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TVT-SECUR: SECURE OR INSECURE?

Hypothesis / aims of study

The purpose of this study was to assess the feasibility of using 3D/4D ultrasound to visualise mid-urethral tapes after insertion⁽¹⁾. The surgical management of female stress urinary incontinence (SUI) has been revolutionized since the introduction of the tension free vaginal tape in1995. The tension-free vaginal tape (TVT) has been a highly efficacious surgical treatment for female stress urinary incontinence with the longest follow up of 7.6 years, demonstrating cure rate of 81% and improvement in 16%.

With the growing popularity of this procedure, many other versions of the mid-urethral tapes were introduced with minor variations. Another minor leap in the progress is tunneling the tape through the obturator foramen instead of the retropubic region, for example, Tension free vaginal tape-Obturator (TVT-O) (2). This has the added advantage of avoiding entry into the pelvis, hence reducing the likelihood of injury to the bladder, intraperitoneal viscera and major vessels. Medium term cure rates are similar to that of TVT.

Recently a newer and shorter tape (Tension free vaginal tape-Secur (TVT-S)) was introduced.

We performed 3D/4D ultrasound on 5 patients who had TVT-S inserted for SUI (3).

Study design, materials and methods

3D/4D U/S was performed using Voluson 730 Expert on 5 patients who had TVT-S inserted more than 6 months ago. The patient's bladder was filled to 300mls with normal saline for the U/S.

Results

We were able to see the placement of TVT-S easily. The position of the tape yielded important reasons that could substantiate the failure of mid-urethral sling surgeries.

Interpretation of results

Some of the tapes were placed with unequal arms. Some tapes were twisted.

Concluding message

Trans-perineal 3D/4D ultrasound provides a unique and revolutionary technique to visualize the mid-urethral tapes after insertion.

References

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Specify source of funding or grant	Yes. KK Research Grant
Is this a clinical trial?	Yes
Is this study registered in a public clinical trials registry?	Yes
Specify Name of Public Registry, Registration Number	Centralised Institutional Review Board, EC200703042
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	Centralised Institutional Review Board
Was the Declaration of Helsinki followed?	Yes
Was informed consent obtained from the patients?	Yes