

USEFULNESS OF 1-HOUR PAD TEST AS PREOPERATIVE DIAGNOSTIC ASSESSMENT FOR STRESS URINARY INCONTINENCE IN KOREAN SOCIETY.

Hypothesis / aims of study

The aim of this study was to determine whether or not the 1-hour pad test demonstrates the objective severity of stress urinary incontinence.

Study design, materials and methods

Twenty nine female patients with stress urinary incontinence symptom were prospectively evaluated with a 1-hour pad test as recommended by the International Continent Society and they also underwent videourodynamics to determine the ALPP(abdominal leak point pressure). The patients were divided into 2 groups by the ALPP: group A (n=16) was the low leak point pressure group (ALPP≤100cmH₂O), and group B (n=13) was the high leak point pressure group (ALPP > 100cmH₂O or no leakage). A pad gain ≤2g was considered a negative pad test. Student's t-test was done to evaluate the difference of urine leakage between the two groups.

Results

The mean amount of urine leakage measured by the 1-hour pad test for groups A and B were 7.5g and 4.1g, respectively, and there was statistically significant differences between two groups (p=0.048). Six (20.7%) women did not leak during the ALPP measurement and 12 (41.4%) women had a negative pad test. Among the 6 women with no leakage on the ALPP, 3 had a positive pad test and 3 had a negative pad test. Among the 12 women with a negative pad test, 9 had leakage and 3 had no leakage during the ALPP measurement.

Interpretation of results

The 1-hour pad test was not more sensitive to demonstrate leakage than the ALPP. However, these data suggest that the 1-hour pad test did demonstrate the objective severity of stress urinary incontinence.

Concluding message

The 1-hour pad test is useful to assess the severity of urinary incontinence and the improvement of treatment.

References

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Is this a clinical trial?	Yes
Is this study registered in a public clinical trials registry?	No
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	SoonChunHyang University IRB
Was the Declaration of Helsinki followed?	Yes
Was informed consent obtained from the patients?	Yes