1 Guidance

1.1 Current evidence suggests that there are no major safety concerns associated with retrograde urethral sphincterometry. However, there is a lack of evidence on the diagnostic utility of this procedure (that is, the extent to which knowledge of its results improves patients' outcomes) and it should be performed only in the context of good-quality research.

2 The procedure

2.1 Indications

2.1.1 Stress urinary incontinence is the involuntary leakage of urine during exercise or certain movements such as coughing, sneezing and laughing. It is usually caused by weak or damaged muscles and connective tissues in the pelvic floor and urethral sphincter.
2.2 **Outline of the procedure**

2.2.1 Retrograde urethral sphincterometry (RUS) involves placing a cone-shaped device a short distance (about 5 mm) into the external urethral meatus. The device then infuses fluid at a controlled rate into the urethra. The pressure required to open the urethral sphincter is displayed on the device.

2.3 **Efficacy**

2.3.1 Preliminary data on the use of this procedure in women with stress urinary incontinence found that there was a weak relationship between the results of this test and other standard tests. In a trial of 258 symptomatic women the mean retrograde urethral pressure as measured by RUS was 71 cm H\textsubscript{2}O, and the mean values were reported as decreasing with increasing symptom severity. In another study, the mean retrograde urethral pressure was found to be 112.6 cm H\textsubscript{2}O in 61 asymptomatic women.

2.3.2 The impact of this procedure on patient outcomes is currently unclear. For more details, refer to the Sources of evidence.

2.3.3 The Specialist Advisors noted that efficacy outcomes are yet to be established.
2.4 Safety

2.4.1 In a study of 258 women, pain (2%) and dysuria (2%) were the two most commonly reported complaints. A total of 12 adverse events were noted in a study of 61 asymptomatic women who had RUS; these included lower back pain (2%), discomfort (2%), urethral pain (3%), dysuria (3%), urinary urgency (3%), urinary frequency (3%) and transient incontinence (3%). For more details, refer to the Sources of evidence.

2.4.2 The Specialist Advisors noted urinary tract infection and mild discomfort as potential adverse events.

3 Further information

Sources of evidence

The evidence considered by the interventional procedures advisory committee is described in the interventional procedure overview of retrograde urethral sphincterometry.

Information for patients

NICE has produced information for patients and carers on this procedure. It explains the nature of the procedure and the guidance issued by NICE, and has been written with patient consent in mind.

Update information

Minor changes since publication

January 2012: minor maintenance.

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Endorsing organisation

This guidance has been endorsed by Healthcare Improvement Scotland.