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## **FIVE-YEAR OUTCOMES OF THE IRIS PROCEDURE FOR TREATMENT OF FEMALE STRESS URINARY INCONTINENCE: COMPARISON WITH TVT PROCEDURE**

### Hypothesis / aims of study

We evaluated the long-term efficacy and safety of the IRIS (innovative replacement of incontinence surgery, B. Braun Korea & Dow Medics Co., Ltd, Seoul) procedure and compared it with TVT (tension free vaginal tape) for the treatment of female stress urinary incontinence.

### Study design, materials and methods

We included all 111 consecutive women who underwent IRIS (n=51) or TVT (n=60) between January 2002 and December 2003 and followed them up for at least 5 years postoperatively. We analyzed the 5-year success rate and postoperative complications in the IRIS procedure and compared to the results of TVT procedure.

### Results

The 5-year success rate was 92.2% for the IRIS and 93.3% for the TVT, and the satisfaction rates were 90.2% and 85.0%, respectively. Intraoperative complications for the IRIS group included 4 cases of bladder perforation, and there were 5 cases of bladder perforation for the TVT group. The postoperative complications for the IRIS group included 6 patients with *de novo* urgency, 2 patients with mesh exposure and one patient showed temporary urinary retention. Nine patients with the TVT developed *de novo* urgency, 2 patients with mesh exposure and 2 patients showed temporary urinary retention.

### Interpretation of results

On the basis of our results, the IRIS may be an effective and safe procedure as compared to the TVT for more than 5 years.

### Concluding message

### References

1. Yoon CJ, Jung HC. Two-year follow up of IRIS procedure for surgical treatment of female stress urinary incontinence. J Korean Continence Soc 2006;10:44-8
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<b><i>Is this a clinical trial?</i></b>	<b>Yes</b>
<b><i>Is this study registered in a public clinical trials registry?</i></b>	<b>Yes</b>
<b><i>Specify Name of Public Registry, Registration Number</i></b>	<b>Yeungnam University</b>
<b><i>What were the subjects in the study?</i></b>	<b>HUMAN</b>
<b><i>Was this study approved by an ethics committee?</i></b>	<b>Yes</b>
<b><i>Specify Name of Ethics Committee</i></b>	<b>Yeungnam University IRB center</b>
<b><i>Was the Declaration of Helsinki followed?</i></b>	<b>Yes</b>
<b><i>Was informed consent obtained from the patients?</i></b>	<b>Yes</b>