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LONG TERM OUTCOMES OF TENSION-FREE VAGINAL TAPE IN THE TREATMENT OF STRESS INCONTINENCE IN FEMALES WITH NEUROPATHIC BLADDERS

Hypothesis / aims of study

To evaluate the long-term outcomes of tension-free vaginal tape (TVT) for the treatment of urinary stress incontinence in females with neuropathic bladders secondary to spinal cord pathology

Study design, materials and methods

A retrospective analysis identified 11 female patients with spinal cord pathology, treated with TVT for stress urinary incontinence, between November 1997 and December 2000. All the procedures were performed by a single specialist surgeon. All patients had pre-operative evaluation including video-cystometrogram (VCMG). Long-term outcome assessment of continence included evidence of urine leakage on VCMG and/or reporting of leakage necessitating pad-usage. Complete cure was defined as no leakage on VCMG and no use of pads. Partial cure was defined as evidence of urine leak on VCMG with >50% reduction in the number of pads used.

Results

Of the 11 patients, 4 were lost to follow-up. Spinal cord pathology in these women included: (i) traumatic spinal cord injuries (n=3); (ii) post lumbar spinal surgery (n=3); (iii) spinal stenosis (n=1). The mean age was 58yrs (range 48 to 76yrs). The mean duration of follow-up was 7.9yrs (range 7.5 to 9.3yrs). Four patients (57%) were completely dry with no usage of pads reported and verified with no demonstrable urine leak on VCMG. Two patients (29%) achieved partial cure. Only 1 patient (14%) reported worsening symptoms and an increase in pad usage at 9.3 years. She had been partially cured at 5 year follow-up.

Interpretation of results

More than half the patients achieved complete cure with maintenance of continence following TVT insertion in this group of patients. This was observed at a minimum of 7.5 years (mean 7.9 years) following the procedure.

Concluding message

In females with spinal cord pathology who have stress urinary incontinence necessitating a definitive intervention, insertion of TVT should be considered a desirable treatment modality with durable outcomes.

Specify source of funding or grant	nil
Is this a clinical trial?	No
What were the subjects in the study?	HUMAN
Was this study approved by an ethics committee?	No
This study did not require eithics committee approval because	retrospective analysis of approved treatment
Was the Declaration of Helsinki followed?	Yes
Was informed consent obtained from the patients?	Yes