382

Tsai C1, Hung M1

1. Department of Obstetrics and Gynecology, Taichung Veterans General Hospital, Taichung, Taiwan

A COMPARISON OF ONE-YEAR OUTCOMES OF TWO SINGLE INCISION VAGINAL MESH PROCEDURES FOR PROLAPSE REPAIR

Hypothesis / aims of study

Use of vaginal meshes for treatment of pelvic organ prolapse (POP) remains controversial. In recent years, there is a trend toward reduce the volume/weight of meshes and single incision surgery. Two methods of single incision mesh involved either lateral/apical anchoring (ElevateTM, lightweight mesh) or non-anchoring (ProsimaTM) had been launched. The aim of this study was to assess safety and efficacy after the above two procedures.

Study design, materials and methods

A cohort study was conducted at a tertiary referral center between Aug 2010 and Feb 2014. Patients who underwent transvaginal pelvic reconstruction surgery using single incision vaginal mesh were recruited. A detailed comparison of one-year outcomes, including anatomic outcome and surgical complications was made. Data were analysed with univariate methods or multivariate logistic regression analysis accordingly.

Results

One hundred and forty-two patients (51 in ElevateTM group and 91 in ProsimaTM group) were recruited and analyzed. Demographic data and surgical result were presented in **Table 1.** Most of these parameters were similar between groups. Mesh erosion rate was lower but not statistically significant in ElevateTM group (5.9% vs. 12.8% in ElevateTM and ProsimaTM group, respectively, P=0.379). Other complications were rare and comparable between groups. Objective anatomic success (POP stage \leq 1) rate was higher in ElevateTM group at one year follow-up, although not statistically significant (86% and 78.2% after ElevateTM and ProsimaTM repair, respectively, P=0.264). Value of post-operation Ba point was significantly greater in ProsimaTM repair group (-2.59cm vs -1.18cm in ElevateTM and ProsimaTM group, respectively, p=0.004). Subgroup analysis about anatomical success/failure was made in **Table 2**. Those in the anatomical failure group had a greater value of pre-operative a point (3.38cm in failure group while 2.26cm in success group, p=0.032).

Interpretation of results

The Elevate™ prolapse repair system had a better 1-year anatomical cure rate of the anterior compartment than Prosima™, which might reflect mesh anchoring prevents anterior vaginal wall descent. Operative and postoperative experiences were similar between groups; however, lightweight mesh seems to have less mesh exposure according to this study.

Concluding message

Single-incision vaginal mesh surgery for treatment of POP seems to be safe and efficient, but have limitations when used to manage severe anterior vaginal wall prolapse. Non-anchoring vaginal mesh may be less powerful for anterior compartment support. Besides, lightweight mesh is important when considering reduce mesh erosion rate.

Table 1. Preoperative characteristics and surgical results of patients who underwent pelvic reconstructive surgery using

Prosima(N=91) or Elevate(N=51).

,		Elevate (N=51)	
Range	Value	Range	P value
(38-78)	65.1 ± 7.1	(49~79)	0.001*
(20.0-32.0)	24.8 ± 3.3	(16.9~33.1)	0.334*
(4-9)	5.4 ± 2.2	(4~17)	0.477*
(3-8)	3.9 ± 2.3	(3~11)	0.342*
9 (90-190)	166.2 ± 28.0	(120~210)	<0.001*
.5 (50-800)	141.2 ± 80.1	(50~500)	0.347*
(30/75)	47.5%	(19/40)	0.581#
(9/62)	5.9%	(3/51)	0.537#
(6/85)	13.7%	(7/51)	0.224#
(11/91)	5.9%	(3/51)	0.379#
(68/87)	86%	(43/50)	0.264#
05	-2.59 ± 0.93		0.044*
09	-6.65 ± 2.08		0.180*
34	-3 ± 0		0.290*
	Range (38-78) (20.0-32.0) (4-9) (3-8) (90-190) .5 (50-800) (30/75) (9/62) (6/85) (11/91) (68/87)	Range Value (38-78) 65.1 ± 7.1 (20.0-32.0) 24.8 ± 3.3 (4-9) 5.4 ± 2.2 (3-8) 3.9 ± 2.3 (90-190) 166.2 ± 28.0 (50-800) 141.2 ± 80.1 (30/75) 47.5% (9/62) 5.9% (6/85) 13.7% (11/91) 5.9% (68/87) 86% 05 -2.59 ± 0.93 09 -6.65 ± 2.08	Range Value Range $(38-78) 65.1 \pm 7.1 (49-79)$ $(20.0-32.0) 24.8 \pm 3.3 (16.9-33.1)$ $(4-9) 5.4 \pm 2.2 (4-17)$ $(3-8) 3.9 \pm 2.3 (3-11)$ $(90-190) 166.2 \pm 28.0 (120-210)$ $.5 (50-800) 141.2 \pm 80.1 (50-500)$ $(30/75) 47.5\% (19/40)$ $(9/62) 5.9\% (3/51)$ $(6/85) 13.7\% (7/51)$ $(11/91) 5.9\% (3/51)$ $(68/87) 86\% (43/50)$ -2.59 ± 0.93 -6.65 ± 2.08

Table 2. Patient characteristics about surgical failure (post-op POP stage ≥ 2) or success using single incision vaginal mesh at 1 year follow-up.

Patient characteristics	Failure group(N=26)		Success group (N=116)		
	Value	Range	Value	Rage	P value
Mean age (year)	62.9 ± 9.8	(48-78)	62.0 ± 7.7	(38~79)	0.924*
Median parity	3	(1-7)	3	(1~6)	0.589*
Mean body mass index (kg/m²)	25.1 ± 3.0	(18.9-31.5)	25.2 ± 3.2	(16.9~33.1)	0.722*
% Prior hysterectomy	15.4%	(4/26)	16.4%	(19/116)	1.000#
% Prior prolapse repair	23.1%	(6/26)	7.8%	(9/116)	0.078#
% Concomitant hysterectomy	40%	(6)	47.5%	(42)	0.180#
% De novo stress incontinence	7.7%	(2/)	8.6%	(10/)	1.000#
% De novo urgency incontinence	3.8%	(1/)	10.3%	(12/)	0.264#
% Vaginal mesh extrusion	11.5%	(3/26)	9.5%	(11/116)	1.000#
Pre-op POPQ parameters					
Ba (cm)	3.4 ± 1.9	(0~+7)	2.6 ± 1.9	(-2~+8)	0.032*
C (cm)	0.9 ± 3.7	(-4~+8)	1.0 ± 2.9	(-6~+8)	0.832*
Bp (cm)	-0.1 ± 2.2	(-3~+7)	0.0 ± 2.1	(-3~+8)	0.617*
Bp (cm)	-0.1 ± 2.2	(-3~+7)	0.0 ± 2.1	(-3~+8)	0.617

^{*:} Mann-Whitney test; #: Fisher's exact test.

Disclosures

Funding: Non. Clinical Trial: Yes Public Registry: No RCT: No Subjects: HUMAN Ethics Committee: CE11280 at Taichung Veterans General Hospital, Taiwan. Helsinki: Yes Informed Consent: Yes