



Circular HSC (SQSD) (NICE IPG775) 21/23

**Subject: NICE Interventional Procedures Guideline
IPG775 - Biodegradable subacromial
spacer insertion for rotator cuff tears**

For action by:

SPPG – for distribution to:

Family Practitioner Services Leads – for cascade to relevant
Family Practitioner groups

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Director of Public Health and Medical Director – for cascade
to relevant staff
Director of Nursing and AHPs – for cascade to relevant
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Chief Executives of HSC Trusts – for distribution to:

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Chief Executive of Regulation & Quality Improvement Authority - for
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Chief Executive of NIBTS

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Chief Executive/Postgraduate Dean, NIMDTA

Clinical Education Centre, BSO

NI NICE Implementation Facilitator

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Summary of Contents:

This guidance provides evidence-based recommendations on
biodegradable subacromial spacer insertion for rotator cuff tears.
This involves inserting a balloon-shaped device between the top of
the shoulder blade and the upper arm bone to reduce pain and
improve shoulder function.

Enquiries:

Any enquiries about the content of this Circular should be addressed
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Related documents:

HSC (SQSD) 14/22

Superseded documents

None

Status of Contents:

For action

Implementation:

Immediate

Additional copies:

Available to download from

<https://www.health-ni.gov.uk/topics/safety-and-quality-standards/national-institute-health-and-care-excellence-nice>

Dear Colleague

**NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE (NICE):
INTERVENTIONAL PROCEDURES GUIDELINE IPG775 - BIODEGRADABLE
SUBACROMIAL SPACER INSERTION FOR ROTATOR CUFF TEARS**

The National Institute for Health and Care Excellence (NICE) produces guidance as part of the Interventional Procedure Programme. Clinicians who have developed new procedures send these to NICE to evaluate their effectiveness. Cost is not considered as part of this evaluation. Solely for this programme, NICE communicates directly with HSC organisations in Northern Ireland and consequently Departmental bi-monthly circulars are normally only required. An exception has been made for this due to the evidence identified on this procedure.

The advice in the NICE guideline states:

When debridement is a suitable option, biodegradable subacromial spacer insertion for rotator cuff tears should not be used.

HSC organisations also should note that the Department expects them to comply with this guidance in their delivery of services to patients when debridement is a suitable option.

Indications and current treatments

People who have rotator cuff tears may have shoulder pain and weakness, with reduced shoulder function, leading to a reduced quality of life. Rotator cuff tears can be caused by an injury or can develop gradually. They can be minor or severe depending on the degree of damage to the tendons. Minor tears to the rotator cuff are very common and may not cause any symptoms at all. Diagnosis is usually by ultrasound or MRI.

Conservative treatment may include physical therapy, pharmacological treatments (including pain relief, and topical or oral non-steroidal anti-inflammatory medicines) and corticosteroid injections. If the tear is severe or has not responded to other treatments, surgical interventions such as debridement, rotator cuff repair, subacromial smoothing, tendon transfer or shoulder arthroplasty may be needed.

Outline of the procedure

Inserting a biodegradable subacromial spacer aims to improve pain and restore shoulder function in people who have irreparable rotator cuff tears. The aim is to reduce subacromial friction by lowering the humeral head during shoulder abduction. It is a less invasive and potentially safer alternative to reverse shoulder arthroplasty or tendon transfer and has shorter procedure and rehabilitation times.

The procedure is done under general or regional anaesthesia. The subacromial space is visualised using either arthroscopy or mini-open surgery. The damaged area is surgically cleared. Measurements are taken to determine the size of biodegradable spacer needed. The balloon-like spacer is then inserted into the subacromial space and inflated with saline solution. Once a sufficient volume is reached, the balloon is sealed and left in place. The balloon spacer is made from a biodegradable polymer and resorbs over about 1 year.

The full NICE interventional procedure guideline is available for download at:
<https://www.nice.org.uk/guidance/ipg775>

A handwritten signature in black ink, appearing to read 'Michael McBride', written in a cursive style.

Prof Sir Michael McBride
Chief Medical Officer