscopy as in our study there was no cases with urethral or bladder injury [5].

Intraoperative cystoscopy not done except in one (3.33%) patient of the TOT by traditional tape group with suspect bladder injury during the procedure and reveal no bladder injury. This low incidence of Intraoperative urethral or bladder injury may be due to insertion of urethral catheter and evacuation of the bladder before dissection so avoid urethral and bladder injuries. Minimal dissection, and tape should be inserted at mid urethra without tension (tension free) to avoid positioning of tape with tension place blunt instrument (forceps or scissor) between urethra and the tape during adjustment of tape.

In our study there was only one case with lateral vaginal wall injury during dissection in the early cases of study of traditional tape group (which may be due to progression in learning curve of the procedure for surgeon) which repaired immediately by vicryl 3/0; compared to Debodinance, [5] reported that perforation of the lateral wall of the vagina occurred in one case from 50 patients [7].

Lukacz and Co-workers, [5] in study included 103 patients evaluated the effect of TVT procedure on post-voiding residual urine and found that there was no clinically significant increase in post-voiding residual urine in comparison of our study we consider that post void residual urine less than 50cc estimated by ultrasound is insignificant there were no perioperative statistically significant difference of both group [8].

Procedure outcome differs in the literatures One of the papers used a cough stress test and a

As regards traditional tape group, 13 patients (43.33%) were cured; two patients (6.67%) improved with no failed cases and STM group 14 patients (46.67%) were cured; one patient (3.33%) improved with no failed cases. These results were comparable to earlier studies summarized in table 13 with cure rate ranging from 83-95%.

Delorme and colleagues [12] in study included 32 patients revealed a cure rate of 90.6%, with a further 9.4% improvement with no failure using trans-obturator tape.

In study done by deTayrac and colleagues [12]. About The outcome after TOT and TVT procedures was evaluated Their study included 30 TOT procedures. The cure rate in their study for TOT cases was 90%, with a further 3.3% improvement, and a failure rate of 6.7%. The mean follow-up was 12 months.

Cindolo et al., [12] reported that 96% of cases satisfied by result of surgery in their study which was conducted on 80 women complaining of SUI. The mean follow-up was 4 months [9].

Evaluation of outcome of TOT procedure in 94 patients with SUI One year follow-up. By Mellier and colleagues, [12] revealed 95% cure rate, with a further 4% improvement, and a failure rate of 1%. The mean follow-up was 12 months ranging from 2-20 months.

Thom and Klutke, [12]. In study on the outcome after TOT procedure included 47 women complaining of SUI. The cure rate in their study was 96%. The mean follow-up was 18 months.

Summary and Conclusion:

TOT (outside in technique) using both STM and traditional tape are minimally invasive techniques and are equally effective and safe in treatment of pure female SUI, with short hospital stay which reduces the cost of surgery. Both techniques could be used in treatment of pure SUI with good outcome.

These results are also comparable with the results of TOT that were reported in the previous literature.

Furthermore, this technique is more economical due to the cheap ordinary polypropylene mesh.

So, it should be considered as a low-cost alternative to the available commercial kits in the treatment of female SUI, mainly for public health systems with few financial resources.

Randomized controlled studies with a longer duration of follow-up are needed to confirm our results.

Conflict of interest: None.

Funding source: None.

References


تقييم إصلاح السلس البولي الإجهاد للإناث

بطاقة الشريط الراقي لمنع المثانة المعتد
مقارنة بالتصميم يدوياً

السلس البولي هو من أكثر المشكلات الصحية شيوعاً وخاصة في فترة النساء اللواتي مرن بتجربة الولادة مراراً أو تعرضن لولادة عسرة.

واللأثير كبرى في المجال الاجتماعي.

ويعرف سلس البول بأنه عدم القدرة على التحكم في البول مما يؤدي إلى تسررب البول إرادياً والذي يمثل مشكلة صحية واجتماعية.

وسلس البول أنواع متعددة أكثرها شيوعاً:

1- سلس البول الإجهاد: وهو فقدان القدرة على التحكم في البول إرادياً مع أي مجهود مفاجئ مثل الضحك والسعال والعطس.

2- سلس البول الإجهاد: وهو فقدان التحكم في البول إرادياً بعد إ焓س المفاجئ بالحاجة إلى البول.

3- سلس البول المختلط: وهو مزيج من سلس البول الإجهاد والسلس البولي الإجهاد.

4- سلس البول الفائق: يحدث عندما تتعذر المثانة على التفرز الكامل مما يسبب في تسررب البول الفائض بشكل مستمر وهذا النوع من سلس البول نادرًا ما يصيب السيدات.

5- سلس البول الوظيفي: هذا ينتج عن عوامل خارجية مثل الوقوف الطويل ويتطلب هذا النوع في الشياطين المتقدمة.

طرق علاج السلس البولي الإجهاد تتضمن:

- تدريب ضعلات البول (العلاج الطبيعي لعضلات البول)، العلاج بالدورة، الشريط الراقي لمنع المثانة، الجراحة، وحديثاً العلاج بالليزر.

قد تمت هذه الدراسة على ثلاثين مريضة من مجموعتين من سلس البول الإجهاد في الفترة من فبراير 2017 إلى فبراير 2019 وتم تقسيم المرضى عشوائياً إلى مجموعتين تضم كل مجموعة 15 مريضة.

- مجموعة (A): وتم علاج المرضى بواسطة الشريط الراقي لمنع المثانة المعتد.

- مجموعة (B): وتم علاج المرضى بواسطة الشريط الراقي لمنع المثانة المعتد.

وقد تمت ملخص المرضى لأخذ التاريخ المرضي والإسهام الإكلينيكي، وتم فحص الحوادث اللازمة لهم وتشمل:

- التحاليل المجهزة: تحيل بول، صورة دم كاملة، وظائف يكي وكبد، وقياس معدلات النزيف.

- الإضافات: العمليات.

- فحص دراسة البول.

وتم تقسيم جميع المرضى قبل وأثناء وبعد العملية من أخذ الملاحظات وعمل الجداول اللازمة لوصول إلى النتائج المطلوبة.

نتائج: وجدنا أن متوسط زمن إجراء العملية متقارب في المجموعتين ولم يحدث نزيف يطلب نقل مرضى الجراحة لأسعى المريضة من المجموعة. وحول أثر الجراحة تجارب الجراح الخارجي لحالة من مجموعة (A) وتم إصلاحها أثناء العملية ولم تترك أي آثار جانبية بعد ذلك.

أثبتت الدراسة أن كلا الطرقتين فعالة في علاج حالات السلس البولي الإجهادي من حيث النجاح والتحسين والشفاء.

وقد أثبت البحث تقارب نسب النجاح ونسبة المضاعفات لكل الطرقتين من النسبة المطلوبة عالمياً.

نستنتج مما سبق أن هذه الطرقية في استخدام الشريط الراقي التي يتم تصميمها يدويًا بواسطة الجراح تعتبر طريقة ناجحة وآمنة وأقل تكلفة في علاج حالات السلس البولي الإجهادي لدى السيدات وفي المرضى الذين من النتائج الجيدة التي أثبتتها هذه الطرقية العلاجية إلا أنه لا تزال الحاجة إلى إجراء المزيد من الأبحاث خلال فترة زمنية أطول وعدد أكبر من المرضى.