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Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considered (Review)



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[Diagnostic Test Accuracy Review]

Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considered

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ABSTRACT

Background

Shoulder pain is a very common symptom. Disorders of the rotator cuff tendons due to wear or tear are among the most common causes of shoulder pain and disability. Magnetic resonance imaging (MRI), magnetic resonance arthrography (MRA) and ultrasound (US) are increasingly being used to assess the presence and size of rotator cuff tears to assist in planning surgical treatment. It is not known whether one imaging method is superior to any of the others.

Objectives

To compare the diagnostic test accuracy of MRI, MRA and US for detecting any rotator cuff tears (i.e. partial or full thickness) in people with suspected rotator cuff tears for whom surgery is being considered.

Search methods

We searched the Cochrane Register of Diagnostic Test Accuracy Studies, MEDLINE, EMBASE, and LILACS from inception to February 2011. We also searched trial registers, conference proceedings and reference lists of articles to identify additional studies. No language or publication restrictions were applied.

Selection criteria

We included all prospective diagnostic accuracy studies that assessed MRI, MRA or US against arthroscopy or open surgery as the reference standard, in people suspected of having a partial or full thickness rotator cuff tear. We excluded studies that selected a healthy control group, or participants who had been previously diagnosed with other specific causes of shoulder pain such as osteoarthritis or rheumatoid arthritis. Studies with an excessively long period (a year or longer) between the index and reference tests were also excluded.

Data collection and analysis

Two review authors independently extracted data on study characteristics and results of included studies, and performed quality assessment according to QUADAS criteria. Our unit of analysis was the shoulder. For each test, estimates of sensitivity and specificity



from each study were plotted in ROC space and forest plots were constructed for visual examination of variation in test accuracy. Metaanalyses were performed using the bivariate model to produce summary estimates of sensitivity and specificity. We were unable to formally investigate potential sources of heterogeneity because of the small number of studies.

Main results

We included 20 studies of people with suspected rotator cuff tears (1147 shoulders), of which six evaluated MRI and US (252 shoulders), or MRA and US (127 shoulders) in the same people. Many studies had design flaws, with the potential for bias, thus limiting the reliability of their findings. Overall, the methodological quality of the studies was judged to be low or unclear. For each test, we observed considerable heterogeneity in study results, especially between studies that evaluated US for the detection of full thickness tears and studies that evaluated MRA for the detection of partial thickness tears. The criteria for a positive diagnostic test (index tests and reference standard) varied between studies.

Meta-analyses were not possible for studies that assessed MRA for detection of any rotator cuff tears or partial thickness tears. We found no statistically significant differences in sensitivity or specificity between MRI and US for detecting any rotator cuff tears (P = 0.13), or for detecting partial thickness tears (P = 1.0). Similarly, for the comparison between MRI, MRA and US for detecting full thickness tears, there was no statistically significant difference in diagnostic performance (P = 0.7). For any rotator cuff tears, the summary sensitivity and specificity were 98% (95% CI 92% to 99%) and 79% (95% CI 68% to 87%) respectively for MRI (6 studies, 347 shoulders), and 91% (95% CI 83% to 95%) and 85% (95% CI 74% to 92%) respectively for US (13 studies, 854 shoulders). For full thickness tears, the summary sensitivity and specificity were 94% (95% CI 85% to 98%) and 93% (95% CI 83% to 97%) respectively for MRI (7 studies, 368 shoulders); 94% (95% CI 80% to 98%) and 92% (95% CI 83% to 97%) respectively for MRA (3 studies, 183 shoulders); and 92% (95% CI 82% to 96%) and 93% (95% CI 81% to 97%) respectively for US (10 studies, 729 shoulders).

Because few studies were direct head-to-head comparisons, we could not perform meta-analyses restricted to these studies. The test comparisons for each of the three classifications of the target condition were therefore based on indirect comparisons which may be prone to bias due to confounding.

Authors' conclusions

MRI, MRA and US have good diagnostic accuracy and any of these tests could equally be used for detection of full thickness tears in people with shoulder pain for whom surgery is being considered. The diagnostic performance of MRI and US may be similar for detection of any rotator cuff tears. However, both MRI and US may have poor sensitivity for detecting partial thickness tears, and the sensitivity of US may be much lower than that of MRI. The strength of evidence for all test comparisons is limited because most studies were small, heterogeneous and methodologically flawed, and there were few comparative studies. Well designed studies that directly compare MRI, MRA and US for detection of rotator cuff tears are needed.

PLAIN LANGUAGE SUMMARY

Diagnostic tests for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considered

This summary of a Cochrane review presents what we know from research about the accuracy of imaging tests to detect tears of the rotator cuff tendons in the shoulder.

The rotator cuff is a group of tendons involved in the positioning and moving of the shoulder joint. The rotator cuff lets people lift their arm and reach overhead. In a lot of people, wear and tear of the rotator cuff tendons is a normal part of ageing and they may not have symptoms. However, many people will develop pain in their shoulder at some point as the tendons degenerate further and tears in the rotator cuff tendons develop. There may also be inflammation of the shoulder tendons or bursa (a sac with internal gliding surfaces that helps the shoulder to move). Often the pain is made worse by sleeping on the affected shoulder and moving the shoulder in certain directions. Often there will be pressure on the tendons by the overlying bone when lifting the arm up. This is called impingement. It may become difficult to use the shoulder in every day activities, sports or work.

If the pain does not go away by itself or with treatments such as steroid injections or physiotherapy, surgery may be performed. Imaging tests such as magnetic resonance imaging (MRI), magnetic resonance arthrography (MRA) and ultrasound (US) are used to assess the presence and size of rotator cuff tears to assist in the planning of the surgery.

Rotator cuff tears can be classified as full or partial thickness tears based on the extent or size of the tears. No test is 100% accurate in identifying tears or assessing their size. The accuracy of the tests is commonly assessed by the sensitivity of the test (the proportion of people who had a tear according to the test, among patients with tears), and specificity (the proportion of people without tears on the test, among patients with no tears).

We searched electronic databases up to February 2011, as well as trial registers, conference proceedings and reference lists of articles, for studies comparing diagnostic tests for people with suspected rotator cuff tears. Our review included 20 studies (1147 shoulders). Many studies had design flaws, which limited the reliability of their findings. We found that MRI, MRA and US may have similar accuracy for detecting the presence of full thickness tears. For identifying any tears (no distinction between partial or full thickness) or identifying partial thickness tears, MRI and US may also have similar accuracy. However, it appears that compared with US, MRI may be more sensitive in



identifying partial thickness tears. With these results we can conclude that all three imaging tests (MRI, MRA and US) may help decisions regarding referral for surgery for people with suspected full thickness tears. Information on adverse effects of using these tests was not reported by the included studies.

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Summary of findings 1. Summary of findings table

What is the bes	it imaging modality for rotator cuff tears?								
Patient popu- lation	Patients with shoulder pain suspected of having a rotator cuff tear for whom surgery is being considered.								
Prior testing	Clinical examination.								
Settings	Secondary or tertiary care.								
Index tests	Magnetic resonance imaging (MRI), magnetic resonance arthroscopy (MRA) and ultrasonography (US).								
Reference standard	Arthroscopy and/or open (including mini-open) surgery findings.								
Target condi- tion	Rotator cuff tears: any tear or full or partial thickness tears.								
Importance	Imaging tests are usually performed to determine the characteristics of the rotator cuff tears in order to plan surgery.								
Included studies	We included 20 (1147 shoulders) prospective accuracy studies that evaluated at least of one of the tests.								
studies	Six of the 20 studies reported results for 2 tests evaluated in the same patients.								
Quality con-	Patient characteristics and study design were poorly reported.								
cerns	Most of the QUADAS items were scored unclear for many studies.								
Limitations	We observed considerable variation in results between studies, especially for US studies.								
	Criteria for test positivity (index tests and reference standard) varied between studies.								
	We could not formally investigate potential sources of heterogeneity due to the number of studies available for each test or because most studies reported the same value for a covariate.								
	Our findings were based on small studies with poor reporting of patient characteristics and study design.								
	Because there were few comparative studies, test comparisons relied on indirect evidence which may be confounded by differences in patient and study design characteristics.								
	No study evaluated MRA, MRI and US in the same population.								

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Test	Studies	Cases/Shoulders	Summary sensitivi- ty (95% CI)	Summary specifici- ty (95% CI)	Consequences in a						
					Median preva- lence% (range) ¹	Missed cases ²	Over				
							treated ²				
Any rotator cu	ff tears										
MRI	6	263/347	98 (92, 99)	79 (68, 87)	80 (34 to 96)	2	4				
US	13	626/854	91 (83, 95)	85 (74, 92)	-	7	3				
MRA ³	3										
Full thickness tears											
MRI	7	193/368	94 (85, 98)	93 (83, 97)	56 (17 to 80)	3	3				
US	10	4	3								
MRA	3	3	4								
Partial thickne	ess tears										
MRI	6	83/347	74 (59, 85)	93 (84, 97)	14 (3 to 54)	4	6				
US	8 121/660 52 (33, 70) 93 (85, 97) 7										
MRA ³	4	65/233	-	-	-	-	_				
Comparisons o	of the imaging te	sts for each type of tear									
Comparison	Findings										
MRIand US for	We did not per	form meta-analysis of MRA	studies.								
any rotator cuff tears	The sensitivity	of MRA ranged between 72	% and 100% and the spe	cificities between 5% an	nd 82% in the three stu	dies.					
	There was a 7%	6 difference in the sensitivi	ties of MRI and US, and a	6% difference in specific	cities. The differences v	were not statistically	significant (P = 0.13).				
	Given a mediar	n prevalence of 80%, 80 ou	t of 100 patients will have	any rotator cuff tears.							
	Of the 80 cases	, two will be missed if test	ed using MRI or seven will	be missed if tested usin	g US.						

	Of the 20 patients without any rotator cuff tears, four will undergo unnecessary surgery if tested using MRI or three If tested using US.
MRI, US and MRA for full thickness tears	MRA, MRI and US had very similar sensitivities and specificities. There was no evidence to suggest a difference in sensitivity or specificity between the tests (P = 0.7).
MRI and US for partial thickness tears	We did not perform meta-analysis of MRA studies. The sensitivity of MRA studies ranged between 62% and 80% and the specificities between 47% and 100% in the four studies. Comparing MRI and US, the specificities were found to be similar and despite the large difference in sensitivity (22%), there was no evidence to suggest a difference in sensitivity or specificity between both tests (P = 1.0). Given a median prevalence of 14%, 14 out of 100 patients will have partial thickness tears. Of the 14 cases, two will be missed if tested using MRI but if tested using US, seven will be missed. Of the 20 patients without any rotator cuff tears, four patients will undergo unnecessary surgery if tested using MRI or three patients If tested using US.

Conclusions: MRI, US and MRA have good diagnostic accuracy for detection of full thickness tears and may perform similarly.

The performance of MRI and US may be comparable for detection of partial thickness tears or for detection of any rotator cuff tears.

The strength of the evidence for all test comparisons is limited because most studies were small and methodologically flawed, and there were few comparative studies.

Well designed studies that directly compare the three tests for detection of rotator cuff tears are needed.

There was limited evidence on the best test to diagnose subscapularis tendons tears.

- ¹ The median prevalence and range were computed using all the studies that evaluated each target condition.
- ² Missed and over treated numbers were computed using the median prevalence for each target condition.
- ³ Meta-analyses were not performed for studies that evaluated partial thickness tears and those of any rotator cuff tears because there were few studies and considerable heterogeneity in estimates of sensitivity or specificity.



BACKGROUND

Target condition being diagnosed

The rotator cuff is composed of the subscapularis, supraspinatus, infraspinatus and teres minor tendons; the long head of the biceps tendon also contributes to the cuff. The role of the rotator cuff is to stabilise the humeral head into the glenoid cavity, preventing the upward migration of the humeral head. The four muscles are recruited during different arm movements. The subscapularis is recruited in internal rotation, the supraspinatus in elevation, and the infraspinatus and teres minor in external rotation (Clark 1992; Favard 2007; Matsen 2008).

Rotator cuff tendinopathy can lead to progressive failure of the rotator cuff, typically progressing from partial to a full thickness tear of the supraspinatus tendon then extending into the infraspinatus tendon or the subscapularis tendon, or both. A review by Lewis 2009 concluded that the pathoaetiology of rotator cuff tears is multifactorial and that tears are correlated with a combination of extrinsic and intrinsic factors, but that more research is necessary to fully understand the aetiology of rotator cuff tears. The extrinsic factors (i.e. those external to the rotator cuff) can be divided into anatomical factors, such as the shape of the acromion (i.e. curved or hooked) and coracoacromial ligament, os acromiale and acromial spurs (Baring 2007; Bigliani 1991; Lewis 2009; Neer 1972; Neer 1983; Nho 2008), and environmental factors including aging, shoulder overuse, smoking, obesity and some metabolic disorders such as diabetes (Chen 2003; Galatz 2006; Harryman 2003; Lewis 2009; Nho 2008; Wendelboe 2004). The intrinsic factors include, among others, repetitive microtrauma, areas of hypoperfusion in the tendons, inflammation and cellular changes in the tendons such as disorganisation of the architecture of collagen (Biberthaler 2003; Levy 2008; Lewis 2009; Nirschl 1989; Rees 2008).

Shoulder pain is very common, with an incidence of 9.5 per 1000 patients in primary care in Cambridge, UK, where amongst them 85% presented with rotator cuff tendinopathy (Ostör 2005). Disorders of the rotator cuff tendons due to either wear or tear are among the most common causes of shoulder pain and disability. In Japan, the prevalence of rotator cuff tears is 20.7% in the general population and 36% in patients with shoulder pain (Yamamoto 2010). More than 4.5 million physician visits occurred and approximately 40,000 inpatient surgeries were performed for rotator cuff problems in the United States in 2002 (Oh 2007).

The diagnosis of rotator cuff tears is mainly based on the patient's history and physical examination. The value of physical examination of the shoulder has been addressed in another Cochrane review (Hanchard 2013). The clinical manifestations vary widely (Duckworth 1999; Matsen 2008). Acute, traumatic full thickness cuff tears may present with sudden onset of weakness during elevation of the arm after a trauma in which the arm has been forced to the side (like a fall with the arm out to the side or on catching a heavy falling object with the arm extended) (Matsen 2008). Chronic degenerative cuff defects may present with progressive pain and weakness, with concomitant loss of active motion. Pain in the lateral area of the shoulder is commonly present at night. Passive motion initially remains full until the pain limits active motion (Baring 2007; Matsen 2008). However, there are many people with degenerative rotator tears who are asymptomatic (Reilly 2006; Zanetti 2000).

Decisions about whether to order a diagnostic test include consideration of whether the results are likely to affect treatment. Plain radiographs of the shoulder may be useful to differentiate rotator tears from osteoarthritis of the glenohumeral or acromioclavicular joints and calcific tendonitis. Ultrasonography (US), magnetic resonance imaging (MRI) and magnetic resonance arthrography (MRA) are increasingly being used to detect rotator cuff tears, although who orders these tests may vary by setting. In some settings, these tests are mainly ordered by specialists but in other settings they are being ordered by primary care physicians or clinicians (Al-Shawi 2008; Miller 2008). In the context of specialist care, US or MRI, or both, are usually performed to determine the characteristics of the rotator cuff tears in order to plan surgery. In some settings, however, there has been a significant rise in the number of diagnostic US being performed in primary care. For example, in Australia there has been a more than fourfold increase, from 104,252 in the year 2000 to 2001 to 440,172 in 2008 to 2009 (Medicare Australia 2010). However, the utility of the test to affect treatment in primary care is unknown.

Tears of the rotator cuff can be classified in several ways: duration (acute or chronic), aetiology (traumatic or degenerative) or size (partial or full thickness). Radiologists often describe the size of tear in millimetres or centimetres or descriptively as small, medium, large or massive. All three factors (duration, aetiology and size) influence treatment decisions (Kuhn 2007).

Acute full thickness rotator cuff tears are uncommon and account for less than 10% of all rotator cuff tears. People with acute full thickness tears usually present with a history of acute trauma, such as a fall or dislocation, and immediate pain and weakness. Prompt surgical treatment, ideally within six weeks, is the recommended treatment (Rees 2008). For all other full thickness rotator cuff tears, surgical treatment is usually reserved for those who fail to improve after a period of conservative treatment, although the most effective surgical intervention and its timing remain uncertain (Coghlan 2009; Dunn 2005; Oh 2007; Rees 2008). For example, a delay in surgical repair of a large tear may allow the injured tendon to retract and the muscle to atrophy (Matsen 2008; Oh 2007). On the other hand, asymptomatic tears are common; these are chronic tears that normally do not compromise the function of the shoulder. A recent review reported the prevalence of full thickness tears in 2553 unselected cadavers as 30% (Rees 2008). Furthermore, the pathogenesis and progression to symptomatic tears remains unclear (Rees 2008). In addition, in contrast to acute full thickness tears, symptoms due to acute or chronic partial thickness cuff tears frequently improve with conservative interventions (Matava 2005; Matsen 2008).

While spontaneous healing of a partial thickness tear is unlikely in most cases, the explanation for the 'cure' with conservative treatment is due to the likely resolution of the accompanying inflammation over time and may also be related to the residual cuff muscles compensating for the mechanical deficiency of the torn cuff (Fukuda 1996; Fukuda 2003; Matava 2005; Matsen 2008). As with full thickness tears, no simple treatment algorithm for partial thickness rotator cuff tears exists. Surgical treatment, however, is normally indicated for people with persisting symptoms despite conservative treatment and in whom imaging suggests the presence of a partial thickness tear or tears. The ideal timing of surgical intervention also remains unclear (Fukuda 2003; Matava 2005). However, case series and anecdotal evidence suggest that



satisfactory results are usually achieved with surgery provided there is a good blood supply to the tendon, contact between the torn ends, absence of retraction and adequate trophic quality of the muscle (Fukuda 2003).

Another recognised category of tears is massive complete tears, in which a large area of the humeral head is uncovered (Wolfgang 1974). Post 1983 defined a massive tear as greater than 5 cm. These tears, which are difficult to repair, are more commonly found in women over 65 years of age and are associated with advanced atrophy, degeneration and progressive fatty infiltration of the rotator cuff muscles (Dines 2007; Gerber 2000). Treatment options for these massive, retracted tears are limited as they are often deemed irreparable. In younger people, tendon transfers may be considered (Neri 2009).

The indications for surgical treatment of rotator cuff tears have not been fully defined. A systematic review of surgical treatment for rotator cuff disease (including tears), which included 14 trials, was unable to draw firm conclusions about the effectiveness of surgery (Coghlan 2009). Nonetheless, the review suggested that there were no significant differences in outcomes between open or arthroscopic surgery and non-surgical treatment (Coghlan 2009). Many studies have demonstrated that the size of the tear is correlated to the final outcome; partial or small full thickness tears usually have a satisfactory surgical result (Bianchi 2005; Bryant 2002; Fotiadou 2008).

Index test(s)

Currently, US, MRI or MRA are usually performed in patients contemplating surgery for rotator cuff tears to determine the characteristics of the tears. With the improvement of technology, the accuracy of these imaging tests is considered to have improved significantly over time, enabling useful assessment of the size and extent of the rotator cuff tear when planning surgery (Rees 2008).

US is a diagnostic imaging technique used to visualise deep structures of the body by recording the echoes of pulsed ultrasonic waves directed into the tissues and reflected by tissue planes to the transducer. These echoes are converted into 'pictures' of the tissues under examination. Seltzer 1979 was the first to describe ultrasonographic evaluation of rotator cuff diseases. US of the shoulder is utilised in secondary, tertiary and, increasingly, primary healthcare settings to evaluate the integrity of the rotator cuff. It consists of a non-invasive examination that has practically no adverse effects and allows dynamic visualisation of the tendons during movement of the shoulder (Al-Shawi 2008). However, operator dependence and a long learning curve are frequently considered to be its limitation (O'Connor 2005; Rutten 2006), principally in view of partial thickness tears for which Le Corroller 2008 described a high interobserver variability.

MRI uses a powerful magnetic field to align the hydrogen atoms of water and other molecules in the body. Pulses of radiofrequency are applied which excite the magnetised atoms. These movements of hydrogen atoms, which vary in different tissues, are captured and the signal can be manipulated to build up an image of the body (Witte 2003). The first article about the use of MRI in the shoulder was published in 1986 (Kneeland 1986). Since then, this technique has been widely used in secondary and tertiary healthcare practice. MRI is a non-invasive method of imaging that is unique in allowing high resolution images in multiple planes. It is a static

examination that may be enhanced by an intra-articular injection of radiopaque dye (this is called magnetic resonance arthrography). The radiopaque dye acts as contrast material that helps to delineate intra-articular structures and outline abnormalities. MRA of the shoulder is also useful for assessing the rotator cuff integrity. In comparison with conventional MRI, MRA may improve diagnostic performance in detecting shoulder diseases; however, any potential benefit from MRA must be weighed against the invasiveness and additional discomfort caused by the procedure.

MRI and MRA have some absolute contraindications, such as the presence of intracerebral aneurysm clips, cardiac pacemakers, automatic defibrillators, biostimulators, implanted infusion devices, cochlear implants and metallic orbital foreign bodies (Witte 2003). They are also expensive and time consuming procedures. The strength of the magnet, the sequences used in the examinations and the person (e.g. consultant radiologist, musculoskeletal radiologist or trainee) interpreting and reporting the test may all affect the results.

Summary of diagnostic pathway

The evaluation of patients with suspected rotator cuff tear(s) should initiate with a full history of the patient's complaints and a thorough clinical examination of the shoulder. Decisions for using an imaging diagnostic test may be supported by whether the results are likely to affect treatment. For example, MRI, MRA or US might confirm a possible full thickness tear. The three index tests considered can also be used as triage tests in people suspected of having partial thickness tears. People whose tests were positive can be treated as having partial tears, while people with rotator cuff symptoms whose tests were negative can undergo further diagnostic procedures, such as diagnostic arthroscopy.

Reference tests

The reference tests for diagnosis of rotator cuff tears are invasive. The most common reference test is diagnostic arthroscopy. Arthroscopy is a minimally invasive surgical procedure that involves insertion of an arthroscope, a type of fibre-optic endoscope, into the joint through a small incision. This allows the surgeon to inspect and probe the articular (joint) and bursal side of the rotator cuff tendons, to assess accurately the rotator cuff insertion (footprint) and to perform a general examination of the shoulder joint in order to identify and treat other potential lesions (Dinnes 2003; Matava 2005). However, limitations associated with diagnostic shoulder arthroscopy include the need for anaesthesia, hospital admission and some interobserver variation in the classification of tears (Kuhn 2007).

Open surgery (including mini-open) has also been used as a reference test although it is more limited than arthroscopy because joint surface or inferior surface tears are difficult to access and identify using an open approach. Thus open surgery is less accurate than arthroscopy for detecting partial rotator cuff tears.

Rationale

US, MRI and MRA are increasingly being used to assess the presence and size of rotator cuff tears to assist in planning surgical treatment. Improved techniques have resulted in increased reliance on these tests, in place of a separate diagnostic arthroscopy, although arthroscopic examination of the shoulder joint is still commonly performed as part of surgical treatment. US, MRI and MRA are



operator and reader dependent. It is not known whether any one test is superior to either of the two others or whether performing US and MRI or US and MRA enhances their value (Swen 1999). It is also not known whether these diagnostic tests provide useful additional information compared with diagnostic arthroscopy, which is an accepted part of the surgical treatment. While, the units costs of MRI and MRA are greater than US, the cost-effectiveness of the three tests has not been determined.

We identified two relevant systematic reviews with meta-analyses that assessed diagnostic imaging tests for rotator cuff disease (De Jesus 2009; Dinnes 2003). The literature search in both reviews was restricted to English language only. Dinnes 2003 evaluated the diagnostic accuracy of clinical testing of US, MRI and MRA for detecting rotator cuff tears using both surgical and non-surgical tests as the reference standard. The authors included 38 studies that assessed the accuracy of US, 29 studies that assessed the accuracy of MRI and 6 studies that assessed the accuracy of MRA and concluded that US or MRI were equivalent for detecting full thickness rotator cuff tears, although MRI was more expensive and US may be better at detecting partial tears. The search date for the review was October 2001. A later review with a search date in September 2007, De Jesus 2009, conducted a meta-analysis comparing the diagnostic accuracy of US and MRI for rotator cuff tears using surgery as the reference standard. This systematic review included 65 studies but the appraisal of the methodological quality of the included studies was unclear or inadequate. De Jesus 2009 concluded that US is as accurate as MRI for both full and partial thickness rotator cuff tears and also suggested that US may be the most cost-effective imaging test for detecting rotator cuff tears.

Important technological improvements in US, MRI and MRA have been made since the search dates of both systematic reviews, and new studies evaluating US, MRI and MRA have been published. Our review involves an updated search for diagnostic accuracy studies for rotator cuff tears and will not be restricted to English language publications.

OBJECTIVES

To compare the diagnostic test accuracy of magnetic resonance imaging (MRI), magnetic resonance arthrography (MRA) and ultrasonography (US) for detecting any rotator cuff tears (i.e. partial or full thickness) in people with shoulder pain for whom surgery is being considered.

We divided our objectives as follows.

- To compare the diagnostic accuracy of US, MRI and MRA for diagnosing any rotator cuff tears (partial or full thickness)
- To compare the diagnostic accuracy of US, MRI and MRA for diagnosing full thickness rotator cuff tears (one or more tendons)
- To compare the diagnostic accuracy of US, MRI and MRA for diagnosing partial thickness rotator cuff tears

Investigation of sources of heterogeneity

We planned to investigate the following potential sources of heterogeneity:

• Type of tear: acute traumatic and chronic degenerative

 Type of reference standard: open (including mini-open) surgery or arthroscopy

METHODS

Criteria for considering studies for this review

Types of studies

All diagnostic accuracy studies that compared one or more of the index tests with one or both of the reference tests in patients suspected of having a partial or full thickness rotator cuff tear were included. We only included results from full reports of prospective studies. Studies with an excessively long period of time (i.e. a year or longer) between the index and reference tests were excluded because there is evidence that rotator cuff tears can progress over time (Mall 2010; Melis 2010); however, the rate of progression is not clearly defined.

We included articles in English and languages for which a full translation could be obtained. Non-English articles where a full translation could not be obtained are cited in the Characteristics of studies awaiting classification but not included in the review.

For studies reported in multiple publications, we included only the most recent or complete report. References to the other publications were cited under the same study identifier.

Participants

We included people with shoulder pain suspected of having a rotator cuff tear for whom surgery was being considered. Studies that included healthy controls or participants who had been previously diagnosed with other specific shoulder pain (e.g. shoulder instability, osteoarthritis, rheumatoid arthritis, frozen shoulder, benign or malignant tumours or referred pain) were excluded. Studies that included participants with shoulder pain, but in which it was unclear if all the participants were suspected of having rotator cuff tears, were also excluded.

Index tests

Studies that assessed the accuracy of US, MRI or MRA were included.

Target conditions

We included studies that evaluated the index tests for detection of at least one of three target conditions:

- presence of any rotator cuff tears (partial or full thickness);
- · presence of full thickness tears;
- · presence of partial thickness tears.

To standardise classification for this review, rotator cuff tears were dichotomised as absence or presence of any, full and partial thickness tears.

Reference standards

We required arthroscopy or open (including mini-open) surgery findings to be the reference standards.



Search methods for identification of studies

Electronic searches

We searched relevant computerised databases for eligible diagnostic studies: MEDLINE (PubMed) (1966 to March 2011), EMBASE (Elsevier) (1980 to February 2011), LILACS (Bireme) (1982 to February 2011) and the Cochrane Register of Diagnostic Test Accuracy Studies (February 2011). We also searched DARE (Database of Abstracts of Reviews of Effects), the HTA Database (Health Technology Assessments Database) and the MEDION database (February 2011) for other related diagnostic test accuracy reviews, and we checked the reference lists of those reviews that were relevant for additional studies. We also searched the US Health Services Research Projects in Progress and the UK Clinical Research Network Portfolio Database for ongoing and recently completed studies. When possible, non-English articles were assessed through translation by a native speaker.

We used a sensitive search strategy as recommended by the Cochrane Collaboration for MEDLINE (PubMed), EMBASE (Elsevier) and LILACS (Bireme) (De Vet 2008). See Appendix 1 for the MEDLINE and EMBASE search strategies.

Searching other resources

We checked the reference lists of articles, reviews and textbooks for relevant primary diagnostic studies and systematic reviews. We handsearched abstracts of the British Elbow and Shoulder Society annual meetings (2005 to July 2011) and American Academy of Orthopaedic Surgeons annual meetings (2005 to July 2011). We also contacted experts in the field.

Data collection and analysis

We used the methods suggested in the Cochrane Handbook for Systematic Reviews of Diagnostic Test Accuracy (Deeks 2009).

Selection of studies

Two review authors (ML and RJ) independently screened the titles and abstracts of retrieved records to identify potentially relevant studies for inclusion. Duplicates were removed and the remaining references were examined. Studies which clearly did not meet the inclusion criteria were excluded, and copies of the full text of potentially relevant references were obtained. ML and RJ independently assessed full text reports and determined inclusion or exclusion of the studies. Any uncertainties or disagreements were resolved by discussion and, when necessary, by adjudication from a third author (RB).

Data extraction and management

Two review authors (ML and RJ) independently collected the available data using a piloted data extraction form without masking of study authors or other identifying information. A third review author (RB) was consulted for resolution of any disagreements. When necessary, we sent requests to study authors for additional information or data. Diagnostic accuracy studies that reported insufficient data for construction of two-by-two tables were excluded from the review.

We retrieved the following data.

 General information: title, journal, year, publication status, country of study, period of study, primary objective and study

- design (i.e. prospective versus retrospective and consecutive versus non-consecutive).
- 2. Sample size: number of participants meeting the criteria and total number screened.
- Baseline characteristics: baseline diagnosis, age, sex, dominant arm, nature of onset (e.g. traumatic or non-traumatic), duration of symptoms, prior treatment, inclusion and exclusion criteria.
- 4. Target condition as reported.
- 5. Index test: description of technique, criteria for positive result, timing of test and expertise of the clinician or technician performing the test.
- Reference standard test: description of technique, criteria for positive result, time from index to reference test and expertise of the clinician or technician performing the test.
- 7. Adverse effects or complications due to index test(s) and reference standard test(s).
- 8. Number of true positives (TP), true negatives (TN), false positives (FP) and false negatives (FN). We extracted data for operational definitions for category of tear (e.g. partial, full or any thickness tears). Multiple outcome categories are often reported for rotator cuff tears: partial thickness tear, full thickness tear and no tears (i.e. three-by-three tables). Currently available methods for evaluating diagnostic tests rely on dichotomised disease status. Therefore, for the assessment of each target condition, we dichotomised rotator cuff tears using a strategy based on the options for treatment. To create two-by-two tables for partial thickness tears, data for full thickness tears were included with those for no tears. We did not exclude data for any category. We included data for partial thickness tears with those for full thickness tears to create two-by-two tables for any tears.

Assessment of methodological quality

The methodological quality of the included studies was assessed independently by two review authors (ML and RJ) and disagreement on study quality was resolved by a third review author (RB). At the same time as data extraction, the methodological quality of selected studies was assessed using a modified version of the QUADAS checklist (Whiting 2003), following the guidelines provided in Chapter 9 of the Cochrane Handbook for Systematic Reviews of Diagnostic Test Accuracy (Reitsma 2009). Appendix 2 explains how we applied the QUADAS items for assessing the included studies.

Statistical analysis and data synthesis

Our unit of analysis was the shoulder. For each test and target condition, estimates of sensitivity and specificity from each study were plotted in receiver operating characteristic (ROC) space and forest plots for visual examination of variation in test accuracy. Where adequate data were available, we conducted meta-analyses using the bivariate model (Chu 2006; Reitsma 2005). In the bivariate model, the logit-transformed sensitivities and specificities, and the correlation between them across studies are modelled directly. The model accounts for sampling variability within studies and also accounts for between study variability through the inclusion of random-effects. In preliminary metaanalyses for each target condition, we fitted the bivariate model separately for each test. We examined the variance of the randomeffects parameters to consider the magnitude of heterogeneity and to judge whether there were differences in heterogeneity in sensitivities and specificities between tests, before comparing the



tests in a single model for formal assessment of comparative accuracy.

Comparative accuracy studies are scarce (Takwoingi 2013). Therefore, whenever possible, we included all studies of US, MRI and MRA (i.e. an indirect comparison) in the main comparative meta-analysis for each target condition. Due to few studies of MRA and considerable heterogeneity in study results, we only performed pairwise comparisons of MRI and US for detection of partial thickness tears and any tears but compared the three tests for detection of full thickness tears. We compared test accuracy by adding covariate terms for test type to the parameters of the bivariate model to determine which test was superior in terms of sensitivity or specificity or both. The variance coefficients from the preliminary meta-analysis and summary ROC plot for each test indicated differences in heterogeneity between tests and so we extended the bivariate model to allow the variances of the random-effects to vary with test type. We assessed the statistical significance of the difference in sensitivity or specificity between tests by using likelihood ratio tests comparing models with and without the covariate terms in the bivariate model. The summary sensitivities and specificities (i.e. average operating points) were plotted on summary ROC plots with corresponding 95% confidence regions. Summary positive and negative likelihood ratios were derived from functions of the bivariate model parameters, with 95% confidence intervals computed using the delta method.

Indirect comparisons of tests are not ideal and are susceptible to bias because other factors, such as participant and study design characteristics, may confound differences between tests. Thus in secondary analyses, we restricted the test comparisons to only studies that evaluated the tests in the same population. Because the studies were few, we were unable to perform meta-analyses but used linked summary ROC plots where estimates for each of the two tests from each study are joined by a line to illustrate the results. Furthermore, for each target condition, we quantified the difference in sensitivities and specificities between pairs of tests by computing differences in these proportions together with the corresponding 95% CI. Thus we visually and numerically demonstrated the change and consistency of the direction of the change in test performance between the tests. We used the xtmelogit command in Stata version 11.2 (StataCorp, College Station, Texas) to fit the bivariate models.

Investigations of heterogeneity

Heterogeneity was investigated in the first instance through visual examination of forest plots and summary ROC plots. The type of tear and type of reference standard reported in each study were presented on forest plots along with the estimates of sensitivity and specificity. In exploratory analyses, we ordered studies on the forest plots by each of the two covariates in turn and also by sensitivity or specificity to examine the pattern of variation between studies. If there were sufficient data we planned to formally investigate heterogeneity by adding covariates to the bivariate model for each potential source of heterogeneity.

Sensitivity analyses

If there were sufficient studies, we performed sensitivity analyses by comparing results based on all studies with results of subsets of studies that complied (scored 'Yes') with the following methodological quality items of the QUADAS checklist (Whiting 2003).

- · Representative spectrum
- Acceptable reference standard
- Acceptable delay between tests
- · Index test results blinded
- · Reference standard results blinded

We also investigated the effect of unit of analysis by excluding studies that included both shoulders for any individual.

RESULTS

Results of the search

The search strategy identified 3169 references and the handsearch identified an additional three records (Figure 1). Of these, 2902 were excluded by initial screening of reference titles and abstracts. There were 974 duplicates and 1926 were either not relevant or did not meet the inclusion criteria. We were unable to obtain full text articles for two studies because they were not available from libraries or vendors.



Figure 1. Study flow diagram

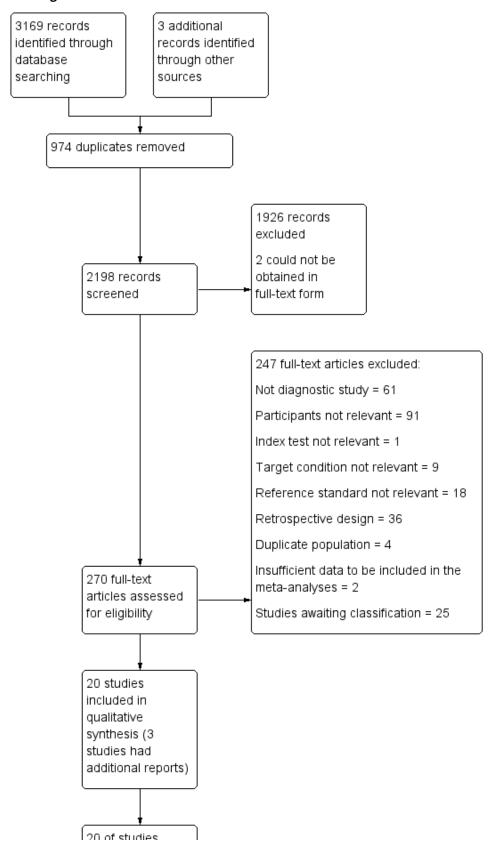




Figure 1. (Continued)

20 of studies included in quantitative synthesis (meta-analysis)

Of the 270 potentially eligible studies that were remaining and for which full reports were obtained (192 were reported in English and 78 in a non-English language), 20 studies met our inclusion criteria and were included in the review. Three of the included studies had additional published data. Two hundred and eighteen studies did not meet our inclusion criteria and were excluded (see Characteristics of excluded studies) and four reported on the same population or a subset of an already excluded study. At the time of publication, we are still awaiting translation of 25 non-English articles that are potentially relevant based upon their title and abstract; these are listed in Studies awaiting classification. Data from these studies will be added in future updates of this review if the studies are found to be eligible for inclusion.

Among the 20 included studies, six (lannotti 2005; Kang 2009; Martin-Hervas 2001; Sipola 2010; Swen 1999; Teefey 2004) evaluated the accuracy of two different tests against the reference standard(s). See the Characteristics of included studies for details of the individual studies.

Methodological quality of included studies

The methodological quality of the 20 included studies was judged to be low or unclear for most categories and is summarised in Figure 2. The quality assessment results for the individual studies can be found in Figure 3 and details are given in the Characteristics of included studies.

Figure 2. Methodological quality graph: review authors' judgements about each methodological quality item presented as percentages across all included studies

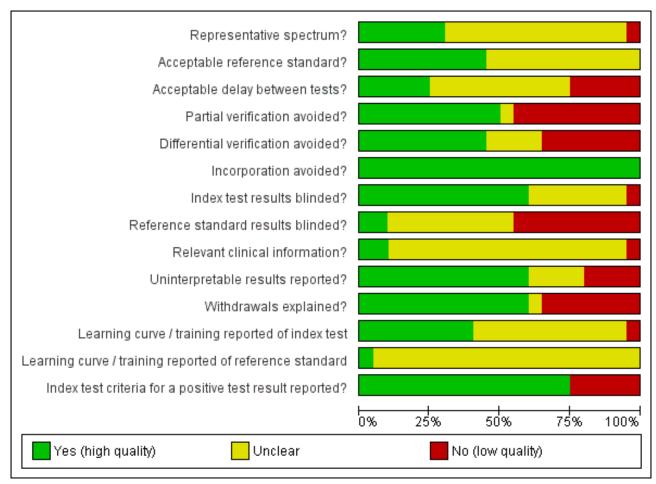




Figure 3. Methodological quality summary: review authors' judgements about each methodological quality item for each included study

	Representative spectrum?	Acceptable reference standard?	Acceptable delay between tests?	Partial verification avoided?	Differential verification avoided?	Incorporation avoided?	Index test results blinded?	Reference standard results blinded?	Relevant clinical information?	Uninterpretable results reported?	Withdrawals explained?	Learning curve / training reported of index test	Learning curve / training reported of reference standard	Index test criteria for a positive test result reported?
De Candia 2002	?	?	•	•	•	•	?	•	?	•	•	?	?	-
Della Sala 1996	?	?	?	•	•	•	?	?	?	•	•	?	?	•
Gagey 1993	?	?	?	•	•	•	?	?	?	•	•	?	?	•
lannotti 2005	•	?	?	?	?	•	•	•	•	?	?	•	?	•
Kang 2009	?	•	•	•	•	•	•	•	?	•	•	•	?	•
Lambert 2009	•	?	•	•	?	•	?	•	?	?	•	?	?	•
Martin-Hervas 2001	?	?	?	•	?	•	•	?	?	?	•	?	?	•
Milosavljevic 2005	?	•	•	•	•	•	•	?	?	•	•	?	?	•
Misamore 1991	?	?	?	•	•	•	•	•	?	•	•	•	?	•
Mohtadi 2004	•	•	•	•	•	•	•	•	?	•	•	?	?	•
Nicoletti 1994	?	•	?	•	?	•	?	?	?	•	•	•	•	•
Sipola 2010	•	?	•	•	•	•	?	?	?	•	•	•	?	•
Stetson 2005	?	•	?	•	•	•	?	•	?	•	•	•	?	
Swen 1998	•	•	?	•	•	•	•	?	?	•	•	•	?	•
Swen 1999	•	•	•	•	•	•	•	?		•	•	?	?	•
Taboury 1992	?	?	?	•	•	•	•	•	?	•	•	?	?	
Teefey 2004	?	•	•	•	•	•	•	•	?	?	•	•	?	•
Venu 2002	?	•	•	•	•	•	•	•	•	•	•	•	?	•
Wallny 2001	•	?	?	•	•	•	•	?	?	•	•	?	?	•
Yen 2004	?	?	•	•	•	•	•		?	•	•	?	?	•



The spectrum of participants (item 1) was judged to be representative in only 6 (30%) of the 20 studies. To be judged representative, studies had to be prospective with consecutive recruitment. The setting had to be secondary or tertiary care and the patients had to present with shoulder pain caused by a suspected rotator cuff tear for which surgery was being considered for treatment. Only half of the studies included an appropriate reference standard (item 2) and avoided partial verification (item 4). The majority (more than 50%) of studies poorly described the following QUADAS items: time period between reference standard and index test (item 3), differential verification bias (item 5), reference standard results blinded (item 8), relevant clinical information (item 9), and learning curve and training reported for both the index and reference standard readers (items 12 and 13) (see Appendix 2 for further explanation of these items). The remaining QUADAS items were well described in 50% to 75% of the included studies: index test results blinded (item 7), un-interpretable results reported (item 10), withdrawals explained (item 11) and index test criteria for a positive result (item 14). Criteria for test positivity was reported by 15 studies and varied between studies; the criteria are presented in detail in the Characteristics of included studies. As we anticipated in our protocol, the answer for 'incorporation avoided' (item 6) was 'Yes' (no bias) for all included studies.

Findings

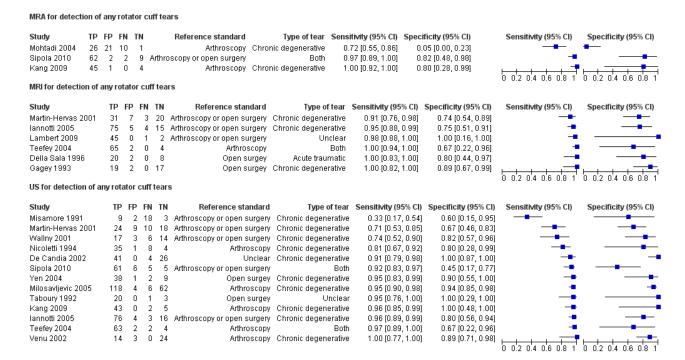
Our meta-analyses were based on indirect comparisons because meta-analyses of studies that directly compared tests were not possible; there were only six comparative studies. No study directly compared MRA and MRI, or all three modalities in the same patients for any of the target conditions. The summary sensitivities and specificities for the tests are shown in Table 1 for each target condition. For MRA, meta-analysis was performed only for studies evaluating detection of full thickness tears due to the few studies and the degree of heterogeneity observed on summary ROC plots for studies evaluating partial thickness tears and any rotator cuff tears.

Two studies (lannotti 2005; Milosavljevic 2005) included both shoulders of one and five patients respectively. The remaining studies reported the same number of patients and shoulders, with the exception of Milosavljevic 2005 where this information was missing.

Detection of any rotator cuff tears

Figure 4 shows the forest plots of the sensitivity and specificity estimates for MRI, US and MRA for the 17 studies that assessed any rotator cuff tears.

Figure 4. Accuracy of MRA, MRI and US for detecting any rotator cuff tears (forest plot)



Six studies, based on 347 shoulders from 346 patients, assessed the diagnostic accuracy of MRI. The median study size was 55 (range 30 to 99), and the median prevalence of any rotator cuff tear was 73% (range 50% to 96%). The sensitivity of MRI reported in the studies ranged from 91% to 100%, and specificity from 67% to 100%. The summary estimates for the sensitivity and specificity of MRI were 98% (95% CI 92% to 99%) and 79% (95% CI 68% to

87%) respectively. The positive and negative likelihood ratios were 5 (95% CI 2 to 10) and 0.03 (95% CI 0.01 to 0.11) respectively.

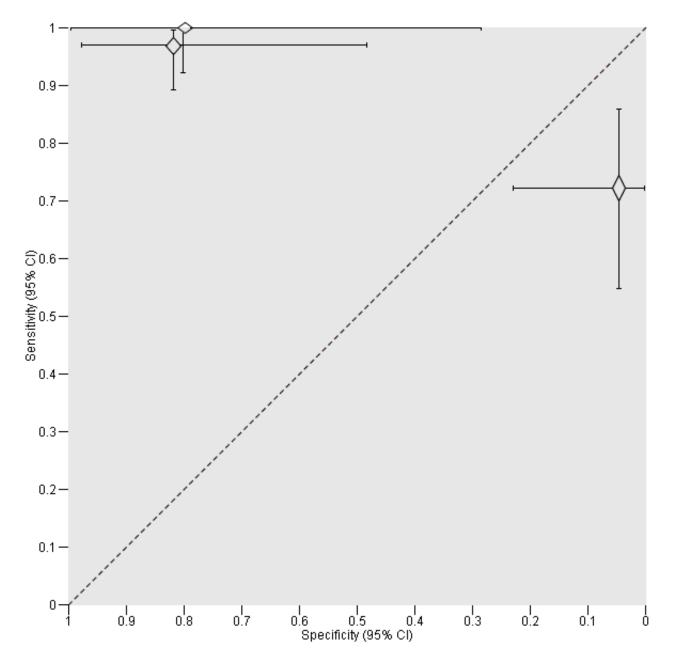
Thirteen studies assessed the accuracy of US to detect any rotator cuff tears. The studies included a total of 854 shoulders from 848 patients with a median study size of 50 (range 24 to 190). The prevalence of any rotator cuff tears in the US studies was 80%



(range 34% to 92%), and the sensitivities ranged from 33% to 100%, specificities from 45% to 100%. The summary sensitivity and specificity of US were 91% (95% CI 83% to 95%) and 85% (95% CI 74% to 92%) respectively. The positive and negative likelihood ratios were 6 (95% CI 3 to 12) and 0.11 (95% CI 0.05 to 0.22) respectively.

Three studies, based on 183 shoulders from 183 participants, assessed the accuracy of MRA for detection of any rotator cuff tears. The median study size was 58 (range 50 to 75), and the median prevalence was 85% (range 62% to 90%). The sensitivity of MRA ranged from 72% to 100%, and specificity from 5% to 80%. Meta-analysis was not performed but study specific estimates of sensitivity and specificity were plotted in ROC space with 95% CI in Figure 5.

Figure 5. Study estimates of sensitivity and specificity with 95% confidence intervals plotted in ROC space for MRA for the detection of any rotator cuff tears



Comparison of MRI and US for detection of any rotator cuff tears

Using the 11 studies that evaluated the accuracy of either MRI or US for detection of any rotator cuff tears, neither test was found to be superior in terms of sensitivity or specificity. Although the

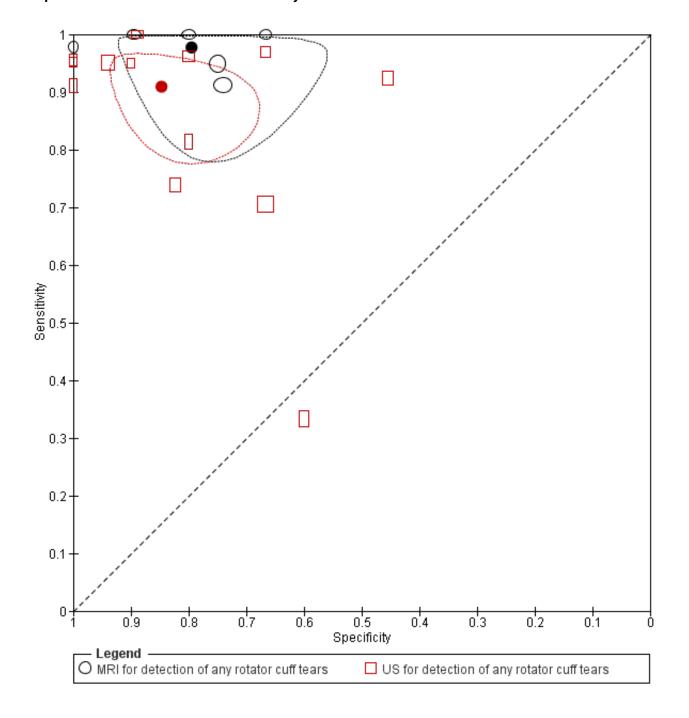
sensitivity of MRI was 7% higher than that of US and the specificity of MRI was 6% lower than that of US (Figure 6; Table 1), there was no statistically significant difference between the two tests (P = 0.13). In the analysis restricted to the three studies (231 shoulders



from 230 patients) that performed head-to-head comparisons of MRI and US within the same patients (Table 2, see Appendix 3 for additional figure), two studies reported higher sensitivity and specificity for MRI compared to US while the other study reported higher sensitivity and specificity for US compared to MRI. For head-

to-head comparisons of MRA and US, there were only two studies (127 shoulders from 127 patients). Both studies reported higher sensitivity for MRA compared to US but the estimates of specificity were conflicting (Table 3).

Figure 6. Study estimates of sensitivity and specificity, and summary points with 95% confidence regions plotted in ROC space for MRI and US for the detection of any rotator cuff tears

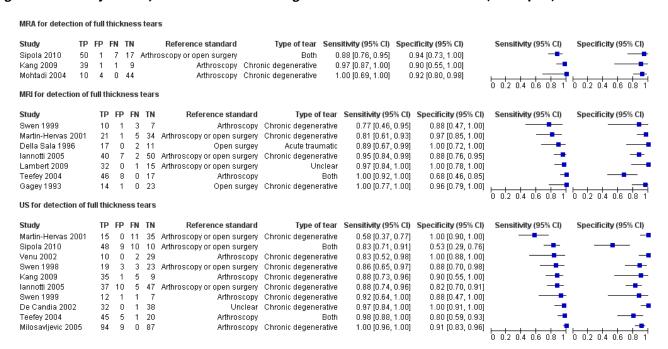




Detection of full thickness rotator cuff tears

The estimates of sensitivity and specificity for the 14 studies that evaluated either MRI, US or MRA for the detection of full thickness rotator cuff tears are shown in Figure 7.

Figure 7. Accuracy of MRA, MRI and US for detecting full thickness rotator cuff tears (forest plot)



Seven studies, based on 368 shoulders from 367 patients, assessed the diagnostic accuracy of MRI. The median study size was 48 (range 21 to 99), and the median prevalence of full thickness rotator cuff tear was 62% (range 37% to 69%). The sensitivities ranged from 77% to 100%, and specificities ranged from 68% to 100%. The summary sensitivity and specificity of MRI were 94% (95% CI 85% to 98%) and 93% (95% CI 83% to 97%) respectively. The positive and negative likelihood ratios were 13 (95% CI 6 to 29) and 0.06 (95% CI 0.02 to 0.16) respectively.

Ten studies (729 shoulders from 723 patients) assessed the accuracy of US to detect full thickness tears. The median study size was 66 (range 21 to 190), and the median prevalence was 48% (range 29% to 80%). Sensitivities ranged from 58% to 100%. Specificities ranged from 53% to 100%. The summary sensitivity and specificity of US were 92% (95% CI 82% to 96%) and 93% (95% CI 81% to 97%) respectively. The positive and negative likelihood ratios were 12 (95% CI 5 to 34) and 0.09 (95% CI 0.04 to 0.20) respectively.

Three studies (the same studies that assessed any rotator cuff tears) assessed the accuracy of MRA to detect full thickness tears with sensitivities ranging from 88% to 100% and specificities ranging

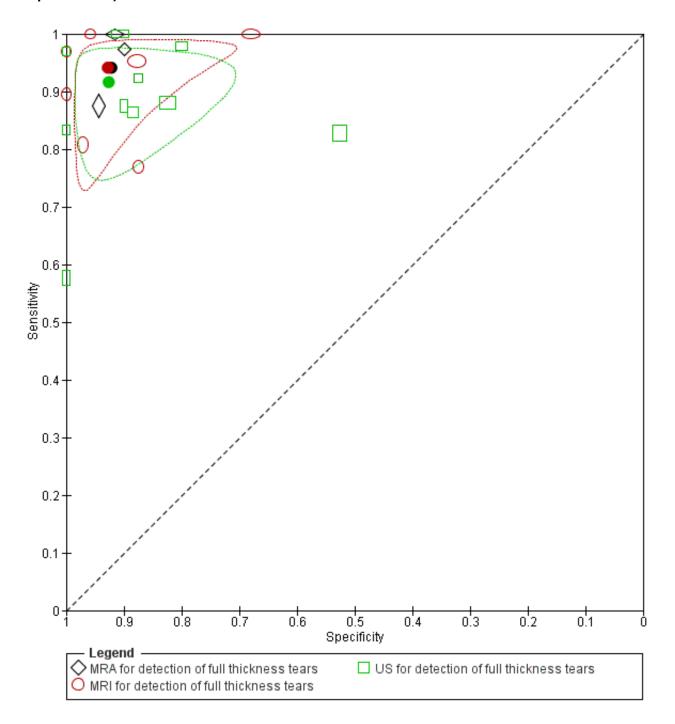
from 90% to 94%. The median prevalence was 76% (range 17% to 80%). The summary sensitivity and specificity of MRA were 94% (95% CI 80% to 98%) and 92% (95% CI 83% to 97%) respectively. The positive and negative likelihood ratios were 12 (95% CI 5 to 30) and 0.06 (95% CI 0.02 to 0.23) respectively.

Comparison of MRI, MRA and US for detection of full thickness rotator cuff tears

Based on the 14 studies that assessed the accuracy of MRI, MRA or US for detection of full thickness rotator cuff tears, the summary sensitivities and specificities of MRI, MRA and US were found to be very similar (Figure 8; Table 1). There was no statistically significant difference in sensitivity or specificity (P = 0.7). Four studies (252 shoulders from 251 patients) directly compared MRI and US (Table 2, see Appendix 3 for additional figure) within the same patients, with no consistency among the studies as to which test was superior in terms of either sensitivity or specificity. Two studies (127 shoulders from 127 patients) directly compared MRA and US (Table 3). Both studies reported higher sensitivity for MRA compared to US. One of the two studies also reported a higher specificity while the other study reported no difference.



Figure 8. Study estimates of sensitivity and specificity, and summary points with 95% confidence regions plotted in ROC space for MRA, MRI and US for the detection of full thickness rotator cuff tears



Detection of partial thickness rotator cuff tears

Figure 9 shows the estimates of sensitivity and specificity for the 13 studies that evaluated either MRI, MRA or US for the detection of partial rotator cuff tears.



Figure 9. Accuracy of MRI, US and MRA for detecting partial thickness rotator cuff tears (forest plot)

MRA for detection of partial thickness tears														
Study	TP	FP	FN	TN		Reference standard		Type of tear	Sensit	ivity (95% CI)	Speci	ificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Mohtadi 2004	16	17	10	15		Arthroscopy	Chroni	ic degenerative	0.6	2 [0.41, 0.80]	0.	.47 [0.29, 0.65]		
Sipola 2010	5	8	2	60	Art	hroscopy or open surgery		Both	0.7	1 [0.29, 0.96]	0.	.88 [0.78, 0.95]		-
Stetson 2005	21	0	6	23		Arthroscopy	Chroni	ic degenerative	0.7	8 [0.58, 0.91]	1.	.00 [0.85, 1.00]	_	-
Kang 2009	4	2	1	43		Arthroscopy	Chroni	ic degenerative	0.8	0 [0.28, 0.99]	0.	.96 [0.85, 0.99]		
MRI for detection	of	oarti	al th	ickn	iess	tears							0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1
Study		TF	FF	F	N T	N Reference sta	ndard	Type of t	tear 9	Sensitivity (95	% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Martin-Hervas 200	01	2	1.3	3.	4 4	O Arthroscopy or open s	uraery (Chronic degenera	ative	0.50 [0.16,	0.841	0.75 [0.62, 0.86]		
Teefey 2004		12	. 1		7 5	1 Arthro	scopy	E	Both	0.63 [0.38,		0.98 [0.90, 1.00]		-
lannotti 2005		27	, 6	11	0 5	6 Arthroscopy or open s	urgery (Chronic degenera	ative	0.73 [0.56,	0.86]	0.90 [0.80, 0.96]	-	-
Lambert 2009		12	1		1 3	4 Arthroscopy or open s	urgery	Unc	lear	0.92 [0.64,	1.00]	0.97 [0.85, 1.00]		-
Della Sala 1996		1	Z	1 1	0 2	5 Open s	urgey	Acute traum	natic	1.00 [0.03,	1.00]	0.86 [0.68, 0.96]		-
Gagey 1993		5	5 1		0 3	2 Open s	urgey (Chronic degenera	ative	1.00 [0.48,	1.00]	0.97 [0.84, 1.00]		
US for detection	of p	artia	l thi	ckne	ess	tears							0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1
Study		TF	FF	F	N	TN Reference sta	andard	Type of	ftear	Sensitivity (95	5% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Martin-Hervas 200	01	1	17	,	7	36 Arthroscopy or open s	surgery	Chronic degener	rative	0.13 [0.00,	, 0.53]	0.68 [0.54, 0.80]	-	-
Sipola 2010		1	9	3	7	60 Arthroscopy or open s	surgery		Both	0.13 [0.00,	0.53]	0.87 [0.77, 0.94]	-	-
Kang 2009		2	. 6	5 :	3	40 Arthr	oscopy	Chronic degener	rative	0.40 [0.05,	, 0.85]	0.89 [0.76, 0.96]		-
Milosavljevic 2005	5	17	, 2	1:	3 1	58 Arthr	оѕсору	Chronic degener	rative	0.57 [0.37,	, 0.75]	0.99 [0.96, 1.00]		•
Teefey 2004		13	3 2	2 1	6	50 Arthr	оѕсору		Both	0.68 [0.43,	, 0.87]	0.96 [0.87, 1.00]		-
lannotti 2005		28	3 7	1	1	55 Arthroscopy or open s	surgery	Chronic degener	rative	0.70 [0.53,	, 0.84]	0.89 [0.78, 0.95]		-
De Candia 2002		8) () :	3	59 l	Inclear	Chronic degener	rative	0.75 [0.43,	, 0.95]	1.00 [0.94, 1.00]		-
Venu 2002		2	2 6	5 1	0	34 Arthr	oscopy	Chronic degener	rative	1.00 [0.16,	, 1.00]	0.87 [0.73, 0.96]	0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1

All six studies (347 shoulders from 346 participants) that assessed the accuracy of MRI for the detection of any rotator cuff tears also assessed partial thickness tears. The median prevalence of partial thickness tears was 20% (range 3% to 37%). Sensitivities ranged from 50% to 100% and specificities ranged from 75% to 98%. The summary sensitivity and specificity of MRI were 74% (95% CI 59% to 85%) and 93% (95% CI 84% to 97%) respectively. The positive and negative likelihood ratios were 10 (95% CI 4 to 26) and 0.28 (95% CI 0.17 to 0.48) respectively.

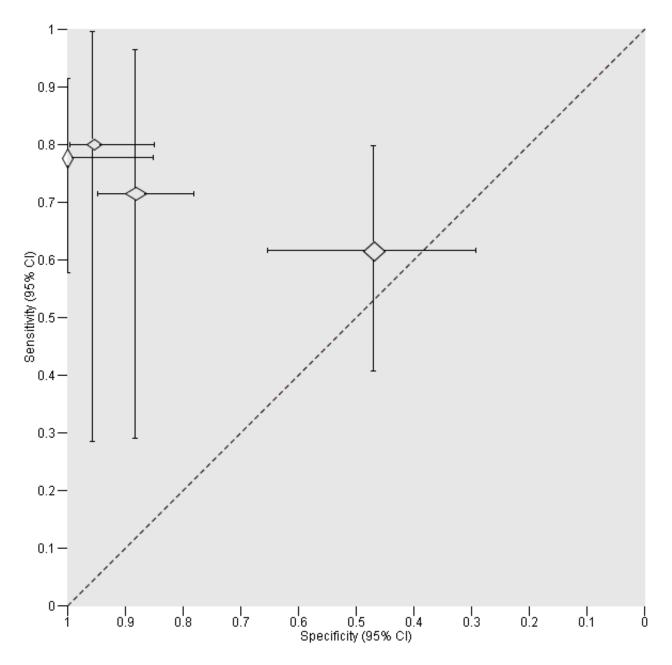
Eight studies (660 shoulders from 654 participants) assessed the accuracy of US to detect partial thickness tears with sensitivities ranging from 13% to 100% and specificities ranging from 68% to

100%. The median prevalence was 14% (range 5% to 37%). The summary sensitivity and specificity of US were 52% (95% CI 33% to 70%) and 93% (95% CI 85% to 97%) respectively. The positive and negative likelihood ratios were 8 (95% CI 3 to 19) and 0.52 (95% CI 0.33 to 0.80) respectively.

Four studies, based on 233 shoulders from 233 participants, assessed the accuracy of MRA to detect partial thickness tears with sensitivities ranging from 62% to 80% and specificities ranging from 47% to 100%. The median prevalence was 27% (range 9% to 54%). Meta-analysis was not performed but study specific estimates of sensitivity and specificity were plotted in ROC space with 95% CI in Figure 10.



Figure 10. Study estimates of sensitivity and specificity with 95% confidence intervals plotted in ROC space for MRA for the detection of partial thickness rotator cuff tears



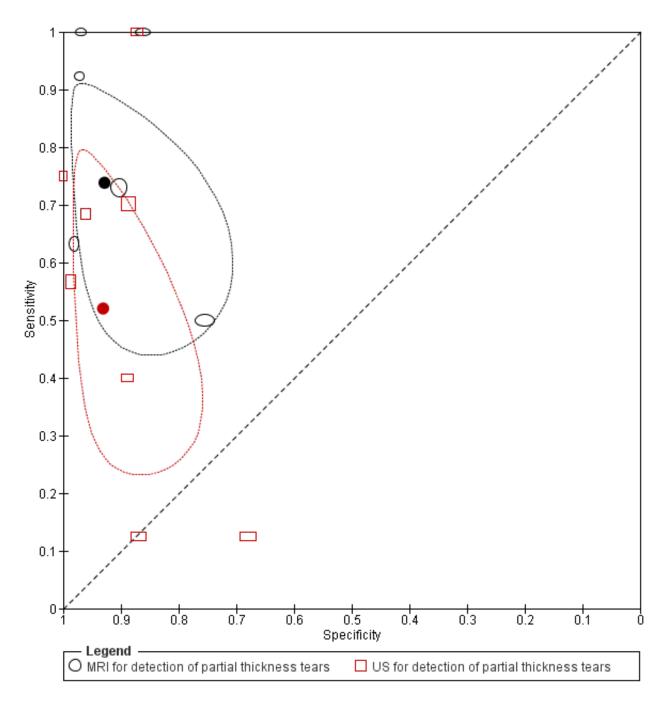
Comparison of MRI and US for detection of partial thickness rotator cuff tears

The diagnostic accuracy of MRI and/or US for detecting partial thickness tears was assessed in 11 studies. There was no statistically significant difference in sensitivity or specificity (P = 1.0) (see Table 1). The individual study estimates of sensitivity and

specificity, with summary points and 95% confidence regions, for each test are shown in ROC space (Figure 11). The sensitivities for MRI and US were generally lower for detecting partial thickness tears than for detecting any or full thickness rotator cuff tears. The sensitivity of US for detecting partial thickness tears was only 52% (95% CI 33% to 70%).



Figure 11. Study estimates of sensitivity and specificity, and summary points with 95% confidence regions plotted in ROC space for MRI and US for the detection of partial thickness rotator cuff tears



The sensitivities and specificities of the three studies that directly compared MRI and US are shown in a ROC space (see Appendix 3 for figure) and differences between the sensitivities and specificities of the tests are presented for each study in Table 2. Two of the studies reported better sensitivity for MRI than US while all three studies reported better specificity for MRI compared to US. Two studies directly compared MRA and US for detection of any rotator cuff tears. Both studies reported better sensitivity and specificity for MRI compared with those of US (Table 3). The same studies also assessed partial thickness tears.

Detection of any subscapularis tendon tears

One study, Mohtadi 2004, assessed the accuracy of MRA for detection of any subscapularis tendon tears, and included 58 shoulders from 58 participants. The study had a prevalence of 33% for subscapularis tendon tears. The sensitivity and specificity of MRA were 79% (95% CI 54% to 94%) and 72% (95% CI 55% to 85%) respectively.



Investigation of heterogeneity

The type of tear and the reference standard used in each study are shown by forest plots for each target condition in Figure 4, Figure 7 and Figure 9. The studies on each plot were ordered according to sensitivity and specificity to demonstrate any pattern in the observed estimates of test accuracy. Based on these descriptive analyses and the magnitude of the variances of the random-effects parameters, we observed greater variability in sensitivity and specificity across studies of US than across studies of MRI or MRA. We were unable to formally investigate potential sources of heterogeneity because the number of studies available for each test was either inadequate or the same value of a covariate was reported by most studies.

Sensitivity analyses

There were few studies of MRI and MRA, and so we could not perform sensitivity analyses for these tests. We performed sensitivity analyses for US for each of the target conditions. We were only able to investigate the impact of two (acceptable reference standard and index test results blinded) of the five quality items we had specified because few studies scored 'Yes' on the other three items (representative spectrum, acceptable delay between tests, and reference test results blinded). There were small differences in sensitivity and/or specificity (Appendix 4). The largest difference was observed between the summary sensitivity of US for detecting partial thickness tears based of all studies (52%, 95% CI 33% to 70%) and the summary sensitivity (62%, 45% to 77%) based on only studies where the reference standard was acceptable. However, the confidence intervals were comparable and the specificities were similar. The exclusion of studies that did not meet either criteria made no difference to our findings. Two studies included both shoulders for six participants and one study did not report the number of participants so it is unclear whether more than one shoulder was included per participant. We investigated the impact of the unit of analysis on the findings for MRI and US by excluding the three studies, thus assuming the individual as the unit of analysis; the results were found to be consistent with the main analyses based on shoulders.

DISCUSSION

Summary of main results

This review summarised the evidence for the diagnostic accuracy of MRI, MRA and US for detecting rotator cuff tears in people with shoulder pain who were suspected of having a rotator cuff tear and for whom surgery was being considered. These imaging tests are usually carried out to determine the characteristics of the rotator cuff tear in order to plan surgery. We included only prospective accuracy studies that evaluated at least one of the tests. We identified 20 studies (1147 shoulders, 1141 participants), of which six evaluated the accuracy of two of the tests within the same participants (paired comparison).

We found no evidence to suggest differences in the sensitivities and specificities of MRI and US for detecting any rotator cuff tears or partial thickness tears. Similarly, we found no evidence to suggest differences in the sensitivities and specificities of MRI, MRA and US for detecting full thickness tears. The estimates were very similar and the tests demonstrated good discriminatory ability for detecting full thickness tears, with sensitivities and specificities of 92% and above. MRI and US had lower sensitivity for partial

thickness tears than for any rotator cuff tears or full thickness tears, with US having a sensitivity of only 52% (95% CI 33% to 70%); this indicates that US may be only marginally better than chance in excluding a partial thickness tear. The specificities of the three tests were generally good except for detection of any rotator cuff tears. The estimates of sensitivity and specificity for any rotator cuff tears suggest that in a population of 100 people with shoulder pain suspected of having a rotator cuff tear and for whom surgery is being considered, if the prevalence was 80%, investigation with MRI may miss two cases (2/80, 3%), while investigation with US may miss seven cases (7/80, 9%). Among patients without a rotator cuff tear (20 out of 100), four patients tested using MRI may have a rotator cuff tear wrongly detected (4/20, 20%) and may undergo unnecessary surgery. A similar number (3/20, 15%) may be overtreated if US is used. The summary of all results are provided in Summary of findings 1.

It is important to emphasise that our review specifically addressed imaging of the rotator cuff by MRI, MRA or US in people with shoulder pain suspected of having a rotator cuff tear and for whom surgery is being considered, and therefore our results are not generalisable to people who present with shoulder pain in primary care where the prevalence of rotator cuff tears may be lower but importantly the prevalence of asymptomatic tears or people with shoulder pain not contemplating surgery could be much higher. Asymptomatic changes in the rotator cuff are common and increase with age and many observed abnormalities might not require specific treatment (Awerbuch 2008). Despite studies continuing to show that primary care practitioners display an over-reliance upon early imaging for shoulder pain (Buchbinder 2013; Johal 2008; Patel 2011), at the present time, guidelines for the management of shoulder pain in primary care do not advocate imaging for shoulder pain unless there is a suggestion of serious pathology (Bussières 2007; Geraets 2009).

The unit of analysis used in evaluating the diagnostic accuracy of a test is likely to have an impact on the estimates of sensitivity and specificity of the test. Our unit of analysis was the shoulder. However, only six out of 1080 participants had both shoulders included in 19 of the 20 included studies; it was unclear in one study (Martin-Hervas 2001) whether the number of shoulders was the same as the number of participants. With the exception of lannotti 2005 and Milosavljevic 2005, the studies reported the same number of participants and shoulders. Both lannotti 2005 and Martin-Hervas 2001 compared the accuracy of MRI and US while Milosavljevic 2005 evaluated only US. In sensitivity analyses, we examined the impact of the unit of analysis by excluding the two studies that included both shoulders for any participant and the one study where it was unclear if the number of shoulders was the same as the number of participants. Overall, findings from the sensitivity analyses were consistent with findings from the main analyses.

Strengths and weaknesses of the review

This review was planned and conducted following criteria and methods set out in a published protocol (Lenza 2011). Our results were based on a comprehensive and sensitive literature search that aimed to identify all published studies. We used wide search terms and several electronic databases, not limited by language, and we excluded search filters for diagnostic terms, as they have limited utility (De Vet 2008). Other strengths of this review are our quality assessment of studies and our synthesis of studies



with similar methodological features into a meta-analytic summary based on recommended methods. To increase the applicability and reliability of the summary findings, we included only prospective studies that investigated people with shoulder pain due to a suspected rotator cuff tear and for whom surgery was being considered. We excluded retrospective studies because of their potential for high risk of spectrum and verification bias (Bossuyt 2003; Van der Schouw 1995).

Our review has some limitations. Our findings were based on small studies with poor reporting of participant characteristics and study design. Most of the QUADAS items were scored as unclear for many studies. For example, only 25% of the included studies reported the time interval between the index tests and the reference standard. For some analyses, we observed considerable heterogeneity in sensitivity and/or specificity, which may be due to several factors including variation in the criteria for a positive diagnostic test for both the index tests and the reference standard, technical details of the tests, variation in population, and variation in operator or reader experience. The three diagnostic tests are known to be operator and reader dependent which may account for some of the observed variation between studies, especially for studies of US which were found to be very heterogeneous. We could not formally investigate potential sources of heterogeneity due to the number of studies available for each test or because most studies reported the same covariate value. Our comparative meta-analyses were based mainly on non-comparative studies because only a small number of studies made direct comparisons between the tests. Consequently, it is possible that observed differences between tests may be confounded by differences in participant and study design characteristics. It is unclear to what extent these limitations influenced our findings.

An important weakness of this review is that due to resource limitations, 25 potentially eligible studies published in non-English languages are still awaiting translation. Good quality translation will be required to reliably extract data from these papers due to the complexity of diagnostic accuracy studies. The studies contain more than 2900 participants that could potentially provide data for analyses and they will be considered for inclusion in a future update of the review.

Comparison with existing reviews

We identified six previous systematic reviews of imaging tests to detect rotator cuff tears (De Jesus 2009; Dinnes 2003; Kelly 2009; Ottenheijm 2010; Shahabpour 2008; Smith 2012). Our review limited inclusion to prospective studies whereas the other systematic reviews allowed the inclusion of retrospective studies. Our literature search failed to identify a study (Ruiz Santiago 2000) which was included in the review by Smith 2012. However, this study would not have been eligible for inclusion in our review because arthrography or arthrographic computed tomography was also used as an index test.

Previous reviews reported similar results. De Jesus 2009 compared US with MRI for detecting rotator cuff tears using surgery as the reference standard. De Jesus 2009 included 65 studies and concluded that US was as accurate as MRI for diagnosing both full and partial thickness rotator cuff tears. Dinnes 2003 assessed the diagnostic accuracy of clinical testing, US and MRI for detecting rotator cuff tears using surgical and non-surgical tests as the reference standard (results also reported in Kelly 2009). Dinnes

2003 concluded that US and MRI were equivalent for detecting full thickness rotator cuff tears, and that MRI may be better at detecting partial thickness tears than US. Shahabpour 2008 also concluded that US and MRI were equivalent for detecting full thickness rotator cuff tears. However, in contrast Shahabpour 2008 concluded that MRA and US may be more accurate at detecting partial thickness tears than MRI. We did not pool MRA studies for detection of partial thickness tears. While our results suggested that MRI may be more sensitive than US, the difference was not statistically significant.

Ottenheijm 2010 assessed the accuracy of US for detecting subacromial diseases in patients presenting in primary and secondary care settings (search date 2001 to June 2010). This systematic review included 23 studies and reported pooled sensitivity and specificity values that were comparable with our results for detecting full thickness tears. Ottenheijm 2010 reported a sensitivity of 95% for detecting full thickness tears compared to 92% (95% CI 82% to 96%) in our systematic review and a specificity of 93% compared with 93% (95% CI 81% to 97%) in our systematic review. However, for detection of partial thickness tears, Ottenheijm 2010 reported a much higher pooled sensitivity of 72% compared with our finding of 52% (95% CI: 33% to 70%). Smith 2012, which included both retrospective and prospective studies, assessed the diagnostic accuracy of MRI and identified 44 studies published up to May 2011. This systematic review reported pooled sensitivity and specificity values that were similar to our results for detecting full thickness tears and partial thickness tears. Smith 2012 reported a pooled sensitivity of 91% (95% CI 86% to 94%) for detecting full thickness tears which was comparable to our result of 94% (95% CI 85% to 98%). Smith 2012 reported a pooled specificity of 97% (95% CI: 96% to 98%) for detecting full thickness tears which is similar to our specificity of 93% (95% CI 83% to 97%). Smith 2012 reported a pooled sensitivity of 80% (95% CI 79% to 84%) for detecting partial thickness tears which is comparable to our sensitivity of 74% (95% CI 59% to 85%); and a pooled specificity of 95% (95% CI 94% to 97%) which is similar to our specificity of 93% (95% CI 84% to 97%). Overall, the results are generally consistent across the different reviews even though there were differences in inclusion criteria and review methods. Despite our study being the most up-to-date published systematic review, we included a much smaller number of studies (20 studies) than some of the previous reviews because we restricted our analyses to only prospective studies thus reducing the risk of spectrum and verification bias.

Applicability of findings to the review question

The applicability of our findings is limited because only 30% of the included studies reported an adequately representative spectrum of consecutive patients from secondary or tertiary care. Furthermore, partial verification was avoided in only 50% of the studies. MRI, MRA and US may have similar accuracy for detecting full thickness rotator cuff tears. The sensitivity of both MRI and US for partial thickness rotator cuff tears appeared to be much lower than their sensitivity for any rotator cuff tears or for full thickness tears. While the difference in sensitivity between MRI and US for detecting partial thickness tears was not statistically significant, US showed a much lower sensitivity (52%) than MRI (74%). A sensitivity of 52% suggests that US may not be any better than chance for detecting partial thickness rotator cuff tears. The specificities of the three tests were generally high except for the detection of any rotator cuff tears.



In many countries, US is less time consuming and less expensive and more readily available in secondary and tertiary care than MRI or MRA. Despite MRI and MRA being comparable for detection of full thickness rotator cuff tears, the choice of test may depend upon cost and availability. As the scope of this review was to limited to test accuracy, we were not able to determine if applying any imaging test prior to surgery results in different surgical interventions or benefits in terms of pain relief and shoulder function following surgery.

AUTHORS' CONCLUSIONS

Implications for practice

The diagnostic performance of MRI and US depends on the extent (i.e. partial or full thickness) of rotator cuff tears. Our findings suggest that MRI, US and MRA have good diagnostic accuracy and any of these tests could equally be used for detection of full thickness tears in people with shoulder pain for whom surgery is being considered. MRI and US also have good sensitivity for detecting any rotator cuff tears but poor sensitivity for detection of partial thickness tears. The validity and generalisability of our findings are limited because they were based on small, heterogeneous, non-comparative studies with methodological flaws.

Implications for research

There is a lack of good quality prospective cohort studies that directly compare the accuracy of MRI, MRA and US shoulder imaging tests for people in secondary and tertiary care, with suspected rotator cuff tears, for whom surgery is being considered. Consequently, further studies are needed in order to evaluate the comparative accuracy of these imaging tests in such circumstances. Future studies should use a blinded design and should limit the amount of time between the index and reference tests as much as possible because there is evidence that rotator cuff tears can progress over time. We suggest that arthroscopy be used as the reference standard test because it is accurate for assessing the articular and bursal side of the rotator cuff. The results of the index test(s) and reference standard should be interpreted by experienced operators.

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CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

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* Indicates the major publication for the study

De Candia 2002

Clinical features and settings

Inclusion criteria: Participants with clinical suspicion of rotator cuff tear who underwent surgery

Exclusion criteria: Not reported

Duration of symptoms: Not reported



De Candia 2002 ((Continued)
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Previous treatments: Not reported

Care setting: Not reported

Participants

Place of study: Udine, Italy

Period of study: January 2000 to December 2000 **Number of participants eligible**: 157 participants

Number of participants enrolled IT and RS:

- US and surgery: 71 participants

Data available for analyses:

- US and surgery: 71 participants

Age (range): 34 to 80 years

Male/Female: 31/40

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To determine the values of the US real time compound imaging in the evaluation of supraspinatus tendon in subacromial impingement disease

Study design: Prospective, accuracy cohort study

Unclear whether consecutive recruitment

Language: English

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Surgery, unclear whether arthroscopy or open surgery

- Description of technique: Not reported

- Criteria for a positive result: Not reported

Index and comparator tests

Index test(s): US

- Description of technique:

Scanner: 7 to 12 MHz linear-array probe applying the soon CT digital algorithm

Technique: Images were obtained in static and dynamic evaluations as described in (Martino 1998; Teefey 2000)

Patient position: Static evaluation was performed on the patient's arm in standard position; dynamic evaluation was performed first with the patient's arm positioned from the internal rotation and extended position to abduction and internal rotation (forearm flexed and the back face of fingertips pointing to the scapula); the second part of the evaluation was performed by moving the patient's arm in adduction and keeping the internal rotation

- Criteria for a positive result: Not reported
- Time from symptoms to index test: Not reported
- <u>Time from index test to reference standard</u>: Index test was performed on the day before reference standard

Follow-up

Adverse events due to index test(s): Not reported



De Candia 2002 (Continued)

Adverse events due to reference standard test(s): Not reported

Notes

Part of population of this study was also reported in De Candia 2003 Although De Candia 2003 is more updated than this study, there were no extra data available to be included in the analyses

The rotator cuff tears were focused on only supraspinatus tendon tears

A two-by-two table of the ITs and RS was given, which tallied with the reported summary data

Item	Authors' judgement	Description
Representative spectrum? All tests	Unclear	Unclear whether consecutive recruitment
Acceptable reference standard? All tests	Unclear	The reference standard was surgery (unclear whether arthroscopy or open surgery) and the target conditions were presence of any rotator cuff tears, full thickness tears and partial thickness tears
Acceptable delay between tests? All tests	Yes	Index test was performed on the day before reference standard
Partial verification avoided?	No	Not all the patients who received the index test underwent a reference standard to verify their true disease status
All tests		Of the 157 eligible participants, only 71 (45.2%) underwent to reference standard
Differential verification avoided? All tests	No	The result of the index test probably influenced the choice of the reference standard
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Reference standard results blinded? All tests	No	The results of the index tests were probably known to the person interpreting the reference tests
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	No	The results of 86 (54.8%) patients were not reported
Withdrawals explained? All tests	No	Some of the eligible patients who entered the study did not complete it and these patients were not accounted for
Learning curve / training reported of index test? All tests	Unclear	The interpreters of index tests were two radiologists; however, the training/expertise was not described



De Candia 2002 (Continued)		
Learning curve / training	Unclear	

reported of reference standard?
All tests

Insufficient information was given to permit judgement

Index test criteria for a positive test result reported?
All tests

No Not reported

Della Sala 1996

Clinical features and settings

Inclusion criteria: Patients with recent trauma without documented articular bone defect radiologically, shoulder pain and disability persisting after appropriate conservative treatment, clinical examination suggestive of rotator cuff tears and/or impingement

Exclusion criteria: Patients with suspected shoulder instability

Duration of symptoms: Not reported **Previous treatments**: Not reported **Care setting**: Tertiary or secondary

Participants Place of study: Trento, Italy

Period of study: January 1993 to December 1994 **Number of participants eligible**: 80 participants

- MRI and open surgery: 30 participants

Number of participants enrolled IT and RS:

Data available for analyses:

- MRI and open surgery: 30 participants

Age: mean 50.1 years (range 21 to 71 years)

Male/Female: 23/7

Dominant arm: Not reported

Nature of onset: Traumatic and chronic injury

Study design **Primary objective**: Not reported

Study design: Unclear whether prospective design. Non-consecutive recruitment

Language: Italian

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Open surgery

- Description of technique:

Open surgery: as described by Neer 1983

- Criteria for a positive result: Not reported



Della Sala 1996 (Continued)

Index and comparator tests

Index test(s): MRI

- Description of technique:

MRI unit: 1.0 T scanner

Sequences and Planes: Spin-echo T1-weighted (TR/TE: 786/17) in coronal and sagittal oblique planes;

and TR/TE 450/12 in axial plane

Patient position: Not reported

- Criteria for a positive result:

Full-thickness tears: an increase signal on the T1-weighted in the entire extension of the rotator cuff

Partial thickness tears: an increased signal in not whole extension of the cuff

- Time from symptoms to index test: Not reported

- <u>Time from index test to reference standard</u>: Not reported

Follow-up Adverse events due to index test(s): Not reported

Adverse events due to reference standard test(s): Not reported

Notes Raw data were given and it was possible to back-calculate this from the reported summary data

Item	Authors' judgement	Description
Representative spectrum? All tests	Unclear	Insufficient information was given to permit judgement
Acceptable reference standard? All tests	Unclear	The reference standard was open surgery and the target conditions were presence of any rotator cuff tears, full thickness tears and partial thickness tears
Acceptable delay between tests? All tests	Unclear	The study did not report the time elapsed between the index tests and reference standard
Partial verification avoided? All tests	No	Not all the patients who received the index test underwent a reference standard to verify their true disease status Of the 80 eligible participants, only 30 received the reference standard
Differential verification avoided?	No	The result of the index test probably influenced the choice of the reference standard
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Reference standard results blinded? All tests	Unclear	Insufficient information was given to permit judgement



Della Sala 1996 (Continued)		
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	No	The results of 50 (62.5%) patients were not reported
Withdrawals explained? All tests	No	Some of the eligible patients who entered the study did not complete it and these patients were not accounted for
Learning curve / training reported of index test? All tests	Unclear	Insufficient information was given to permit judgement
Learning curve / training reported of reference standard? All tests	Unclear	Insufficient information was given to permit judgement
Index test criteria for a positive test result reported? All tests	No	Not reported

Gagey 1993

agey 1000				
Clinical features and set- tings	Inclusion criteria : Patients with a rotator cuff syndrome resistant to any medical treatment and indication for surgery			
	Exclusion criteria: Not reported			
	Duration of symptoms: Not reported			
	Previous treatments: Not reported			
	Care setting: Tertiary			
Participants	Place of study: Paris, France			
	Period of study: 15 months period			
	Number of participants eligible: 38 participants			
	Number of participants enrolled IT and RS:			
	- MRI and open surgery: 38 participants			
	Data available for analyses:			
	- MRI and open surgery: 38 participants			
	Age (mean): 47 years			
	Male/Female: 14/24			
	Dominant arm: Not reported			
	Nature of onset: Not reported			



Gage	y 1993	(Continued)
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Study design **Primary objective**: To compare the results of the MRI with the open surgery

Study design: Prospective accuracy cohort study. Unclear whether consecutive recruitment

Language: French

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Open surgery

- Description of technique: Not reported

- Criteria for a positive result:

Visual identification of the tears by the surgeon

Index and comparator tests

Index test(s): MRI

- Description of technique:

MRI unit: 1.5 T surface circular coil

Sequences: Spin-echo T2-weighted (TR/TE: 2000/25 to 75; TR/TE 1500/25 to 75) and TR/TE 300 to

500/20.

Planes: Sagittal and coronal

Patient position: Not reported

- Criteria for a positive result:

Increased signal on T2-weighted images

- Time from symptoms to index test: Not reported

- Time from index test to reference standard: Not reported

Follow-up

Adverse events due to index test(s): Not reported

Adverse events due to reference standard test(s): Not reported

Notes

Mr Jean-Philippe Regnaux and Mr Ludovic Trinquart kindly translated into English and extracted the

data of this study

The same population of this study was also reported in Gagey 1991

No two-by-two table of the ITs and RS was given, but it was possible to back-calculate this from the re-

ported summary data

Item	Authors' judgement	Description
Representative spectrum? All tests	Unclear	Insufficient information was given to permit judgement
Acceptable reference standard? All tests	Unclear	The reference standard was open surgery and the target conditions were presence of any rotator cuff tears, full thickness tears and partial thickness tears
Acceptable delay between tests?	Unclear	The study did not report the time elapsed between the index tests and reference standard



Gagey 1993 (Continued) All tests		
Partial verification avoided? All tests	Yes	All patients who received the index test went on to receive verification of their disease status using a reference standard
Differential verification avoided? All tests	Yes	All patients received the same reference standard, regardless of the result of their index test
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Reference standard results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Yes	The number of results reported agrees with the number of patients recruited
Withdrawals explained? All tests	Yes	No participants were excluded from the analysis
Learning curve / training reported of index test? All tests	Unclear	Insufficient information was given to permit judgement
Learning curve / training reported of reference stan- dard? All tests	Unclear	Insufficient information was given to permit judgement
Index test criteria for a positive test result report-	Yes	The study reported the definition of a positive index test result

lannotti 2005

ed? All tests

Clinical features and settings

Inclusion criteria: Participants with a clinical diagnosis of rotator cuff symptoms, consisting of pain,

decreased function, and/or weakness

Exclusion criteria: Not reported

Duration of symptoms: Not reported

Previous treatments: Not reported

Care setting: Tertiary and secondary



lannotti 2005 (Continued)

Participants

Place of study: Cleveland, Ohio, USA

Period of study: Not reported

Number of participants eligible: 98 participants (99 shoulders)

Number of participants enrolled IT and RS:

- MRI and arthroscopy or open surgery: 98 participants (99 shoulders)

- US and arthroscopy or open surgery: 98 participants (99 shoulders)

Data available for analyses:

- MRI and arthroscopy or open surgery: 98 participants (99 shoulders)

- US and arthroscopy or open surgery: 98 participants (99 shoulders)

Age: Not reported

Gender: Not reported

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To define the accuracy of US, when performed in an orthopaedic surgeon's office, for the diagnosis of rotator cuff tears

Study design: Prospective, consecutive, accuracy cohort study with fully paired direct comparison between MRI and US

Language: English

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Shoulder arthroscopy or open surgery

- Description of technique(s): Not reported
- Criteria for a positive result:

Full thickness tears: a gap in the tendon; the involved tendons were measured with a calibrated probe or ruler, and the total tendon gap was measured in centimetres prior to débridement of the tendon edges

Partial thickness tears: on either the bursal or the articular surface was identified as tendon-fraying and loss of tendon substance

The size of the partial thickness tears was measured after débridement of the frayed portions of the tendon

Index and comparator tests

Index test(s): MRI and US

- Description of technique:

MRI

MRI unit: 1.5-T magnet

Sequences and Planes: T1 and T2-weighted image sequences in the sagittal and coronal oblique and axial planes

Patient position: Not reported



lannotti 2005 (Continued)

US

Scanner: 7.5 MHz transducer

Technique: Static and dynamic examinations

Images were obtained in transverse plane scans of the biceps tendon; longitudinal and parallel scans of the subscapularis tendon; perpendicular and parallel scans of the supraspinatus and infraspinatus tendons

Patient position: Both the patient and the examiner seated on backless stools facing each other

The patient positioned the arm at the side with the elbow bent to 90°

- Criteria for a positive result:

MRI: Full thickness tears: a fluid filled the gap in the tendon on the T2-weighted sagittal or coronal oblique images

Partial thickness tears: an increase signal on the T1-weighted images, with brighter signal on the T2weighted paired image

The location of the tear was defined by the tendon(s) involved

US: Full thickness tears: a gap in the tendon substance with retraction with increased echogenic signal from the exposed articular cartilage of the humeral head

Partial thickness tears: an increase echogenic signal intensity or a focal decrease in the thickness of the

The location of the tear was defined by the tendon(s) involved

- Time from symptoms to index test: Not reported
- Time from US to MRI: Not reported
- <u>Time from index test to reference standard</u>: Not reported

Follow-up	Adverse events due to index test(s): Not reported
	Adverse events due to reference standard test(s): Not reported
Notes	A two-by-two table of the ITs and RS was given, which tallied with the reported summary data

A two-by-two table of the ITs and RS was given, which tallied with the reported summary data

Item	Authors' judgement	Description
Representative spectrum? All tests	Yes	Tertiary or secondary care, participants with suspected of having any rotator cuff tears
		The study was prospective and recruitment was consecutive
Acceptable reference standard? All tests	Unclear	The reference standard was arthroscopy or open surgery and the target conditions were presence of full thickness tears and partial thickness tears
Acceptable delay between tests? All tests	Unclear	The study did not report the time elapsed between the index tests and reference standard
Partial verification avoided?	Unclear	The study did not report how many patients had US and did not proceed to surgery



lannotti 2005 (Continued) All tests		
Differential verification avoided? All tests	Unclear	Insufficient information was given to permit judgement
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The study was prospective and the results of the index tests were interpreted before the reference standard
Reference standard results blinded? All tests	No	The results of the index tests were known to the person interpreting the reference tests
Relevant clinical information? All tests	Yes	The authors had knowledge of history, physical findings and radiographs
Uninterpretable results reported? All tests	Unclear	Insufficient information was given to permit judgement
Withdrawals explained? All tests	Unclear	Insufficient information was given to permit judgement
Learning curve / training reported of index test? All tests	No	Performace of index tests was not by radiologist or surgeon and they had only two training sections
Learning curve / training reported of reference stan- dard? All tests	Unclear	Insufficient information was given to permit judgement
Index test criteria for a positive test result report- ed? All tests	Yes	The study reported the definition of a positive index test result

Kang 2009

Clinical features and set-	
tings	

Inclusion criteria: Participants with clinical findings of impingement and suspected rotator cuff tear referred for MRA

Exclusion criteria: Participants with previous rotator cuff repair, dislocation, previous humeral fracture, and infectious or inflammatory arthritis were excluded from the study

Participants who showed clinical improvement while scheduled for surgery and refused it

Duration of symptoms: Not reported **Previous treatments**: Not reported

Care setting: Not reported

Participants Place of study: Seoul, Korea



Kang 2009 (Continued)

Period of study: February 2007 to August 2008

Number of participants eligible: 128 participants

Number of participants enrolled IT and RS:

- MRA and arthroscopy: 50 participants

- 3D-US and arthroscopy: 50 participants

Data available for analyses:

- MRA and arthroscopy: 50 participants

- 3D-US and arthroscopy: 50 participants

Age: mean 55.6 years (range 22 to 78 years)

Male/Female: 32/18

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To compare the diagnostic performance of three-dimensional (3D) US and MRA for both the detection of supraspinatus tendon tears and the quantification of their size, with arthroscopic findings used as the standard

Study design: Prospective, consecutive, accuracy cohort study with fully paired direct comparison between MRA and 3D-US

Language: English

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Shoulder arthroscopy

- Description of technique:

In a partial thickness tear or in which the initial arthroscopic findings differed from those reported on the imaging, the rotator cuff was examined from both the articular side and the bursal side

Cuff tear size was measured with a calibrated probe using the posterior portal to measure the anterior–posterior dimension and the lateral portal to measure the medial–lateral dimension

- Criteria for a positive result:

The presence or absence of a full or partial thickness tear was noted at the arthroscopy

An estimate of tear size was performed by the location of the medial edge of the tear

Index and comparator tests

Index test(s): MRA and 3D-US

- Description of technique:

MRA

MRI unit: 3.0 T magnet with a dedicated shoulder coil

Sequences and Planes: Fat-suppressed T1-weighted spin-echo images (TR/TE, 650 to 750/12) in the transverse plane, sagittal oblique plane and coronal oblique plane. T2-weighted turbo spin-echo (TSE) images (4000 to 4500/70) in the sagittal oblique and coronal oblique plane

Contrast and procedure: 12 to 15 mL of diluted gadopentetate dimeglumine with a concentration of 2.0 mmol/L

The procedure involved direct intra-articular injection with fluoroscopic guidance



Kang 2009 (Continued)

Patient position: Supine with the arm in neutral position

3D US

Scanner: 8 to 15 MHz with a dedicated 3D-volume transducer

Technique: Images were obtained in longitudinal scans of supraspinatus tendon 3D-US data were transferred to a separate workstation which was equipped with various post-processing software that allowed display and interactive analysis of the 3D data

In the section mode the volume data were visualised in three orthogonal scan planes, i.e., longitudinal, transverse, and the C-plane (parallel to the surface of the transducer)

Patient position: Patients with the arm in internal rotation, as the patient placed his or her arm on the buttock

- Criteria for a positive result:

MRA: Full-thickness tears: the extension of the contrast medium through the entire thickness of the rotator cuff or presence of the contrast medium in the subacromial–subdeltoid bursa or both

Partial thickness tears: no communication between the glenohumeral joint and the subacromial–sub-deltoid bursa

3D-US: Full thickness tear: a hypoechoic zone extending through the entire substance of the cuff or segmental or complete loss of rotator cuff substance with visualised tear margins or non-visualisation of the cuff

Partial thickness tear: a focal hypoechoic or anechoic defect in the tendon involving either the bursal or the articular surface and manifesting in both longitudinal transverse planes

- Time from symptoms to index test: Not reported
- <u>Time from 3D-US to MRA</u>: The index tests were performed sequentially on the same day beginning with 3D-US and ending with the MRA
- Time from index test to reference standard: mean 24.9 days (range 4 to 99 days

Follow-up	Adverse events due to index test(s): Not reported	
	Adverse events due to reference standard test(s): Not reported	
Notes	The rotator cuff tears were focused on only supraspinatus tendon tears	
	A two-by-two table of the ITs and RS was given, which tallied with the reported summary data	

Item	Authors' judgement	Description
Representative spectrum? All tests	Unclear	Population was patients with suspected of having any rotator cuff tears The study was prospective and recruitment was consecutive
		The care setting was not specified
Acceptable reference standard? All tests	Yes	The reference standard was arthroscopy and the target conditions were presence of full thickness tears and partial thickness tears
Acceptable delay between tests? All tests	Yes	The average interval between reference standard and index test was less than one month



Kang 2009 (Continued)		
Partial verification avoided	No	Not all the patients who received the index test underwent a reference standard to verify their true disease status
All tests		Of the 128 eligible participants only 50 (39%) underwent to reference standard
		16 patients (12.5%) refused surgery as they had improved and it was unclear why the other patients did not receive the reference standard
Differential verification avoided? All tests	No	Probably the result of the index test influenced the choice of the reference standard
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The index tests were interpreted before and without knowledge of the reference standard results
Reference standard results blinded? All tests	No	The results of the index tests were probably known to the person interpreting the reference tests
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	No	The results of 78 (41%) patients were not reported
Withdrawals explained? All tests	No	Some of the eligible patients who entered the study did not complete it and these patients were not accounted for
Learning curve / training reported of index test?	Yes	The interpreter of MRA was a musculoskeletal radiologist with 15 years experience
All tests		The interpreters of US were two radiologists with respectively 5 and 10 years experience performing musculoskeletal US
Learning curve / training reported of reference stan- dard? All tests	Unclear	Insufficient information was given to permit judgement
Index test criteria for a positive test result report- ed? All tests	Yes	The study reported the definition of a positive index test result

Lambert 2009

Clinical features and settings

Inclusion criteria: Not reported

Exclusion criteria: Not reported

Duration of symptoms: Not reported

Previous treatments: Not reported



Lam	bert	2009	(Continued,
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Care setting: Tertiary or secondary

Participants

Place of study: Dijon, France

Period of study: November 2005 to June 2007

Number of participants eligible: 192 participants

Number of participants enrolled IT and RS:

- MRI and arthroscopy or open surgery: 48 participants

Data available for analyses:

- MRI and arthroscopy or open surgery: 48 participants

Age (mean): 56 years
Gender: Not reported

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To demonstrate the value of 3.0 T MRI for the detection of rotator cuff tendon tears

Study design: Prospective, accuracy cohort study Unclear whether consecutive recruitment

Language: English

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Shoulder arthroscopy or open surgery

- <u>Description of technique</u>: Not reported

- Criteria for a positive result: Not reported

Index and comparator tests

Index test(s): MRI

- Description of technique:

MRI unit: 3.0 T scanner with a shoulder coil

Sequences and Planes: Fat suppressed TSE T2-weighted in three planes (TR/TE 3000/39)

A sagittal T1-weighted sequence to detect fatty muscle atrophy

Patient position: Not reported

- Criteria for a positive result:

Full thickness tears: presence of hyperintense fluid signal with a communication between the gleno-humeral joint and subacromial space

Partial thickness tears: hyperintense fluid signal or irregularity at the articular or bursal surface of the tendon

- Time from symptoms to index test: Not reported

- Time from index test to reference standard: mean 77.6 days (range 22 to 169 days)

Follow-up

Adverse events due to index test(s): Not reported



Lambert 2009 (Continued)

Adverse events due to reference standard test(s): Not reported

Notes A two-by-two table of the ITs and RS was given, which tallied with the reported summary data

Item	Authors' judgement	Description
Representative spectrum? All tests	No	Unknown why patients had MRI shoulder as included all people who had one at their institution of a period of time November 2005 to June 2007
Acceptable reference standard? All tests	Unclear	The reference standard was shoulder arthroscopy or open surgery and the target condition were presence of any rotator cuff tears, full thickness tears and partial thickness tears
Acceptable delay between tests? All tests	No	The average interval between reference standard and index test was 77.6 days
Partial verification avoided? All tests	No	Not all the patients who received the index test underwent a reference standard to verify their true disease status. Of the 192 eligible participants, only 48 underwent to reference standard
Differential verification avoided? All tests	Unclear	Insufficient information was given to permit judgement
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Reference standard results blinded? All tests	No	The results of the index tests were known to the person interpreting the reference tests
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Unclear	Insufficient information was given to permit judgement
Withdrawals explained? All tests	No	Some of the eligible patients who entered the study did not complete it and these patients were not accounted for
Learning curve / training reported of index test? All tests	Unclear	Insufficient information was given to permit judgement
Learning curve / training reported of reference stan- dard? All tests	Unclear	Insufficient information was given to permit judgement



Lambert 2009 (Continued)

Index test criteria for a positive test result reported?
All tests

Yes

The study reported the definition of a positive index test result

Martin-Hervas 2001

Clinical features and settings

Inclusion criteria: Patients with shoulder pain and limited movement

Exclusion criteria: Patients with claustrophobia, metallic implants, and pacemaker

Duration of symptoms: Not reported **Previous treatments**: Not reported

Care setting: Not reported

Participants

Place of study: Madrid, Spain

Period of study: During 1998

Number of participants eligible: 140 shoulders

Number of participants enrolled IT and RS:

- MRI and arthroscopy or open surgery: 61 shoulders

- US and arthroscopy or open surgery: 72 shoulders

Data available for analyses:

- MRI and arthroscopy or open surgery: 61 shoulders

- US and arthroscopy or open surgery: 61 shoulders

Age: Not reported

Male/Female: 25/36

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To compare the accuracy of US and MRI in the diagnosis of rotator cuff injuries (focusing on supraspinatus tears) using arthroscopy or open surgery findings as the gold standard

Study design: Prospective accuracy cohort study with fully paired direct comparison between US and

MRI

Unclear whether consecutive recruitment

Language: English

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Shoulder arthroscopy or open surgery

- Description of technique: Not reported

- Criteria for a positive result: Not reported



Martin-Hervas 2001 (Continued)

Index and comparator tests

Index test(s): MRI and US

- Description of technique:

MRI

MRI unit: 0.5 T superconducting magnet

Sequences: Spin echo T1-weighted sequences for coronal and oblique plane images and gradient echo T2*-weighted sequences for axial and oblique coronal images; when the supraspinatus tendon showed a suggestive increased signal intensity, spin echo T2-weighted sequences were performed

Planes: Axial, oblique coronal and oblique sagittal images

Patient position: Patient in a supine position and the arm in a neutral position

US

Scanner: 7.5 MHz high-resolution linear electronic transducer

Technique and Patient position: Images were obtained in transverse and longitudinal plane scans on the anterior plane of a shoulder with a neutrally rotated humerus to visualise bicipital and subscapularis bursae and axilla

Next, sections of the shoulder were performed with internal humeral rotation, and the transducer was moved laterally to visualise the supraspinatus tendon and subacromial bursa

The last images were obtained in the posterior plane with the humerus in a neutral position to visualise the infraspinatus and teres minor tendons

- Criteria for a positive result:

MRI: Full thickness tears: hypersignal on the T1- and T2-weighted images or any irregularity in the borders of the entire thickness of the tendon

Partial thickness tears: any irregularity within the tendon or at the bursal or joint surfaces

US: Full thickness tears: complete absence of the tendon, focal atrophy, a concave border, liquid-filled hypoechoic bands, and/or lineal hyperechoic bands

Partial thickness tears: heterogeneous tendon with hypoechoic areas (> 3 mm) that do not reach both sides of the tear and an irregular or indented border

- Time from symptoms to index test: Not reported
- Time from MRI and US: Not reported
- Time from index test to reference standard: Less than 6 months

Follow-up	Adverse events due to index test(s): Not reported	
	Adverse events due to reference standard test(s): Not reported	
Notes	The rotator cuff tears were focused on only supraspinatus tendon tears	
	No two-by-two table of the ITs and RS was given, but it was possible to back-calculate this from the reported summary data	

Item	Authors' judgement	Description
Representative spectrum? All tests	Unclear	Tertiary or secondary care, participants with suspected of having any rotator cuff tears



Martin-Hervas 2001 (Continued)	
		The study was prospective
		It was unclear whether consecutive recruitment
Acceptable reference standard? All tests	Unclear	The reference standard was arthroscopy or open surgery and the target conditions were presence of any rotator cuff tears, full thickness tears and partial thickness tears
Acceptable delay between tests? All tests	Unclear	The interval between tests was not clearly reported
Partial verification avoided? All tests	No	Not all the patients who received the index test underwent a reference standard to verify their true disease status
Differential verification avoided? All tests	Unclear	Insufficient information was given to permit judgement
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The study was prospective and the results of the index tests were interpreted before the reference standard
Reference standard results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Unclear	Insufficient information was given to permit judgement
Withdrawals explained? All tests	No	Some of the eligible patients who entered the study did not complete it and these patients were not accounted for
Learning curve / training reported of index test? All tests	Unclear	The interpreter of index test was a musculoskeletal radiologist Experience was not reported
Learning curve / training reported of reference stan- dard? All tests	Unclear	Insufficient information was given to permit judgement
Index test criteria for a positive test result report- ed? All tests	Yes	The study reported the definition of a positive index test result



Milosavljevic 2005

Clinical features and settings

Inclusion criteria: Participants with shoulder symptoms longer than 3 months duration and clinical findings of impingement and suspected rotator cuff tears were referred for US of the shoulder

The patients had pain at rest and during motion, the pain-provoking test was positive, and some patients had weakness of the rotator cuff muscles

Exclusion criteria: Not reported

Duration of symptoms: Participants with shoulder symptoms longer than 3 months duration

Previous treatments: Not reported

Care setting: Not reported

Participants

Place of study: Uppsala, Sweden

Period of study: February 1999 to October 2002

Number of participants eligible: 185 participants (190 shoulders)

Number of participants enrolled IT and RS:

- US and arthroscopy: 185 participants (190 shoulders)

Data available for analyses:

- US and arthroscopy: 185 participants (190 shoulders)

Age: mean 57 years (range 22 to 78 years)

Male/Female: 114/71

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To evaluate the accuracy of high-resolution shoulder US compared with arthroscopy in a large group of consecutive patients with clinically suspected rotator cuff disease

Study design: Prospective consecutive accuracy cohort study

Language: English

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Shoulder arthroscopy

- Description of technique:

Patient in the beach-chair position under general anaesthesia

Posterior and anterior portal were used

The cartilage of the humeral head and the glenoid fossa, the labrum ligament complex, the biceps tendon, the intraarticular portion of the subscapular tendon, and the underside of the rotator cuff were inspected

- Criteria for a positive result:

In the same manner as for the US findings, i.e. intact cuff, full thickness tears, or partial thickness tears (see below)

Index and comparator tests

Index test(s): US

- Description of technique:



Milosavljevic 2005 (Continued)

Scanner: 10 MHz linear-array transducer

Technique: All tendons were examined in longitudinal and transversal plane

Patient position: Both patient and examiner seated on rotatable chairs without armrests

The examiner faced the patient and was seated at the patient's right side

- Criteria for a positive result:

Full thickness tears: defect (hypoechoic zone) extending through the entire substance of the cuff; focal, mixed hyper- and hypoechoic lesion extending through the entire substance of the cuff; focal thinning with visible margins of the tear; and non-visualisation of the cuff

Partial thickness tears: mixed hyper- and hypoechoic focus or a hypoechoic lesion visualised in two orthogonal imaging planes located within the tendon substance but not extending to the surface or with either articular or bursal extension

- Time from symptoms to index test: More than 3 months
- Time from index test to reference standard: mean 6 months (range 1 day to 18 months)

Follow-up	Adverse events due to index test(s): Not reported
	Adverse events due to reference standard test(s): Not reported
Notes	A two-by-two table of the ITs and RS was given, which tallied with the reported summary data

Item	Authors' judgement	Description
Representative spectrum? All tests	Unclear	Population was patients with suspected of having any rotator cuff tears The study was prospective and recruitment was consecutive
		The care setting was not specified
Acceptable reference standard? All tests	Yes	The reference standard was arthroscopy and the target conditions were presence of any rotator cuff tear, full thickness tears and partial thickness tears
Acceptable delay between tests? All tests	No	The average interval between reference standard and index test was 6 months
Partial verification avoided? All tests	Yes	All patients who received the index test went on to receive verification of their disease status using a reference standard
Differential verification avoided? All tests	Yes	All patients received the same reference standard regardless of the result of their index test
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	No	When there was disagreement between index test and reference standard findings the results of index test (US) were re-evaluated to explain discrepancy
Reference standard results blinded?	Unclear	Insufficient information was given to permit judgement



Milosavl	jevic 2005	(Continued)
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М	u		U		-	LZ

Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Yes	The study was prospective, recruitment was consecutive and results were reported for all initially included participants
Withdrawals explained? All tests	Yes	No participants were excluded from the analysis
Learning curve / training reported of index test? All tests	Unclear	The interpreter of index tests was a radiologist Training and expertise were not described
Learning curve / training reported of reference standard? All tests	Unclear	The reference standards were performed by three shoulder surgeons Training and expertise were not described
Index test criteria for a positive test result report- ed? All tests	Yes	The study reported the definition of a positive index test result

Misamore 1991

Clinical features and settings

Inclusion criteria: Participants with symptoms and physical findings consistent with chronic tendinitis or with degeneration or a tear of the rotator cuff

Patients who had signs indicating stage II or stage III impingement were included

Exclusion criteria: Patients who had an acute injury or who had symptoms for less than one year were excluded

Patients were excluded if they had a previous operation on the affected shoulder, if they had any associated disorders of the shoulder (such as arthritis or instability), or if they had cervical radiculopathy or peripheral neuropathy

Duration of symptoms: Not reported **Previous treatments**: Not reported

Care setting: Not reported

Participants

Place of study: Indianapolis, Indiana, USA

Period of study: January 1988 to June 1989

Number of participants eligible: 82 participants

Number of participants enrolled IT and RS:

- US and arthroscopy or open surgery: 32 participants

Data available for analyses:

- US and arthroscopy or open surgery: 32 participants



Age: mean 47 years (range 35 to 65 years)

Male/Female: 26/6

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To compare the accuracy of arthrography compared with US in the evaluation of thirty-two patients who had a degenerative lesion of the rotator cuff.

Study design: Prospective consecutive accuracy cohort study

Language: English

Target condition and reference standard(s)

Target conditions: Presence of full thickness tears and partial thickness rotator cuff tears

Reference standard(s): Shoulder arthroscopy or open surgery

- <u>Description of technique</u>: Not reported
 <u>Criteria for a positive result</u>: Not reported
- Index and comparator tests

Index test(s): US

- Description of technique: Not reported
- Criteria for a positive result:

Full thickness tear: an obvious defect localised to the tendon of the rotator cuff was seen or alternatively when there was no echo of the rotator cuff

An abnormality of echogenicity alone was not considered to be a tear

- <u>Time from symptoms to index test</u>: Not reported
- Time from index test to reference standard: Not reported

Follow-up

Adverse events due to index test(s): Not reported

Adverse events due to reference standard test(s): Not reported

Notes

No two-by-two table of the ITs and RS was given, but it was possible to back-calculate this from the reported summary data for detecting any rotator cuff tears

Item	Authors' judgement	Description
Representative spectrum? All tests	Unclear	Population was patients with suspected of having any rotator cuff tears The study was prospective and recruitment was consecutive The care setting was not specified
Acceptable reference standard? All tests	Unclear	The reference standard was arthroscopy or open surgery and the target conditions were presence of full and partial thickness tears
Acceptable delay between tests? All tests	Unclear	The study did not report the time elapsed between the index tests and reference standard



Misamore 1991 (Continued)		
Partial verification avoided?	No	Not all the patients who received the index test underwent a reference standard to verify their true disease status
All tests		Of the 82 eligible participants, 32 patients received the reference standard
		For 50 patients the symptoms were not severe enough to justify surgery or satisfactory improvement was achieved with conservative treatment
Differential verification avoided? All tests	No	Probably the result of the index test influenced the choice of the reference standard
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The index tests were interpreted before and without knowledge of the reference standard results
Reference standard results blinded? All tests	No	The results of index tests were probably known to the person interpreting the reference standard
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Yes	Results were reported for all initially included participants
Withdrawals explained? All tests	Yes	The number and reasons of all withdrawals from the study were explained
Learning curve / training reported of index test? All tests	Yes	The interpreters of index tests were radiologists who were skilled in the technique
Learning curve / training reported of reference stan- dard? All tests	Unclear	The reference standards were performed by one of the authors (orthopaedic surgeon)
Index test criteria for a positive test result report- ed? All tests	Yes	The study reported the definition of a positive index test result

Mohtadi 2004

Clinical features and settings

Inclusion criteria: Patients with shoulder pain at night or with overhead activity greater than 3 months duration or both

A minimum of 3 of the following 6 clinical findings: a painful arc of motion in the scapular plane (60° to 120°) of elevation; pain-related weakness on resisted elevation in the scapular plane; Neer's impingement sign; Hawkin's impingement sign; point of maximal tenderness over the supraspinatus tendon; and positive impingement xylocaine test



Mohtadi 2004 (Continued)

Failure of conservative management

The patients consented to undergo shoulder arthroscopy and subacromial decompression

Exclusion criteria: Patients with symptoms of instability

Signs of instability

Point of maximum tenderness over the acromioclavicular joint

Any signs or symptoms consistent with associated cervical spine pathology Previous surgery, arthrog-

raphy, ultrasound, or MRI

Duration of symptoms: More than 3 months of symptoms

Previous treatments: Conservative management (nonsteroidal anti-inflammatory drugs, physiothera-

py, home-based rehabilitation, cortisone injections, and modification of activity)

Care setting: Tertiary or secondary

Participants

Place of study: Calgary, Alberta, Canada

Period of study: 1998 to 2000

Number of participants eligible: 73 participants

Number of participants enrolled IT and RS:

- Indirect MRA and arthroscopy: 58 participants

Data available for analyses:

- Indirect MRA and arthroscopy: 58 participants

Age: mean 46.2 years (range 21 to 73 years)

Male/Female: 43/15

Dominant arm: Not reported

Nature of onset: Of these 58 patients, 91.4% reported pain at night and 96.6% reported pain with activ-

ity above shoulder level

Study design

Primary objective: To determine the diagnostic ability of MRI compared with a reference standard, arthroscopy, in patients presenting with shoulder pain consistent with the signs and symptoms of

shoulder impingement

Study design: Prospective, consecutive accuracy cohort study

Language: English

Target condition and reference standard(s)

Target conditions: Presence full thickness tears and partial thickness supraspinatus tendon tears

Presence of any infraspinatus tendon tears

Presence of any subscapularis tendon tears

Reference standard(s): Shoulder arthroscopy

- Description of technique:

In accordance with the standardised 15-point protocol of Snyder classification This included standard posterior and anterior portal examination with subsequent visualisation in the subacromial bursa

The subacromial (bursal) examination was not performed



Mohtad	i 2004	(Continued)
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All surgeries were videotaped

- Criteria for a positive result: Not reported

Index and comparator tests

Index test(s): Indirect MRA

- <u>Description of technique</u>:

MRI unit: 1.5 T with conventional shoulder coil

Sequences and Planes: Axial water density (TR/TR 1000/20) and multi-planar gradient recalled (TR/TE 400/20, flip angle20°)

Oblique coronal fast multi-planar inversion recovery (TR/TE 4600/28, inversion time 150)

Oblique coronal post-gadolinium fat-saturated T1-weighted (TR/TE 400/8) and sagittal T1-weighted (TR/TE 400/8)

Contrast and procedure: Intravenous gadolinium administration

Patient position: Supine with the arm in a neutral position

- Criteria for a positive result: Not reported
- Time from symptoms to index test: More than 3 months of symptoms
- <u>Time from index test to reference standard</u>: Upon entry into the study patients were scheduled to undergo MRI within 1 week before arthroscopy

Follow-up

Adverse events due to index test(s): Not reported

Adverse events due to reference standard test(s): Not reported

Notes

The analyses of rotator cuff tears were focused on only supraspinatus and subscapularis tendons tears

No two-by-two table of the ITs and RS was given, but it was possible to back-calculate this from the reported summary data

Item	Authors' judgement	Description
Representative spectrum? All tests	Yes	Tertiary or secondary care, participants with suspected of having any rotator cuff tears
		The study was prospective and recruitment was consecutive
Acceptable reference standard? All tests	Yes	The reference standard was arthroscopy and the target conditions were presence full thickness tears and partial thickness supraspinatus tendon tears; presence of any infraspinatus tendon tears; presence of any subscapularis tendon tears
Acceptable delay between tests? All tests	Yes	Patients were scheduled to undergo MRI within 1 week before arthroscopy
Partial verification avoided? All tests	No	Not all the patients who received the index test underwent a reference standard to verify their true disease status
		Of the 73 eligible participants, eight cancelled the surgery and seven did not undergo MRI within a week and were excluded but their results were not reported



Mohtadi 2004 (Continued)		
Differential verification avoided? All tests	Yes	All patients received the same reference standard regardless of the result of their index test
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The index tests were interpreted before and without knowledge of the reference standard results
Reference standard results blinded? All tests	Yes	Before and during diagnostic arthroscopy, the surgeon was blinded to the MRI results
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Yes	The study was prospective recruitment was consecutive and results were reported for all initially included participants
Withdrawals explained? All tests	Yes	The number and reasons of all withdrawals from the study were explained
Learning curve / training reported of index test? All tests	Unclear	The interpreter of index tests was a musculoskeletal radiologist Training and expertise were not described
Learning curve / training reported of reference stan-	Unclear	The reference standards were performed by two experienced orthopaedic surgeons
dard? All tests		Training and expertise were not described
Index test criteria for a positive test result report- ed? All tests	No	Not reported

Nicoletti 1994

Clinical features and set- tings	Inclusion criteria : Patients who had shoulder pain and signs and symptoms of rotator cuff tears with failure of conservative treatment	
	Exclusion criteria: Patients with suspected of instability and neurologic symptoms	
	Duration of symptoms : More than 3 months	
	Previous treatments: Physiotherapy	
	Care setting: Tertiary	
Participants	Place of study: Sao Paulo, Brazil	
	Period of study: Not reported	
	Number of participants eligible: 48 participants	



Nicoletti 1994	(Continued)
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Number of participants enrolled IT and RS:

- US and arthroscopy: 48 participants

Data available for analyses:

- US and arthroscopy: 48 participants

Age: mean 48 years (range 19 to 79 years)

Male/Female: 16/32

Dominant arm: 65%

Nature of onset: Not reported

Study design

Primary objective: To evaluate the sensitivity, specificity and accuracy of arthrography and US to de-

tect rotator cuff tears

Study design: Unclear whether prospective design

Unclear whether consecutive recruitment

Language: Portuguese

Target condition and reference standard(s)

Target conditions: Presence any rotator cuff tears

Reference standard(s): Shoulder arthroscopy

- Description of technique:

Patient in lateral position with traction in the operative limb

The posterior and anterior portals were used to visualise the glenohumeral and subacromial spaces

- Criteria for a positive result: Not reported

Index and comparator tests

Index test(s): US

- Description of technique:

Scanner: 5 or 7 MHz linear transducer in real time

Technique and Patient position: As described by Crass 1985

- Criteria for a positive result:

US signs were: focal or diffuse thinning or non-visualisation of tendon(s)

- Time from symptoms to index test: Not reported
- Time from index test to reference standard: Not reported

Follow-up

Adverse events due to index test(s): Not reported

Adverse events due to reference standard test(s): Not reported

Notes

A two-by-two table of the ITs and RS was given which tallied with the reported summary data

Item	Authors' judgement	Description
Representative spectrum?	Unclear	Unclear whether prospective design and consecutive recruitment



Nicoletti 1994 (Continued) All tests		
Acceptable reference standard? All tests	Yes	The reference standard was arthroscopy and the target condition was presence of any rotator cuff tears
Acceptable delay between tests? All tests	Unclear	The study did not report the time elapsed between the index tests and reference standard
Partial verification avoided? All tests	Yes	All patients who received the index test went on to receive verification of their disease status using a reference standard
Differential verification avoided? All tests	Unclear	Insufficient information was given to permit judgement
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Reference standard results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Yes	The number of results reported agrees with the number of patients recruited
Withdrawals explained? All tests	Yes	No participants were excluded from the analysis
Learning curve / training reported of index test? All tests	Yes	The interpreter of index tests was a musculoskeletal radiologist
Learning curve / training reported of reference stan- dard? All tests	Yes	The reference standards were performed by an experienced shoulder surgeon
Index test criteria for a positive test result reported?	Yes	The study reported the definition of a positive index test result

Sipola 2010

All tests

Clinical features and settings Inclusion criteria: Participants with acute or chronic shoulder pain and suspicion of rotator cuff tears



Sipola 2010 (Continued)

Patients who had undergone conservative treatment without sufficient symptom relief

Exclusion criteria: Time elapsed between index test and reference standard was more than 12 months

Duration of symptoms (pain): mean 21 months (range 2 to 144 months

Previous treatments: Conservative treatment including physiotherapy for at least 3 months

Care setting: Tertiary or secondary

Participants

Place of study: Kuopio, Finland

Period of study: Not reported

Number of participants eligible: 79 participants

Number of participants enrolled IT and RS:

- MRA and arthroscopy or open surgery: 75 participants

- US and arthroscopy or open surgery: 77 participants

Data available for analyses:

- MRA and arthroscopy or open surgery: 75 participants

- US and arthroscopy or open surgery: 77 participants

Age: mean 57 years (range 42 to 76 years)

Male/Female: 40/37

Dominant arm: Not reported

Nature of onset: The etiology of suspected tear was traumatic in 22% and degenerative in 78% of the

participants

Study design

Primary objective: To compare the accuracy of US and MRA for the detection and measurement of ro-

tator cuff tears using surgical findings as the standard in a prospective study setting

Study design: Prospective, consecutive, accuracy cohort study with fully paired direct comparison be-

tween MRA and US

Language: English

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Shoulder arthroscopy or mini-open

- Description of technique: Not reported

- Criteria for a positive result:

The size and type (partial/full thickness) of tendon tear was determined and measured from anterior to posterior and from lateral to medial dimensions A sterile ruler or a calibrated arthroscopic probe was

used to define both the anteroposterior and the mediolateral size of the tear

Index and comparator tests

Index test(s): MRA and US

- Description of technique:

MRA

MRI unit: 1.5 T scanner equipped with a flexible surface coil



Sipola 2010 (Continued)

Sequences and Planes: Oblique coronal T1-weighted spin-echo (TR/TE 650/20); a T2-weighted fat-saturated dual-echo fast spin-echo (FSE), 3500/16; oblique sagittal T2-weighted dual-echo FSE, 3500/16, 98; axial T2*-weighted gradient echo two-dimensional FLASH, 580/15, flip angle 15°; a T1-weighted fat-saturated SE, 800/20; T1-weighted fat-saturated spin-echo 800/20 images in the sagittal oblique, coronal oblique and axial planes, and T2-weighted FSE 4500/96 images in the coronal oblique plane

Contrast and procedure: 10 to 20 mL of gadopentetate dimeglumine in a concentration of 469.01 mg/ mL was diluted in 250 mL of saline

The procedure involved direct intra-articular injection

Patient position: Not reported

Seven participants underwent to MRI only (without an intra-articular contrast)

US

Scanner: 7.5 MHz linear-array transducers in real-time

Technique: Images were obtained on the long and short axes of the tendon

Patient position: The subscapularis tendon was evaluated with the forearm rotated externally

The supraspinatus tendon was assessed with the arm on the ipsilateral side The supraspinatus was assessed with the hand behind the patient's back (Crass position) or on the waist (modified Crass position) (Crass 1987; Ferri 2005)

The infraspinatus tendon was assessed with the patient placed the ipsilateral hand across the chest on top of the contralateral shoulder

- Criteria for a positive result:

MRA: Full thickness tears: the contrast agent was detected on the MR image throughout the full thickness of the rotator cuff and/or when the contrast agent was detected in the subacromial bursa

Partial thickness tears: the contrast agent entered the cuff substance without reaching the subacromial bursa

US: Full thickness tears: hypoechoic area or volume loss extended from the bursal surface to the articular surface of the tendon

Otherwise the tear was diagnosed as a partial thickness tear

- Time from symptoms to index test: mean 21 months (range 2 to 144 months
- Time from MRA and US: in the same day
- Time from index test to reference standard: mean 2.3 months (range 0 to 9.5 months)

Follow-up

Adverse events due to index test(s): Of the 77 patients, two (3%) could not undergo MRA due to claustrophobia

Adverse events due to reference standard test(s): Not reported

Notes

A two-by-two table of the ITs and RS was given, which tallied with the reported summary data

Item	Authors' judgement	Description
Representative spectrum? All tests	Yes	Tertiary or secondary care, participants with suspected of having any rotator cuff tears
		The study was prospective and recruitment was consecutive



Sipola 2010 (Continued)		
Acceptable reference standard? All tests	Unclear	The reference standard was arthroscopy or open surgery and the target conditions were presence of any rotator cuff tears, full thickness tears and partial thickness tears
Acceptable delay between tests? All tests	No	The average interval between reference standard and index test was 2.3 months
Partial verification avoided? All tests	Yes	Only two patients (2.5%) were excluded of study because of delay in surgery 12 months due to medical illness
Differential verification avoided? All tests	No	The choice of reference standard (arthroscopy or open surgery) varied between individuals
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Reference standard results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Yes	The study was prospective, recruitment was consecutive and results were reported for all initially included participants
Withdrawals explained? All tests	Yes	The number and reasons of all withdrawals from the study were explained
Learning curve / training reported of index test?	Yes	The interpreter of MRA was a radiologist who had 1 year of experience in musculoskeletal MRI at the beginning of the study
All tests		The interpreters of US were three radiologists each with more than 10 years experience in shoulder US
Learning curve / training reported of reference standard?	Unclear	The reference standards were performed by three experienced orthopaedic surgeons
dard? All tests		Training and expertise were not described
Index test criteria for a positive test result reported? All tests	Yes	The study reported the definition of a positive index test result



Stetson 2005

Clinical features and settings

Inclusion criteria: Patients with chronic shoulder pain who were suspected of having a rotator cuff ab-

normality underwent MRA with use of an intra-articular injection of gadolinium

Exclusion criteria: Not reported

Duration of symptoms: Not reported **Previous treatments**: Not reported

Care setting: Tertiary or secondary

Participants

Place of study: Burbank, California, USA

Period of study: During 2 years

Number of participants eligible: 50 participants

Number of participants enrolled IT and RS:

- MRA and arthroscopy: 50 participants

Data available for analyses:

- MRA and arthroscopy: 50 participants

Age: Not reported

Gender: Not reported

Dominant arm: Not reported

Nature of onset: Not reported

Study design

Primary objective: To detect partial thickness articular-sided rotator cuff tears using an intra-articular

injection of gadolinium and MRI

Study design: Prospective accuracy cohort

Unclear whether consecutive recruitment

Language: English

Target condition and reference standard(s)

Target conditions: Presence partial articular-side thickness tears

Reference standard(s): Shoulder arthroscopy

- Description of technique:

All participants were taken to surgery and underwent a complete 15-point glenohumeral arthroscopic

examination

The presence or absence of articular-sided rotator cuff tears was recorded

- Criteria for a positive result: Not reported

Index and comparator tests

Index test(s): MRA

- Description of technique:

MRI unit: 1.5 T scanner.

Sequences and Planes: Axial proton-density-weighted image with fat suppression, oblique coronal proton-density-weighted image, oblique coronal T2-weighted with fat suppression, oblique sagittal T1-weighted, and oblique sagittal proton-density-weighted image with fat suppression. In addition, axial

T1-weighted with fat suppression, oblique coronal T1-weighted with fat suppression



Stetson 2005 (Continued)

Contrast and procedure: 1.5 mL of gadolinium with normal saline solution intra-articularly into the glenohumeral joint under fluoroscopic control

Patient position: Supine in neutral position and abduction and external rotation images were also acquired

- Criteria for a positive result: Not reported
- Time from symptoms to index test: Not reported
- Time from index test to reference standard: Not reported

Follow-up Adverse events due to index test(s): Not reported Adverse events due to reference standard test(s): Not reported Notes The authors described as false positive the four patients who were incorrectly diagnosed with MRA, as having a full thickness tear, but, at the time of shoulder arthroscopy, they had partial thickness articular-sided tears

To make concordance with our analyses we described these participants as false negative

No two-by-two table of the ITs and RS was given, but it was possible to back-calculate this from the reported summary data

Item	Authors' judgement	Description
Representative spectrum? All tests	Unclear	The study was prospective and the population was participants with shoulder pain and suspected of having any rotator cuff tears. However, it was unclear whether there was consecutive recruitment
Acceptable reference standard? All tests	Yes	The reference standard was arthroscopy and the target condition was presence of partial articular-side rotator cuff tears
Acceptable delay between tests? All tests	Unclear	The study did not report the time elapsed between the index tests and reference standard
Partial verification avoided? All tests	Yes	All patients who received the index test went on to receive verification of their disease status using a reference standard
Differential verification avoided? All tests	Yes	All patients received the same reference standard, regardless of the result of their index test
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Reference standard results blinded? All tests	No	The results of the index tests were probably known to the person interpreting the reference tests
Relevant clinical information?	Unclear	Insufficient information was given to permit judgement



Stetson 2005 (Continued) All tests		
Uninterpretable results reported? All tests	Yes	The number of results reported agrees with the number of patients recruited
Withdrawals explained? All tests	Yes	No participants were excluded from the analysis
Learning curve / training reported of index test? All tests	Yes	The interpreter of index tests was a fellowship-trained musculoskeletal radiologist
Learning curve / training reported of reference stan- dard? All tests	Unclear	Insufficient information was given to permit judgement
Index test criteria for a positive test result report- ed? All tests	No	Not reported

Swen 1998

tings	inclusion criteria: Patients awaiting surgery because of clinically suspected of rotator cuπ tears		
ungs	The clinical diagnosis of rotator cuff tears was based on marked difficulty in initiating abduction of the arm with weakness and limitation of movement Lidocaine was injected below the acromion, if after the injection the strength of the rotator cuff was still decreased; this was considered to indicate rotator cuff tears		
	Exclusion criteria: Patients with neurologic origins of the weakness		

Execusion effection. Facilities with hear otogic origins of the weakings

Duration of symptoms: mean 2.3 years (range 0.3 to 10 years)

Previous treatments: Not reported

Care setting: Not reported, probably tertiary or secondary

Participants Place of study: The Netherlands

Period of study: January 1993 to December 1995

Number of participants eligible: 48 participants

Number of participants enrolled IT and RS:

- US and arthroscopy or open surgery: 48 participants

Data available for analyses:

- US and arthroscopy or open surgery: 48 participants

Age: mean 55 years (range 30 to 76 years)

Male/Female: 28/20

Dominant arm: Not reported



Swen 1998 (Continued)	Nature of onset: Not re	eported	
Study design	Primary objective : To compare the diagnostic value of US performed by the rheumatologist with that of arthrography by a radiologist for otherwise healthy patients with suspected rotator cuff tears		
	Study design: Prospective consecutive accuracy cohort study		
	Language: English		
Target condition and ref-	Target conditions: Pre	esence full thickness tears	
erence standard(s)	Reference standard(s): Shoulder arthroscopy and open surgery		
	- Description of technic	<u>que</u> : Not reported	
	- Criteria for a positive	result:	
	A full-thickness was dia of the cuff	agnosed if free communication was found between the bursal and humeral sides	
Index and comparator	Index test(s): US		
tests	- Description of technic	<u>que</u> :	
	Scanner: 7.5 MHz linear array and the 5.0 MHz curved array transducers		
	<i>Technique</i> : The shoulder was examined in the anterior, lateral, and posterior directions, in both the transverse and the longitudinal planes as described by Van Holsbeeck 1991		
	Patient position: The patients were seated		
	For the anterior approach, the patient's upper arm was visualised in internal rotation, which was achieved by placing the patient's hand behind the back		
	- <u>Criteria for a positive result</u> :		
	Full thickness tears: a discontinuity in the rotator cuff extending from the bursal to the humeral side of the rotator cuff		
	- <u>Time from symptoms to index test</u> : Not reported		
	- <u>Time from index test to reference standard</u> : Not reported		
Follow-up	Adverse events due to	index test(s): Not reported	
	Adverse events due to	reference standard test(s): Not reported	
Notes	A two-by-two table of t	he ITs and RS was given which tallied with the reported summary data	
Table of Methodological Qu	ality		
Item	Authors' judgement	Description	
Representative spectrum? All tests	Yes	Tertiary or secondary care	
		Participants with suspected of having any rotator cuff tears The study was prospective and recruitment was consecutive	
Acceptable reference standard? All tests	Yes	The reference standard was arthroscopy or open surgery and the target conditions were presence of full thickness tears	



Swen 1998 (Continued)		
Acceptable delay between tests? All tests	Unclear	The study did not report the time elapsed between the index tests and reference standard
Partial verification avoided? All tests	Yes	All patients who received the index test went on to receive verification of their disease status using a reference standard
Differential verification avoided? All tests	Yes	All patients received the same reference standard, regardless of the result of their index test
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The study was prospective and the results of the index tests were interpreted before the reference standard
Reference standard results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Yes	The study was prospective, recruitment was consecutive and results were reported for all initially included participants
Withdrawals explained? All tests	Yes	No participants were excluded from the analysis
Learning curve / training reported of index test? All tests	Yes	The interpreter of index tests was a rheumatologist with experience in this technique
Learning curve / training reported of reference stan- dard? All tests	Unclear	Insufficient information was given to permit judgement
Index test criteria for a positive test result report- ed? All tests	Yes	The study reported the definition of a positive index test result

Swen 1999

Clinical features and settings

Inclusion criteria: Patients awaiting surgery because of a clinically suspected rotator cuff tears

The clinical diagnosis of rotator cuff tears was based on marked difficulty in initiating abduction of the arm with weakness and limitation of movement Lidocaine was injected below the acromion, if after the injection the strength of the rotator cuff was still decreased; this was considered to indicate rotator cuff tears

Exclusion criteria: Patients with neurologic origins of the weakness



Swen 1999 (Continued)

Duration of symptoms: mean 2.3 years (range 0.3 to 8 years)

Previous treatments: Not reported

Care setting: Not reported Probably tertiary or secondary

Participants Place of study: The Netherlands

Period of study: Not reported

Number of participants eligible: 21 participants

Number of participants enrolled IT and RS:

- MRI and arthroscopy: 21 participants

- US and arthroscopy: 21 participants

Data available for analyses:

- MRI and arthroscopy: 21 participants

- US and arthroscopy: 21 participants

Age (mean/SD): 54/12 years

Male/Female: 12/9

Dominant arm: Not reported

Nature of onset: In four patients the shoulder complaints could be attributed to trauma

Study design

Primary objective: To evaluate the ability of US and MRI to detect full thickness rotator cuff tears in patients with a clinically suspected rotator cuff tears as a solitary non-inflammatory condition

Study design: Prospective, consecutive, accuracy cohort study with fully paired direct comparison between MRI and US

Language: English

Target condition and reference standard(s)

Target conditions: Presence full thickness tears

Reference standard(s): Shoulder arthroscopy

- Description of technique:

First the arthroscope was introduced in the glenohumeral joint and then into the subacromial space

After introducing the scope into the subacromial space, the bursa was removed to enable examination of the bursal side of the cuff

- Criteria for a positive result:

A full-thickness was diagnosed if free communication was found between the bursal and humeral sides of the cuff

Index and comparator tests

Index test(s): MRI and US

- Description of technique:

MRI

MRI unit: 1.0T system with a dedicated shoulder coil as receiver



Swen 1999 (Continued)

Sequences: T1-weighted (TR/TE 680/15) and a standard T2 coronal spin-echo sequence (TR/TE 3000/15,105 ms)

Planes: Oblique coronal

Patient position: Supine position

US

Scanner: 7.5 MHz linear array and the 5.0 MHz curved array transducers

Technique: The shoulder was examined in the anterior, lateral, and posterior directions, in both the transverse and the longitudinal planes as described by Van Holsbeeck 1991

Patient position: The patients were seated

For the anterior approach, the patient's upper arm was visualised in internal rotation, which was achieved by placing the patient's hand behind the back

- Criteria for a positive result:

MRI: Full-thickness tears: a focal, well-defined area of increased signal intensity on T1-weighted and T2-weighted images that extended through the entire thickness of the tendon

US: Full-thickness tears: a discontinuity in the rotator cuff, extending from the bursal to the humeral side of the rotator cuff

- Time from symptoms to index test: Not reported
- Time from Conventional MRA and 3D isotropic MRA: Not reported
- <u>Time from index test to reference standard</u>: MRI and US were performed within 3 weeks before surgery

Raw data were given and it was possible to back-calculate this from the reported summary data

Follow-up Adverse events due to index test(s): Not reported Adverse events due to reference standard test(s): Not reported The results of index test were interpreted by two experienced musculoskeletal radiologists The data of only one reader (reader 1) were arbitrarily chosen to be included in our analyses

Item	Authors' judgement	Description
Representative spectrum? All tests	Yes	Tertiary or secondary care, participants with suspected of having any rotator cuff tears
		The study was prospective and recruitment was consecutive
Acceptable reference standard? All tests	Yes	The reference standard was arthroscopy and the target condition was presence of full thickness tears
Acceptable delay between tests? All tests	Yes	The index tests were performed within 3 weeks of surgery
Partial verification avoided?	Yes	All patients who received the index test went on to receive verification of their disease status using a reference standard



Swen 1999 (Continued) All tests		
Differential verification avoided? All tests	Yes	All patients received the same reference standard, regardless of the result of their index test
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The study was prospective and the results of the index tests were interpreted before the reference standard
Reference standard results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Relevant clinical information? All tests	No	The interpreters were blinded to history and physical examination
Uninterpretable results reported? All tests	Yes	The study was prospective, recruitment was consecutive and results were reported for all initially included participants
Withdrawals explained? All tests	Yes	No participants were excluded from the analysis
Learning curve / training	Unclear	A rheumatologist and a radiologist, both experienced with this test
reported of index test? All tests		In fact they had different results but this was not examined in this study
Learning curve / training reported of reference standard? All tests	Unclear	The reference standards were performed by a single experienced surgeon
Index test criteria for a positive test result report- ed? All tests	Yes	The study reported the definition of a positive index test result

Taboury 1992

Inclusion criteria: Not reported		
Exclusion criteria: Not reported		
Duration of symptoms: Not reported		
Previous treatments: Not reported		
Care setting: Tertiary		
Place of study: Paris, France		
Period of study: Not reported		
Number of participants eligible: 24 participants		



Ta	boui	у 199	2 (Con	tinued)

Number of participants enrolled IT and RS:

- US and open surgery: 24 participants

Data available for analyses:

- US and open surgery: 24 participants

Age: Not reported

Gender: Not reported

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To compare the results of US to open surgery in order to evaluate the characteris-

tics of the rotator cuff tears

Study design: Prospective accuracy cohort study

Unclear whether consecutive recruitment

Language: French

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears

Reference standard(s): Open surgery
- <u>Description of technique</u>: Not reported

- Criteria for a positive result: Not reported

Index and comparator tests

Index test(s): US

- Description of technique:

Scanner: 5 to 10 MHz linear or vectorial short focal probe

Technique: Static and dynamic examination of rotator cuff tendons

Patient position: Patients seated with the arm in adduction and internal rotation by asking the patients

to place their arm behind their back

- Criteria for a positive result: Not reported

- Time from symptoms to index test: Not reported

- Time from index test to reference standard: Not reported

Follow-up

Adverse events due to index test(s): Not reported

Adverse events due to reference standard test(s): Not reported

Notes

Mr Jean-Philippe Regnaux and Mr Ludovic Trinquart kindly translated into English and extracted the

data of this study

A two-by-two table of the ITs and RS was given, which tallied with the reported summary data

Item	Authors' judgement	Description
Representative spectrum?	Unclear	Unclear whether consecutive recruitment



Ta	bo	ury	1992	(Continued)

ΛΙ	ltes	
ΑI	1165	ı۶

All tests		
Acceptable reference standard? All tests	Unclear	The reference standard was open surgery and the target condition was presence of any rotator cuff tears
Acceptable delay between tests? All tests	Unclear	The study did not report the time elapsed between the index tests and reference standard
Partial verification avoided? All tests	Yes	All patients who received the index test went on to receive verification of their disease status using a reference standard
Differential verification avoided? All tests	Yes	All participants included in the analyses received open surgery, regardless of the results of their index test
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The person undertaking the index test was blinded to the results of the standard reference
Reference standard results blinded? All tests	Yes	The reference standard results were performed blind to the results of the index test
Relevant clinical information? All tests	Unclear	Insufficient information was given
Uninterpretable results reported? All tests	Yes	The number of results reported agrees with the number of patients recruited
Withdrawals explained? All tests	Yes	No participants were excluded from the analysis
Learning curve / training reported of index test? All tests	Unclear	Insufficient information was given
Learning curve / training reported of reference stan- dard? All tests	Unclear	Insufficient information was given
Index test criteria for a positive test result reported? All tests	No	Not reported

Teefey 2004

Clinical features and set-	Inclusion criteria : Acute or chronic shoulder pain accompanied by a high clinical suspicion of rotator
tings	cuff disease



Teefe	y 2004	(Continued)
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Exclusion criteria: Participants with severe claustrophobia, which is a contraindication for magnetic resonance imaging; a previous operation on the shoulder; a humeral fracture; and inflammatory arthri-

tis

Duration of symptoms: Not reported **Previous treatments**: Not reported

Care setting: Not reported, probable tertiary or secondary

Participants

Place of study: St. Louis, Missouri, USA

Period of study: December 1998 and April 2001

Number of participants eligible: 130 participants

Number of participants enrolled IT and RS:

- MRI and arthroscopy: 71 shoulders

- US and arthroscopy: 71 shoulders

Data available for analyses:

- MRI and arthroscopy: 71 shoulders

- US and arthroscopy: 71 shoulders

Age: mean 59 (range 31 to 80 years)

Male/Female: 41/30

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To compare the diagnostic performances of US and MRI for both the detection of a rotator cuff tear and the quantification of its size, with use of arthroscopic findings as the standard

Study design: Prospective, consecutive, accuracy cohort study with fully paired direct comparison between US and MRI

Language: English

Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears

Reference standard(s): Shoulder arthroscopy

- Description of technique:

The rotator cuff was examined from both the articular and the bursal side. A tagged suture (number-1 PDS [polydioxanone]) was placed, during intra-articular viewing, through the suspected region of the cuff tear to guide arthroscopic bursal imaging

- Criteria for a positive result:

The presence or absence of a rotator cuff tear and the size and extent of the tear, when present, were recorded. Specifically, the presence or absence of a full thickness tear or of a bursal or articular-side partial thickness tear and the width (perpendicular to the long axis of the cuff fibres) of any tear that was found were recorded

Index and comparator tests

Index test(s): MRI and US

- Description of technique:



Teefey 2004 (Continued)

MRI

MRI unit: 1.5 T with high field strength with a two-piece shoulder array coil (54 participants); and with flexible local coils (17 participants)

Sequences: Fat-suppressed, fast-spin-echo, proton-density-weighted, spin-echo, or fast-spin-echo and transverse, T2-weighted, fast-spin-echo images with or without fat suppression

Planes: Oblique coronal and oblique sagittal

Patient position: Not reported

US

Scanner: 7.5 to 9 MHz linear-array transducer in real-time

Technique and Patient position: As previously described (Teefey 2000).

- Criteria for a positive result:

MRI: Full thickness tears: complete disruption of all tendon fibres or when the signal within the cuff tendons was isointense compared with fluid on the T2-weighted images and extended from the articular to the bursal surface on one or more images. Partial thickness tears: fluid-intensity signal within the tendons was in contact with only one of the surfaces

US: Full thickness tears: non-visualisation of rotator cuff or a focal defect in the rotator cuff created by a variable degree of retraction of the torn tendon ends. Partial thickness tears: minimal flattening of the bursal side of the rotator cuff (bursal-side tear) or a distinct hypoechoic or mixed hyperechoic and hypoechoic defect visualized in both the longitudinal and the transverse plane at the deep articular side of the rotator cuff (articular-side tear)

- Time from symptoms to index test: Not reported
- <u>Time from US and MR!</u>: MRI was performed on the same day as the US for all but three patients, two of whom had the studies six days apart and one of whom had them one day apart
- Time from index test to reference standard: mean 56 days (range 2 to 190 days)

Follow-up	Adverse events due to index test(s): Not reported
	Adverse events due to reference standard test(s): Not reported
Notes	Part of population (only the participants with full thickness rotator cuff tears) of this study was also reported in Teefey 2005
	A two-by-two table of the ITs and RS was given, which tallied with the reported summary data with a few discrepancy

Item	Authors' judgement	Description
Representative spectrum? All tests	Unclear	Insufficient information was given to permit judgement
Acceptable reference standard? All tests	Yes	The reference standard was arthroscopy and the target condition were presence of any rotator cuff tears, full thickness tears and partial thickness tears
Acceptable delay between tests? All tests	No	The average interval between reference standard and index test was 56 days



Teefey 2004 (Continued)		
Partial verification avoided?	No	Not all the patients who received the index test underwent a reference standard to verify their true disease status
All tests		Of the 130 eligible participants 71 underwent to reference standard
Differential verification avoided? All tests	No	The result of the index test probably influenced the choice of the reference standard
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The person undertaking the index test was blinded to the results of the standard reference
Reference standard results blinded? All tests	No	The results of the index tests were known to the person interpreting the reference tests
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Unclear	Insufficient information was given to permit judgement
Withdrawals explained? All tests	No	Some of the eligible patients who entered the study did not complete it and these patients were not accounted for
Learning curve / training reported of index test?	Yes	The interpreter of MRI was one of six radiologists with extensive experience in musculoskeletal magnetic resonance imaging
All tests		The interpreter of US was one of two radiologists who were very experienced with the technique and who had conducted more than 2500 examinations during a 10-year period
Learning curve / training reported of reference stan- dard? All tests	Unclear	The reference standards were performed by an experienced orthopaedic surgeon
Index test criteria for a positive test result reported? All tests	Yes	The study reported the definition of a positive index test result

Venu 2002

Clinical features and settings

Inclusion criteria: Participants with clinical supraspinatus impingement syndrome and failure of clinical improvements with conservative treatment within one year of onset of symptoms

Exclusion criteria: Not reported

Duration of symptoms: Probably more than one year

Previous treatments: Shoulder physiotherapy and sub-acromial steroid injections



Venu 2002 (Continued)

Care setting: Not reported

Participants Place of study: Eastbourne, UK

Period of study: June 1997 to June 1999

Number of participants eligible: 276 participants

Number of participants enrolled IT and RS:

- US and arthroscopy: 41 participants

Data available for analyses:

- US and arthroscopy: 41 participants

Age: mean 54 years (range 34 to 79 years)

Male/Female: 24/17

Dominant arm: Not reported **Nature of onset**: Not reported

Study design

Primary objective: To determine the accuracy of ultrasound compared with arthroscopy in the evaluation of the symptomatic supraspinatus tendon and to identify whether ultrasound diagnosis was helpful in pre-operative planning

Study design: Prospective, consecutive, accuracy cohort study

Language: English

Target condition and reference standard(s)

Target conditions: Presence of normal tendon, tendinopathy, partial thickness tear, full thickness tear,

and rotator cuff rupture

Reference standard(s): Shoulder arthroscopy

- Description of technique: Not reported

- Criteria for a positive result: Not reported

Index and comparator tests

Index test(s): US

- Description of technique:

Scanner: 5 to 10 MHz using a linear array transducer

Technique: Longitudinal and transverse views of the supraspinatus

Patient position: Patient probably seated with the shoulder internally rotated to visualise the supraspinatus tendon

- Criteria for a positive result:

Tendinopathy: thickened and often decreased echogenicity

Partial thickness tears: a hypo-or hyperechoic tendon defect not involving the full thickness of the ten-

don

Full thickness tears: a hypo or hyperechoic tendon defect involving the full thickness of the tendon

Rupture: the tendon was absent with often only the retracted proximal tendon visualised

- Time from symptoms to index test: More than 1 year



Venu 2002 (Continued)	- <u>Time from index test to reference standard</u> (mean): 6 months
Follow-up	Adverse events due to index test(s): Not reported
	Adverse events due to reference standard test(s): Not reported
Notes	The study reported five categories to classify the tendon (normal tendon, tendinopathy, partial thickness tear, full thickness tear, and rotator cuff rupture)
	In our analyses we classified the categories 'normal' and 'tendinopathy' as normal tendon; and 'full' and 'rupture' as full thickness tear
	No two-by-two table of the ITs and RS was given but it was possible to back-calculate this from the reported summary data

Item	Authors' judgement	Description
Representative spectrum?	Unclear	Population was patients with suspected of having any rotator cuff tears
All tests		The study was prospective and recruitment was reported as consecutive
		The care setting was not specified
Acceptable reference standard? All tests	Yes	The reference standard was arthroscopy and the target conditions were presence of normal tendon, tendinopathy, partial thickness tear, full thickness tear, and rotator cuff rupture
Acceptable delay between tests? All tests	No	The average interval between reference standard and index test was 6 months
Partial verification avoided?	No	Not all the patients who received the index test underwent a reference standard to verify their true disease status
All tests		Of the 276 eligible participants only 41 (15%) received the reference standard
Differential verification avoided? All tests	No	The result of the index test influenced the choice of the reference standard
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The study was prospective and the results of the index tests were interpreted before the reference standard
Reference standard results blinded? All tests	No	The surgeon knew the US diagnosis prior to surgery
Relevant clinical information? All tests	Yes	Clinical data and plain radiographs were available at the time of performing the US examination
Uninterpretable results reported? All tests	No	The results of 235 (85%) patients were not reported



Venu 2002 (Continued)				
Withdrawals explained? All tests	No	Some of the eligible patients who entered the study did not complete it and these patients were not accounted for		
Learning curve / training reported of index test? All tests	Yes	The interpreters of index tests were two radiologists specialised in shoulder US		
Learning curve / training reported of reference stan- dard? All tests	Unclear	Insufficient information was given to permit judgement		
Index test criteria for a positive test result report- ed? All tests	Yes	The study reported the definition of a positive index test result		
Wallny 2001				
Clinical features and set- tings	Inclusion criteria : Participants suffering from shoulder pain with histories and physical examinations suggestive of rotator cuff lesions			
	Exclusion criteria : Participants with prior shoulder surgery or previous fracture of the humeral head			
	Duration of symptoms: Not reported			
	Previous treatments: Not reported			
	Care setting: Tertiary or secondary			
Participants	Place of study: Bonn, Germany			
	Period of study: N	Not reported		
	Number of partic	ipants eligible: 40 participants		
	Number of partic	ipants enrolled IT and RS:		
	- Two-dimensional (2D) US and arthroscopy or open surgery: 40 participants			
	- Tree-dimensional (3D) US and arthroscopy or open surgery: 40 participants			
	Data available for analyses:			
	- 2D US and arthroscopy or open surgery: 40 participants			
	- 3D US and arthroscopy or open surgery: 40 participants			
	Age: mean 54 years (range 38 to 79 years)			
	Male/Female: 25/15			
	Dominant arm: Not reported			
	Nature of onset:	Not reported		

Primary objective: To determine the validity of 3D US in the diagnosis of rotator cuff lesions

Study design: Prospective, consecutive, accuracy cohort study with fully paired direct comparison be-

tween 3D US and 2D US

Study design



Nallny 2001 (Continued)	Language: English			
Target condition and ref-	Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness tears			
erence standard(s)	Reference standard(s): Shoulder arthroscopy or open surgery			
	- <u>Description of technique</u> : Not reported			
	- <u>Criteria for a positive result</u> : Not reported			
Index and comparator	Index test(s): 3-D US and 2-D US			
tests	- Description of technique:			
	Scanner: 10 MHz electronic linear array in broad bandwidth technology, 192 fine pitch elements, frequency ranges: resolution: 4.5 to 13 MHz, penetration: 2.5 to 10 MHz			
	Technique and Patient position: Not reported			
	The region of interest was defined by 2D US before 3D US could be undertaken			
	- <u>Criteria for a positive result</u> :			
	Full thickness tear was defined as: marked thinning, sudden changes of calibre, hyper- and/or hypoe-choic zones and total absence of the cuff			
	Partial thickness tear was defined as: constituting no more than loss of $1/4$ to $1/2$ of full thickness of the intact rotator cuff			
	- <u>Time from symptoms to index test</u> : Not reported			
	- <u>Time from 2D US and 3D US</u> : in the same examination			
	- <u>Time from index test to reference standard</u> : Not reported			
Follow-up	Adverse events due to index test(s): Not reported			
	Adverse events due to reference standard test(s): Not reported			
Notes	The study reported that the target conditions were presence of any rotator cuff tears, full thickness tears and partial thickness tears			
	Only the data for analysing presence of any rotator cuff tears were available			
	The study reported the data of two different types of US (three-dimensional and two-dimensional)			
	Inasmuch as the 2D US examinations are more often used in clinical practice we arbitrarily chose 2D US to be included in our analyses			
	A two-by-two table of the ITs and RS was given which tallied with the reported summary data			

Item	Authors' judgement	Description
Representative spectrum? All tests	Yes	Tertiary or secondary care, participants with suspected of having any rotator cuff tears
		The study was prospective and recruitment was consecutive
Acceptable reference standard? All tests	Unclear	The reference standard was arthroscopy or open surgery and the target conditions were presence of full thickness tears and partial thickness tears



Wallny 2001 (Continued)		
Acceptable delay between tests? All tests	Unclear	The study did not report the time elapsed between the index tests and reference standard
Partial verification avoided? All tests	Yes	All patients who received the index test went on to receive verification of their disease status using a reference standard
Differential verification avoided? All tests	Yes	The indication for surgery was based on the results of clinical assessment and an MRI scan but independent of the result of the index test
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The study was prospective and the results of the index tests were interpreted before the reference standard
Reference standard results blinded? All tests	Unclear	Insufficient information was given to permit judgement
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results re-	Yes	The study was prospective
ported? All tests		Recruitment was consecutive and results were reported for all initially included participants
Withdrawals explained? All tests	Yes	No participants were excluded from the analysis
Learning curve / training reported of index test? All tests	Unclear	Insufficient information was given to permit judgement
Learning curve / training reported of reference stan- dard? All tests	Unclear	The reference standards were performed by a single orthopaedic surgeon
Index test criteria for a positive test result report- ed? All tests	Yes	The study reported the definition of a positive index test result

Yen 2004

Clinical features and settings

Inclusion criteria: Patients with shoulder pain and suspected of rotator cuff tears

Exclusion criteria: Not reported

Duration of symptoms: Not reported **Previous treatments**: Not reported



en 2004 (Continued)	Care setting: Not reported		
Participants	Place of study: Taiwan, China		
	Period of study: Not reported		
	Number of participants eligible: 50 participants		
	Number of participants enrolled IT and RS:		
	- US and open surgery: 50		
	Data available for analyses:		
	- US and open surgery: 50		
	Age: mean 63 years (range 17 to 81 years)		
	Male/Female: 26/24		
	Dominant arm: Not reported		
	Nature of onset: Not reported		
Study design	Primary objective : To prospectively compare the US and operative findings of rotator cuff tears		
	Study design: Prospective accuracy cohort study		
	Unclear whether consecutive recruitment		
	Language: English		
Target condition and ref-	Target conditions: Presence any rotator cuff tears		
erence standard(s)	Reference standard(s): Open surgery		
	- <u>Description of technique</u> : Not reported		
	- <u>Criteria for a positive result</u> : Not reported		
Index and comparator	Index test(s): US		
tests	- <u>Description of technique</u> :		
	Scanner: 7 MHz linear transducer		
	Technique: Longitudinal, transverse and oblique scans of the tendons were used		
	Patient position: Probably patient seated with the arm in		
	External rotation for scanning the subscapularis tendon		
	Neutral position for the long head of the biceps tendon		
	Internal rotation and with the patient's hand behind the back with extreme internal rotation for the supraspinatus tendon		
	Flexion and adduction for infraspinatus and teres minor tendons		
	- <u>Criteria for a positive result</u> :		
	Six US signs were used: non-visualisation;		
	Floating bright spots		
	Focal depression		



Yen 2004	(Continued)
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Focal thinning

Focal hypoechoic cleft

Focal heterogeneous hypoechogenicity

- Time from symptoms to index test: Not reported

- Time from index test to reference standard: within 1 month

Follow-up Adverse events due to index test(s): Not reported

Adverse events due to reference standard test(s): Not reported

Notes A two-by-two table of the ITs and RS was given which tallied with the reported summary data

Item	Authors' judgement	Description
Representative spectrum?	Unclear	Population was patients with suspected of having any rotator cuff tears
All tests		The study was prospective
		It was unclear whether consecutive recruitment
Acceptable reference standard? All tests	Unclear	The reference standard was open surgery and the target condition was presence of any rotator cuff tears
Acceptable delay between tests? All tests	Yes	The reference standard was performed within 1 month after the index test
Partial verification avoided? All tests	Yes	All patients who received the index test went on to receive verification of their disease status using a reference standard
Differential verification avoided? All tests	Yes	All patients received the same reference standard regardless of the result of their index test
Incorporation avoided? All tests	Yes	The index test did not form part of the reference standard
Index test results blinded? All tests	Yes	The index tests were interpreted before and without knowledge of the reference standard results
Reference standard results blinded? All tests	No	The results of the index tests were known to the person interpreting the reference tests
Relevant clinical information? All tests	Unclear	Not reported
Uninterpretable results reported? All tests	Yes	The study was prospective and results were reported for all initially included participants



Yen 2004 (Continued)		
Withdrawals explained? All tests	Yes	No participants were excluded from the analysis
Learning curve / training reported of index test? All tests	Unclear	All of the procedures were performed by one sonologist and the findings were interpreted by two or three sonologists in consensus prior to surgery
Learning curve / training reported of reference standard? All tests	Unclear	Insufficient information was given to permit judgement
Index test criteria for a positive test result reported? All tests	Yes	The study reported the definition of a positive index test result

<: less than

>: more than

IT: Index test

MHz: Megahertz

RS: Reference standard

T: Tesla

T1-weighted: Short TR and short TE sequences T2-weighted: Long TR and long TE sequences

TE: Echo time TR: Repetition time

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Adams 2010	This was a retrospective study with a possible risk of spectrum and verification bias
Al-Shawi 2008	Reference standard not relevant: Arthroscopy or MRI was used as reference standard
Aliabadi 1991	Type of study not relevant: Narrative review
Aliprandi 2006	Participants not relevant: Participants with suspected of chronic or traumatic rotator cuff tear, congenital atraumatic or traumatic glenohumeral instability, traumatic rotator cuff tear and glenohumeral instability, and "frozen shoulder" were enrolled
Allmann 1999	Type of study not relevant: Technique report
Ardic 2006	Reference standard not relevant: MRI was used as reference standard
Auethavekiat 2006	Type of study not relevant: Case report
Awerbuch 2008	Type of study not relevant: Narrative review
Balich 1997	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears
Bencardino 2010	Type of study not relevant: Narrative review



Study	Reason for exclusion	
Blanchard 1999a	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Blanchard 1999b	Type of study not relevant: Questionnaire study	
Blum 1993	Index test not relevant: Arthrographic computed tomography was used as index test	
Boisrenoult 1999	Type of study not relevant: Index test was not compared with reference standard(s)	
Boorstein 1992	Type of study not relevant: Narrative review	
Brandt 1989	Reference standard not relevant: Arthrography or surgery was used as reference standard	
Brasseur 1994	Type of study not relevant: Anatomic description	
Brenneke 1992	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Bryant 2002	Type of study not relevant: The purpose of this study was to determine how well the size of rotator cuff tears could be estimated noninvasively by ultrasonography and MRI and how well arthroscopy could detect the size of rotator cuff tears	
Burk 1989	Reference standard not relevant: Arthrography or surgery was used as reference standard	
Chang 2002	Participants not relevant: Participants with shoulder pain; however, it was unclear if all participants were suspected of having rotator cuff tears	
Chaubal 2007	Type of study not relevant: Narrative review	
Chen 1996	Target condition not relevant: The aim of the study was to determine the MRI findings that are associated with full thickness rotator cuff tears	
Chiodi 1994	Participants not relevant: Selective population that all participants had full thickness tears (100% of prevalence) and the study was retrospective and reported non-consecutive recruitment	
Chiodi 1995	Participants not relevant: Selective population that all participants had rotator cuff tears (100% of prevalence)	
	The study also included patients that were reported in Chiodi 1994	
Chiou 1999	This was a retrospective study with a possible risk of spectrum and verification bias	
Chucair 2008	Type of study not relevant: Narrative review	
Chun 2010	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Crass 1987	Type of study not relevant: Case report	
Crass 1988	This was a retrospective study with a possible risk of spectrum and verification bias	
Cullen 2007	This was a retrospective study with a possible risk of spectrum and verification bias	
Cusmano 2000	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	



Study	Reason for exclusion	
D'Erme 1993	Reference standard not relevant: Surgery or arthrography was used as reference standard	
Davidson 2005	Target condition not relevant: To determine the MRI criteria for predicting rotator cuff tear pattern and method of repair	
Davis 1991	Type of study not relevant: Technique report	
De Muynck 1994	Reference standard not relevant: Arthrography or arthroscopy or open surgery was used as reference	
	standard	
Demouy 1993	Type of study not relevant: Narrative review	
Deutsch 1997	Participants not relevant: Selective population, restricted to subscapularis tendon tear (retrospective, so selected out patients with the diagnosis)	
Dhagat 2002	Type of study not relevant: Index test (US) was not compared with reference standard(s)	
Dinter 2008	Participants not relevant: Participants with shoulder pain; however, it was unclear if all participants were suspected of having rotator cuff tears	
Drakeford 1990	Participants not relevant: Asymptomatic participants were included	
El-Dalati 2005	Insufficient data to be included in the meta-analyses	
El-Kouba 2010	This was a retrospective study with a possible risk of spectrum and verification bias	
Evancho 1988	Reference standard not relevant: Arthroscopy or arthrography as reference standard	
Fabis 1999a	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Fabis 1999b	Participants not relevant: The aim was to evaluate US images of rotator cuff integrity after surgical repair	
Farin 1995	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears	
Farin 1996a	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears	
Farin 1996b	Participants not relevant: Selective population that all participants had full thickness tears (100% of prevalence) and the study was retrospective and reported non-consecutive recruitment	
Farley 1992	Target condition not relevant: The aim of the study was to determine the MRI findings that are associated with full thickness rotator cuff tears	
Ferrari 2002	Participants not relevant: Selective population that all participants had full thickness tears (100% of prevalence) and the study reported non-consecutive recruitment	
Ferri 2005	Target condition not relevant: The aim of the study is to assess the accuracy of the Crass and modified Crass positions	
Flannigan 1990	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	



Study	Reason for exclusion	
Fotiadou 2008	This was a retrospective study with a possible risk of spectrum and verification bias	
Frei 2008	This was a retrospective study with a possible risk of spectrum and verification bias	
Fritz 1992	Type of study not relevant: Letter	
Furtschegger 1988	This was a retrospective study with a possible risk of spectrum and verification bias	
Girard 1995	Type of study not relevant: Narrative review	
Goergen 1996	Type of study not relevant: Technique report	
Goldberg 2003	Reference standard not relevant: Arthrography findings and clinical examination were used as reference standard	
Hedtmann 2002	Type of study not relevant: Narrative review	
Heijne 2004	Type of study not relevant: Editorial letter	
Herold 2006	Participants not relevant: A history of trauma was reported in 17 (33%) of 51 patients	
	Fourteen (27%) of 51 patients had previous shoulder dislocation, and 36 (71%) presented with clinical signs of impingement	
Herzog 1997	Type of study not relevant: Narrative review	
Herzog 1998	Type of study not relevant: Narrative review	
Hodler 1987	Reference standard not relevant: Arthrography was used as reference standard	
Hodler 1988	This was a retrospective study with a possible risk of spectrum and verification bias	
Hodler 1992	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Hollister 1995	Target condition not relevant: The aim of the study was to determine the association between bursal and joint effusion (index tests findings) that are associated with rotator cuff tears	
Homsi 1989	This was a retrospective study with a possible risk of spectrum and verification bias	
Horii 1998	Participants not relevant: Selective population that all participants had full thickness tears (100% of prevalence) and the study was retrospective and reported non-consecutive recruitment	
lannotti 1991	Participants not relevant: Asymptomatic participants were enrolled	
Imhoff 1992	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Imhoff 1993	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Imhoff 1996	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
lovane 2001	Participants not relevant: Selective population that all participants had full thickness tears (100% of prevalence) and the study was retrospective and reported non-consecutive recruitment	



Study	Reason for exclusion	
lyengar 2010	Type of study not relevant: Technique report	
Jacobson 2003	Type of study not relevant: Narrative review	
Jacobson 2004	Target condition not relevant: The aim of the study was to determine which US signs are important for the diagnosis of a surgically identifiable supraspinatus tendon tear	
Jaovisidha 1999	Type of study not relevant: The time elapsed between the index and reference tests was during a 26-month follow-up	
Jeyam 2008	This was a retrospective study with a possible risk of spectrum and verification bias	
Jung 2009	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Jung 2010	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Kaneko 1994	Participants not relevant: A control group without suspected of rotator cuff tears was included	
Kautzner 2008	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Kelly 2009	Type of study not relevant: Diagnostic Test Accuracy review	
Kerkovsky 2008	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Kluger 2003	Target condition not relevant: The aim of this study was to evaluate the accuracy of MRA and US in millimetres for both width and retraction of full-thickness rotator cuff tears, and not to detect the tears	
Kneeland 1987	Reference standard not relevant: Arthroscopy or arthrography was used as reference standard	
Kujat 1986	Type of study not relevant: Technique report	
Kurol 1991	This was a retrospective study with a possible risk of spectrum and verification bias	
Lawson 1991	Type of study not relevant: Narrative review	
Lee 2002	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Lipman 1992	Type of study not relevant: Letter	
Loew 2000	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Lopez 2007	This was a retrospective study with a possible risk of spectrum and verification bias	
Low 1998	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Lucas 1991	Type of study not relevant: Narrative review	
Mack 1988	This was a retrospective study with a possible risk of spectrum and verification bias	



Study	Reason for exclusion	
Magee 2003a	Participants not relevant: Participants with shoulder pain; however, it was unclear if all participants were suspected of having rotator cuff tears	
Magee 2003b	Participants not relevant: Participants with shoulder pain; however, it was unclear if all participants were suspected of having rotator cuff tears	
Magee 2006	Participants not relevant: Participants with shoulder pain; however, it was unclear if all participants were suspected of having rotator cuff tears	
Magee 2007	Participants not relevant: Participants with clinical diagnosis of pain or instability or both were enrolled	
Magee 2009	Participants not relevant: Participants with shoulder pain; however, it was unclear if all participants were suspected of having rotator cuff tears	
Malvestiti 1997	Reference standard not relevant: Arthroscopy or MRI or arthrography was used as reference standard	
Martin 2008	Type of study not relevant: Technique report	
Masaoka 1999	Participants not relevant: Participants who underwent index test after surgery were enrolled	
Masciocchi 1989	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled	
Meister 2004	This was a retrospective study with a possible risk of spectrum and verification bias	
Mendieta-Sevilla 2009	Reference standard not relevant: Surgery or MRI or arthrography or rehabilitation was used as reference	
	standard	
Merl 1996	Type of study not relevant: Narrative review	
Middleton 1993	Type of study not relevant: Letter	
Miller 2008	This was a retrospective study with a possible risk of spectrum and verification bias	
Montrucchio 1997	This was a retrospective study with a possible risk of spectrum and verification bias	
Monu 1994	Participants not relevant: The study included selective participants without rotator cuff tears	
Moosmayer 2005	Participants not relevant: Participants with other shoulder complaints, including symptoms from the long head of the biceps muscle were enrolled	
Moosmayer 2007	Participants not relevant: Participants with shoulder pain; however, it was unclear if all participants were suspected of having rotator cuff tears	
Morrison 1990	Reference standard not relevant: Arthrography or arthroscopy or open surgery was used as reference standard	
Naqvi 2009	This was a retrospective study with a possible risk of spectrum and verification bias	
Narbona 2007	Target condition not relevant: The aim of this study was to detect SLAP lesion in patients with rotator cuff tears	



Study	Reason for exclusion
Needell 1997	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Nelson 1991	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Nogueira-Barbosa 2002	This was a retrospective study with a possible risk of spectrum and verification bias
Norregaard 2002	Participants not relevant: Participants with clinical suspicion of labral or rotator cuff lesion were enrolled
Oh 2009	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Oh 2010	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Ostlere 1997	Type of study not relevant: Narrative review
Ozcakar 2005	Type of study not relevant: Letter
Paavolainen 1994	This was a retrospective study with a possible risk of spectrum and verification bias
Palmer 1993	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Palmer 1994	Type of study not relevant: Narrative review
Parsa 1997	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Pattee 1988	This was a retrospective study with a possible risk of spectrum and verification bias
Patten 1994	Reference standard not relevant: Arthroscopy, arthrography and non-surgical therapy were used as reference standard
Peetrons 1986	Type of study not relevant: Index test was not compared with reference standard(s).
Pfirrmann 1999	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Pfirrmann 2004	Participants not relevant: Asymptomatic participants were included
Pigeau 1992	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears
Poey 1990	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears
Porcellini 1994	Reference standard not relevant: Surgery or arthrography was used as reference standard
Prendergast 1992	Type of study not relevant: Narrative review
Quinn 1995	This was a retrospective study with a possible risk of spectrum and verification bias
Rafii 1990	Participants not relevant: Asymptomatic participants were included



Study	Reason for exclusion
Read 1998	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all
	participants were suspected of having rotator cuff tears
Recht 1993	Type of study not relevant: Narrative review.
Recht 1994	Type of study not relevant: Narrative review.
Reinus 1995	Participants not relevant: Participants with shoulder pain; however, it was unclear if all participants had suspected of having rotator cuff tears.
Roberts 1998	Reference standard not relevant: MRI or arthrography was used as reference standard.
Roberts 2001	Participants not relevant: Participants with other shoulder complaints, including adhesive capsulitis and osteoarthritis were enrolled.
Robertson 1995	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Rouaud 1989	Type of study not relevant: Index test (US) was not compared with reference standard(s)
Rubin 1997	Type of study not relevant: Letter
Rutten 2010a	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears
Rutten 2010b	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears.
Sahin-Akyar 1998	Participants not relevant: Participants with rotator cuff tear and other disorders were enrolled
Sartoris 1992	Type of study not relevant: Narrative review
Sasaki 1990	Participants not relevant: Asymptomatic participants were included
Schneider 2003	Insufficient data to be included in the meta-analyses
Schreinemachers 2009	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Schultz 1994	Type of study not relevant: Letter
Seeger 1988	Type of study not relevant: The study did not describe the comparison between the index test and the reference standard
Sheah 2009	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Shellock 1996	Type of study not relevant: Narrative review
Shellock 2001	Participants not relevant: The authors reported that participants with suspected of 'shoulder pathology' were included, probable included participants with suspected of rotator cuff tears and shoulder instability
Shiv 1990	Type of study not relevant: Index test (US) was not compared with reference standard(s)



Study	Reason for exclusion
Singer 1995	Type of study not relevant: Index test was not compared with reference standard(s)
Singson 1996	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears
Skib 1998	Type of study not relevant: Letter
Soble 1989	This was a retrospective study with a possible risk of spectrum and verification bias
Sonin 1996	This was a retrospective study with a possible risk of spectrum and verification bias
Sonnabend 1997	This was a retrospective study with a possible risk of spectrum and verification bias
Soto Araiza 1998	Reference standard not relevant: Surgery or MRI was used as reference standard
Steinbach 2000	Type of study not relevant: Narrative review
Strauss 1998	This was a retrospective study with a possible risk of spectrum and verification bias
Suder 1994	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Sunde 2001	Type of study not relevant: Letter
Sunde 2008	Type of study not relevant: Letter
Taboury 1995	Participants not relevant: Selective population that all participants had full thickness tears (100% of prevalence) and the study was retrospective and reported non-consecutive recruitment
Takagishi 1993	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Takagishi 1996	This was a retrospective study with a possible risk of spectrum and verification bias
Teefey 2000	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears
Teefey 2009	Type of study not relevant: Case report
Theodoropoulos 2010	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Tirman 1994	Participants not relevant: Selective population of five professional throwing athletes were evaluated; and, these participants had other shoulder complaints, including instability
Torstensen 1999	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Touzard 1991	Reference standard not relevant: Surgery or arthrography was used as reference standard
Toyoda 2005	Participants not relevant: Selective population that all participants had full thickness tears (100% of prevalence) and the study was retrospective and reported non-consecutive recruitment
Traughber 1992	This was a retrospective study with a possible risk of spectrum and verification bias



Study	Reason for exclusion
Traughber 1996	Type of study not relevant: Letter
Traughber 2006	Type of study not relevant: Letter
Tuite 1994	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Tuite 1995	Participants not relevant: It was unclear if all participants were suspected of having rotator cuff tears;
	furthermore, the study reported that part of participants of Tuite 1994 were included
	Thus, participants with other shoulder complaints, including instability were enrolled
Tuite 1998	Participants not relevant: It was unclear if all participants were suspected of having rotator cuff tears; furthermore, the study reported that part of participants of Tuite 1994 were included
	Thus, participants with other shoulder complaints, including instability were enrolled
Tuite 2001	Participants not relevant: It was unclear if all participants were suspected of having rotator cuff tears; probable the study included participants with shoulder instability
Turrin 1997	This was a retrospective study with a possible risk of spectrum and verification bias
Vahlensieck 2001	Type of study not relevant: Letter
Van Dyck 2009	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Van Holsbeeck 1995	This was a retrospective study with a possible risk of spectrum and verification bias
Van Moppes 1995	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Vander Maren 1995	Participants not relevant: Participants with shoulder pain were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears
Vanecek 2000	Type of study not relevant: Narrative review
Waldt 2007	Participants not relevant: Participants with symptomatic shoulder; however, it was unclear if all participants were suspected of having rotator cuff tears
Wallny 1999	Type of study not relevant: Technique report
	The study described an index to improve the accuracy of diagnosis of chronic rotator cuff tears
Walz 2007	Target condition not relevant: The aim of this study was a description of delamination tears of the supraspinatus, subscapularis, infraspinatus or teres minor tendons, as well as for mention of partial or full thickness tears
Wang 1994	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Weinstabl 1988	Type of study not relevant: Technique report
Wiener 1993	This was a retrospective study with a possible risk of spectrum and verification bias



Study	Reason for exclusion
Wilson 1994	Type of study not relevant: Letter
Wnorowski 1997	Participants not relevant: Participants with shoulder problems were enrolled; however, it was unclear if all participants were suspected of having rotator cuff tears
	The study reported that in the majority of the participants the primary diagnosis was unclear after the clinical evaluation
Wu 2003	This was a retrospective study with a possible risk of spectrum and verification bias
Yagci 2001	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Yamakawa 2001	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Yeh 2003	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled
Yeu 1994	This was a retrospective study with a possible risk of spectrum and verification bias
Zehetgruber 2002	This was a retrospective study with a possible risk of spectrum and verification bias
Ziegler 2004	This was a retrospective study with a possible risk of spectrum and verification bias
Zlatkin 1989	This was a retrospective study with a possible risk of spectrum and verification bias
Zlatkin 2004	Participants not relevant: Participants with other shoulder complaints, including instability were enrolled

Characteristics of studies awaiting classification [ordered by study ID]

Engebretsen 1994

Lingcoretacii 1994	
Clinical features and settings	
Participants	Number of participants eligible: 41 participants
	Number of participants enrolled IT and RS:
	- MRI and surgery: 25 participants
Study design	
Target condition and reference	Target conditions: Presence of any rotator cuff tears
standard(s)	Reference standard(s): Surgery
	Unclear whether arthroscopy or open surgery
Index and comparator tests	Index test(s): MRI
Follow-up	
Notes	Awaiting translation - Norwegian article



Engebretsen 1994 (Continued)

The information was collected from titles and abstracts that were reported in English

Farin 1990

Clinical features and settings	
Participants	Number of participants eligible: 301 participants
	Number of participants enrolled IT and RS:
	- US and surgery: 66 participants
Study design	
Target condition and reference	Target conditions: Presence of any rotator cuff tears
standard(s)	Reference standard(s): Surgery
	Unclear whether arthroscopy or open surgery
Index and comparator tests	Index test(s): US
Follow-up	
Notes	Awaiting translation - German article
	The information was collected from titles and abstracts that were reported in English

Guo 2000

Guo 2000	
Clinical features and settings	
Participants	Number of participants eligible: 53 participants
	Number of participants enrolled IT and RS:
	- MRI and surgery: 53 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Open surgery or arthroscopy
Index and comparator tests	Index test(s): MRI
Follow-up	
Notes	Awaiting translation - Chinese article
	The information was collected from titles and abstracts that were reported in English
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Clinical features and settings	
Participants	Number of participants eligible: 49 participants
	Number of participants enrolled IT and RS:
	- US and surgery: 17 participants
Study design	
Target condition and reference standard(s)	Target conditions: Presence of any rotator cuff tears
	Reference standard(s): Open surgery
Index and comparator tests	Index test(s): US
Follow-up	
Notes	Awaiting translation - German article
	-
edtmann 1995 Clinical features and settings	
	The information was collected from titles and abstracts that were reported in English
edtmann 1995 Clinical features and settings	The information was collected from titles and abstracts that were reported in English
edtmann 1995	The information was collected from titles and abstracts that were reported in English Number of participants eligible: 4172 participants
edtmann 1995 Clinical features and settings	The information was collected from titles and abstracts that were reported in English Number of participants eligible: 4172 participants Number of participants enrolled IT and RS:
edtmann 1995 Clinical features and settings Participants	The information was collected from titles and abstracts that were reported in English Number of participants eligible: 4172 participants
edtmann 1995 Clinical features and settings Participants Study design	Number of participants eligible: 4172 participants Number of participants enrolled IT and RS: - US and surgery: 1227 participants
edtmann 1995 Clinical features and settings Participants	Number of participants eligible: 4172 participants Number of participants enrolled IT and RS: - US and surgery: 1227 participants Target conditions: Presence of any rotator cuff tears
edtmann 1995 Clinical features and settings Participants Study design Target condition and reference standard(s)	Number of participants eligible: 4172 participants Number of participants enrolled IT and RS: - US and surgery: 1227 participants Target conditions: Presence of any rotator cuff tears Reference standard(s): Open surgery
edtmann 1995 Clinical features and settings Participants Study design Target condition and reference	Number of participants eligible: 4172 participants Number of participants enrolled IT and RS: - US and surgery: 1227 participants Target conditions: Presence of any rotator cuff tears
edtmann 1995 Clinical features and settings Participants Study design Target condition and reference standard(s)	Number of participants eligible: 4172 participants Number of participants enrolled IT and RS: - US and surgery: 1227 participants Target conditions: Presence of any rotator cuff tears Reference standard(s): Open surgery
edtmann 1995 Clinical features and settings Participants Study design Target condition and reference standard(s) Index and comparator tests	Number of participants eligible: 4172 participants Number of participants enrolled IT and RS: - US and surgery: 1227 participants Target conditions: Presence of any rotator cuff tears Reference standard(s): Open surgery

Number of participants enrolled IT and RS:

- MRI and surgery: 88 participants



Heininger-Biner 20	(Continued)
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Study	ı de	cισn
Juan	y uc	அதா

Study design		
Target condition and reference standard(s)	Target conditions: Presence of any rotator cuff tears	
	Reference standard(s): Surgery	
	Unclear whether arthroscopy or open surgery	
Index and comparator tests	Index test(s): MRI	
Follow-up		
Notes	Awaiting translation - German article	
	The information was collected from titles and abstracts that were reported in English	

Kayser 2005

Clinical features and settings			
Participants	Number of participants eligible: 239 participants		
	Number of participants enrolled IT and RS:		
	- US and surgery: 239 participants		
Study design			
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears		
	Reference standard(s): Shoulder arthroscopy		
Index and comparator tests	Index test(s): US		
Follow-up			
Notes	Awaiting translation - German article		
	The information was collected from titles and abstracts that were reported in English		

Clinical features and se	ettings	
Participants	Number of participants eligible: 40 participants	
	Number of participants enrolled IT and RS:	
	- US and surgery: 40 participants	
	- MRI and surgery: 40 participants	



Kenn 2000	(Continued)
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Target condition and refer-
ence standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness

tears

Reference standard(s): Surgery

Unclear whether arthroscopy or open surgery

Index and comparator tests	Index test(s): MRI and US
Follow-up	
Notes	Awaiting translation - German article
	The information was collected from titles and abstracts that were reported in English

Kumagai 1991

Participants	Number of participants eligible: 30 participants
Participants	Number of participants eligible: 30 participants

Number of participants enrolled IT and RS:
- MRI and surgery: 30 participants

- MRA and surgery: 30 participants

Study design

Target condition and refer-
ence standard(s)

Clinical features and settings

Target conditions: Presence of any rotator cuff tears

Reference standard(s): Surgery

Unclear whether arthroscopy or open surgery

Index and comparator tests Index test

Index test(s): MRI and MRA

Follow-up

Notes

Awaiting translation - Japanese article

The information was collected from titles and abstracts that were reported in English

Kumagai 1992

Cl	inical	features	and	l settings

Participants

Number of participants eligible: 115 participants

Number of participants enrolled IT and RS:

- MRI and surgery: unclear

Study design



Kumaga	i 1992	(Continued)
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Target condition and reference standard(s)

Target conditions: Presence of any rotator cuff tears, full thickness tears and partial thickness

tears

Reference standard(s): Surgery

Unclear whether arthroscopy or open surgery

Index and comparator tests	Index test(s): MRI
Follow-up	
Notes	Awaiting translation - Japanese article
	The information was collected from titles and abstracts that were reported in English

Kumagai 1995

Clinical features and settings	
Participants	Number of participants eligible: 94 participants
	Number of participants enrolled IT and RS:
	- MRI and surgery: 21 participants
Study design	
Target condition and reference stan-	Target conditions: Presence of full thickness tears
dard(s)	Reference standard(s): Surgery
	Unclear whether arthroscopy or open surgery
Index and comparator tests	Index test(s): MRI
Follow-up	
Notes	Awaiting translation - Japanese article

Labanauskaite 2002

Clinical features and settings	
Participants	Number of participants eligible: 31 participants
	Number of participants enrolled IT and RS:
	- US and surgery: 31 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Shoulder arthroscopy



Index and comparator tests	Index test(s): US
Follow-up	
Notes	Awaiting translation - Lithuanian article
	The information was collected from titles and abstracts that were reported in English

Manych 2007 Clinical featu

Clinical features and settings	
Participants	Number of participants eligible: 275 participants
	Number of participants enrolled IT and RS:
	- MRA and surgery: 197 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Shoulder arthroscopy
Index and comparator tests	Index test(s): MRA
Follow-up	
Notes	Awaiting translation - German article
	The information was collected from titles and abstracts that were reported in English

Nagamori 1995

Clinical features and settings	
Participants	Number of participants eligible: 45 participants
	Number of participants enrolled IT and RS:
	- MRI and surgery: 45 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Surgery
	Unclear whether arthroscopy or open surgery
Index and comparator tests	Index test(s): MRI
Follow-up	



Nagamori 1995 (Continued)

Notes Awaiting translation - Japanese article

The information was collected from titles and abstracts that were reported in English

Qu 2008

Clinical features and settings	
Participants	Number of participants eligible: 57 participants
	Number of participants enrolled IT and RS:
	- MRI and surgery: 57 participants
	- MRA and surgery: 57 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Shoulder arthroscopy
Index and comparator tests	Index test(s): MRI and MRA
Follow-up	
Notes	Awaiting translation - Chinese article
	The information was collected from titles and abstracts that were reported in English

Rudolph 2000

Rudotpii 2000	
Clinical features and settings	
Participants	Number of participants eligible: 63 participants
	Number of participants enrolled IT and RS:
	- MRA and surgery: 32 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Shoulder arthroscopy
Index and comparator tests	Index test(s): MRI
Follow-up	
Notes	Awaiting translation - German article
	The information was collected from titles and abstracts that were reported in English



Sa	kura	gi	19	89

Clinical features and settings	
Participants	Number of participants eligible: unclear number of participants
	Number of participants enrolled IT and RS:
	- US and surgery: unclear number of participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Surgery
	Unclear whether arthroscopy or open surgery
Index and comparator tests	Index test(s): US
Follow-up	
Notes	Awaiting translation - Japanese article
	The information was collected from titles and abstracts that were reported in English

Sasaki 1991

Clinical features and settings	
Participants	Number of participants eligible: 30 participants
	Number of participants enrolled IT and RS:
	- MRI and surgery: 15 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Surgery
	Unclear whether arthroscopy or open surgery
Index and comparator tests	Index test(s): MRI
Follow-up	
Notes	Awaiting translation - Japanese article
	The information was collected from titles and abstracts that were reported in English



Clinical features and settings	
Participants	Number of participants eligible: 30 participants
	Number of participants enrolled IT and RS:
	- US and surgery: unclear number of participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Surgery
	Unclear whether arthroscopy or open surgery
Index and comparator tests	Index test(s): US
Follow-up	
Notes	Awaiting translation - German article
	The information was collected from titles and abstracts that were reported in English

Schroder 2003

Clinical features and settings	
Participants	Number of participants eligible: 80 participants
	Number of participants enrolled IT and RS:
	- MRI and surgery: 80 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Open surgery or shoulder arthroscopy
Index and comparator tests	Index test(s): MRI
Follow-up	
Notes	Awaiting translation - German article
	The information was collected from titles and abstracts that were reported in English

Sell 1992

Clinical features and settings	
Participants	Number of participants eligible: 37 participants



Sell 1992 (Continued)	
	Number of participants enrolled IT and RS:
	- MRI and surgery: unclear number of participants
	- US and surgery: unclear number of participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Open surgery or shoulder arthroscopy
Index and comparator tests	Index test(s): MRI and US
Follow-up	
Notes	Awaiting translation - German article
	The information was collected from titles and abstracts that were reported in English

Sperner 1993

Clinical features and settings	
Participants	Number of participants eligible: 375 participants
	Number of participants enrolled IT and RS:
	- US and surgery: 375 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Open surgery
Index and comparator tests	Index test(s): US
Follow-up	
Notes	Awaiting translation - German article
	The information was collected from titles and abstracts that were reported in English

Vahlensieck 1996

Valitelisieck 1990		
Clinical features and se	ettings	
Participants	Number of participants eligible: 25 participants	
	Number of participants enrolled IT and RS:	
	- MRI and surgery: 25 participants	



ahlensieck 1996 (Continued)	
. ,	- US and surgery: 25 participants
	- MRA and surgery: 25 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Shoulder arthroscopy
Index and comparator tests	Index test(s): MRI, US and MRA
Follow-up	
Notes	Awaiting translation - German article
	The information was collected by titles and abstracts that were reported in English
Nallay 2000	
Vallny 2000 Clinical features and settings	
Participants	Number of participants eligible: 25 participants
	Number of participants enrolled IT and RS:
	- US and surgery: 25 participants
Study design	
Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Surgery
	Unclear whether arthroscopy or open surgery
Index and comparator tests	Index test(s): US
Follow-up	
Notes	Awaiting translation - German article
	The information was collected from titles and abstracts that were reported in English
Vang 2009	
Clinical features and settings	

Number of participants enrolled IT and RS:

- MRA and surgery: 40 participants



Wang 2009 (Continued)

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Study	<i>1</i> d	ACI	σr
Juan	, u	CO	51

Target condition and reference standard(s)	Target conditions : Presence of any rotator cuff tears, full thickness tears and partial thickness tears
	Reference standard(s): Shoulder arthroscopy
Index and comparator tests	Index test(s): MRA
Follow-up	
Notes	Awaiting translation - Chinese article
	The information was collected from titles and abstracts that were reported in English

IT: index test

RS: reference standard

MRI: magnetic resonance imaging MRA: magnetic resonance arthrography

US: ultrasound

DATA

Presented below are all the data for all of the tests entered into the review.

Table Tests. Data tables by test

Test	No. of studies	No. of participants
1 MRA for detection of any rotator cuff tears	3	183
2 MRA for detection of full thickness tears	3	183
3 MRA for detection of partial thickness tears	4	233
4 MRI for detection of any rotator cuff tears	6	347
5 MRI for detection of full thickness tears	7	368
6 MRI for detection of partial thickness tears	6	347
7 US for detection of partial thickness tears	8	660
8 US for detection of full thickness tears	10	729
9 US for detection of any rotator cuff tears	13	854
11 MRA for detection of any subscapularis tendon tears	1	58



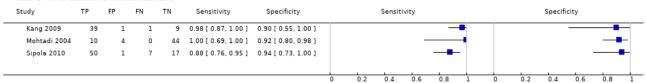
Test 1. MRA for detection of any rotator cuff tears.

Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 1 MRA for detection of any rotator cuff tears

Study		TP	FP	FN	TN	Sensitivity	Specificity			Sensitiv	ity					Specific	ity		
Kang 2	009	45	1	0	4	1.00 [0.92, 1.00]	0.80 [0.28, 0.99]								-			-	
Mohta	di 2004	26	21	10	1	0.72 [0.55, 0.86]	0.05 [0.00, 0.23]			-	-	_		•	_				
Sipola	2010	62	2	2	9	0.97 [0.89, 1.00]	0.82 [0.48, 0.98]					-				_		•	-
								0	0.2	0.4	0.6	0.8	1	0	0.2	0.4	0.6	0.8	1

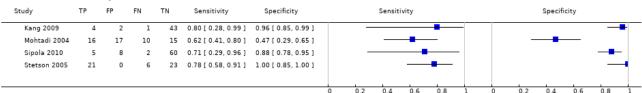
Test 2. MRA for detection of full thickness tears.

Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 2 MRA for detection of full thickness tears



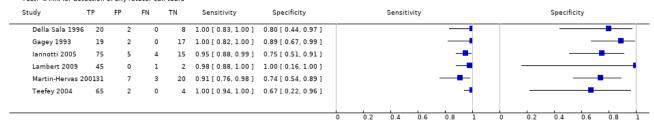
Test 3. MRA for detection of partial thickness tears.

Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 3 MRA for detection of partial thickness tears



Test 4. MRI for detection of any rotator cuff tears.

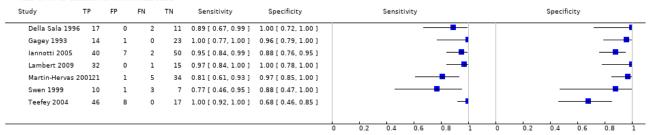
Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 4 MRI for detection of any rotator cuff tears





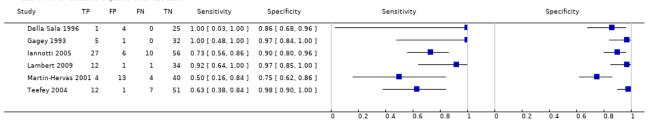
Test 5. MRI for detection of full thickness tears.

Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 5 MRI for detection of full thickness tears



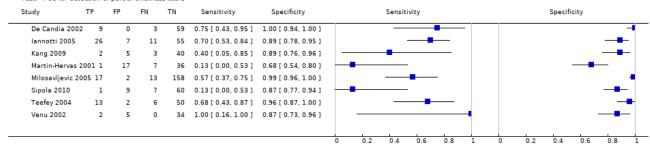
Test 6. MRI for detection of partial thickness tears.

Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 6 MRI for detection of partial thickness tears



Test 7. US for detection of partial thickness tears.

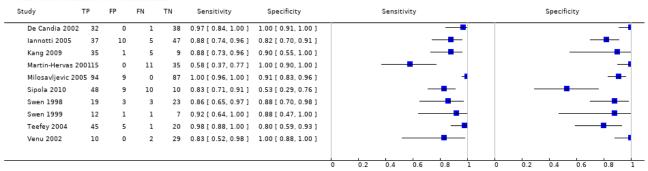
Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 7 US for detection of partial thickness tears





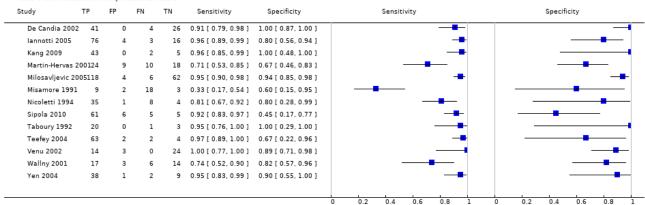
Test 8. US for detection of full thickness tears.

Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 8 US for detection of full thickness tears



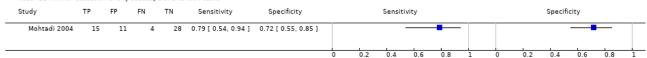
Test 9. US for detection of any rotator cuff tears.

Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 9 US for detection of any rotator cuff tears



Test 11. MRA for detection of any subscapularis tendon tears.

Review: Magnetic resonance imaging, magnetic resonance arthrography and ultrasonography for assessing rotator cuff tears in people with shoulder pain for whom surgery is being considere Test: 11 MRA for detection of any subscapularis tendon tears



ADDITIONAL TABLES

Table 1. Comparison of MRI, US and MRA for detection of any rotator cuff tears (partial or full thickness) using all studies (indirect comparison)

Index test	Studies	Shoulders	Cases	Summary sensitivity (95% CI)	Summary specificity (95% CI)	LR+	LR-	Test ¹
Any rotator	cuff tears							
MRI	6	347	263	98 (92, 99)	79 (68, 87)	5 (2, 10)	0.03 (0.01, 0.11)	P = 0.13
US	13	854	626	91 (83, 95)	85 (74, 92)	6 (3, 12)	0.11 (0.05, 0.22)	
Full thicknes	s tears							
MRI	7	368	193	94 (85, 98)	93 (83, 97)	13 (6, 29)	0.06 (0.02, 0.16)	P = 0.7
MRA	3	183	107	94 (80, 98)	92 (83, 97)	12 (5, 30)	0.06 (0.02, 0.23)	
US	10	729	386	92 (82, 96)	93 (81, 97)	12 (5, 34)	0.09 (0.04, 0.20)	
Partial tears								
MRI	6	347	83	74 (59, 85)	93 (84, 97)	10 (4, 26)	0.28 (0.17, 0.48)	P = 1.00
US	8	660	121	52 (33, 70)	93 (85, 97)	8 (3, 19)	0.52 (0.33, 0.80)	

¹ Likelihood ratio test for evidence of a difference in sensitivity and/or specificity between the tests.

Table 2. Comparison of MRI and US for detection of rotator cuff tears (any, partial or full thickness) limited to studies in which all participants received both MRI and US (direct comparison)

Study	Cases	Non-cases	MRI		US		Difference in	Difference in specificity (95% CI)
			Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)	- sensitivity (95% CI)	
Any rotator cuff	tears							
Iannotti 2005	79	20	95 (88, 99)	75 (51, 91)	96 (89, 99)	80 (56, 94)	-1 (-8, 5)	-5 (-31, 21)
Martin-Hervas 2001	34	27	91 (76, 98)	74 (54, 89)	71 (53, 85)	67 (46, 83)	21 (3, 39)	7 (-17, 32)

LR+ = positive likelihood ratio; LR- = negative likelihood ratio

Table 2. Comparison of MRI and US for detection of rotator cuff tears (any, partial or full thickness) limited to studies in which all participants received both MRI and US (direct comparison) (Continued)

Teefey 2004	65	6	100 (94, 100)	67 (22, 96)	97 (89, 100)	67 (22, 96)	3 (-1, 7)	0 (-53, 53)
Full thickness to	ears							
lannotti 2005	42	57	95 (84, 99)	88 (76, 95)	88 (74, 96)	82 (70, 91)	7 (-5, 19)	5 (-8, 18)
Martin-Hervas 2001	26	35	81 (61, 93)	97 (85, 100)	58 (37, 77)	100 (90, 100)	23 (-1, 47)	-3 (-8, 3)
Swen 1999	13	8	77 (46, 95)	88 (47, 100)	92 (64, 100)	88 (47, 100)	-15 (-42, 12)	0 (-32, 32)
Teefey 2004	46	25	100 (92, 100)	68 (46, 85)	98 (88, 100)	80 (59, 93)	2 (-2, 6)	-12 (-36, 12)
Partial thicknes	s tears							
lannotti 2005	37	62	73 (56, 86)	90 (80, 96)	70 (53, 84)	89 (78, 95)	3 (-18, 23)	2 (-9, 12)
Martin-Hervas 2001	8	53	50 (16, 84)	75 (62, 86)	13 (0, 53)	68 (54, 80)	38 (-4, 79)	8 (-10, 25)
Teefey 2004	19	52	63 (38, 84)	98 (90, 100)	68 (43, 87)	96 (87, 100)	-5 (-35, 25)	2 (-4, 8)

Table 3. Comparison of MRA and US for detection of rotator cuff tears (any, partial or full thickness) limited to studies in which all patients received both MRI and US (direct comparison)

Study	Cases	Non-cases	MRA US¹ Sensitivity (95% Specificity (95% CI) CI) Sensitivity (95% CI) CI)			Difference in sensi- _ tivity (95% CI)	Differ- ence in specificity (95% CI)	
					Specificity (95% CI)	, (55 /5 51)		
Any rotator c	uff tears							
Kang 2009	45	5	100 (92, 100)	80 (28, 99)	96 (85, 99)	100 (48, 100)	4 (-2, 10)	-20 (-55, 15)
Sipola 2010 ²	64	11	97 (89, 100)	82 (48, 98)	92 (83, 97)	45 (17, 77)	4 (-3, 12)	36 (-0.9, 74)
Full thicknes	s tears							

Table 3. Comparison of MRA and US for detection of rotator cuff tears (any, partial or full thickness) limited to studies in which all patients received
both MRI and US (direct comparison) (Continued)

Kang 2009	40	10	97 (87, 100)	90 (55, 100)	88 (73, 96)	90 (55, 100)	10 (-1, 21)	0 (-26, 26)
Sipola 2010 ³	57	18	88 (76, 95)	94 (73, 100)	83 (71, 91)	53 (29, 76)	5 (-8, 18)	42 (17, 67)
Partial thickr	ness tears							
Kang 2009	5	45	80 (28, 99)	96 (85, 99)	40 (5, 85)	89 (76, 96)	40 (-15, 95)	7 (-4, 18)
Sipola 2010 ⁴	7	68	71 (29, 96)	88 (78, 95)	13 (0, 53)	87 (77, 94)	59 (18, 99)	1 (-10, 12)

¹ For the three target conditions, there were 2 additional shoulders for US

² 66 cases for detection of any rotator cuff tears using US

³ 8 cases and 69 non-cases for detection of full thickness tears using US

 $^{^4}$ 58 cases and 19 non-cases for detection of partial thickness tears using US



APPENDICES

Appendix 1. Search strategies

MEDLINE (PubMed)

((Ultrasonography [mh] OR ultrasound [tw] OR ultrasonograph* [tw] OR sonograp*[tw] OR us [sh]) OR (Magnetic Resonance Imaging [mh] OR MR imag*[tw] OR magnetic resonance imag* [tw] OR MRI [tw])) AND (Rotator Cuff [mh] OR rotator cuff* [tw] OR musculotendinous cuff* [tw] OR subscapularis [tw] OR supraspinatus [tw] OR infraspinatus OR teres minor [tw]) AND (Rupture [mh:noexp] OR tear* [tw] OR torn [tw] OR thickness [tw] OR lesion* [tw] OR ruptur* [tw] OR injur* [tw])

Total references = 1551

EMBASE (Elsevier)

1 'echography'/de AND [embase]/lim (124208)

2 ultrasound:ab,ti OR ultrasonograph*:ab,ti OR sonograp*:ab,ti AND [embase]/lim (192495)

3 #1 OR #2 (242499)

4 'nuclear magnetic resonance imaging'/de AND [embase]/lim (277184)

5 (('magnetic resonance' OR mr) NEAR/3 imag*):ab,ti AND [embase]/lim (130882)

6 mri:ab,ti AND [embase]/lim (108797)

7 #4 OR #5 OR #6 (311974)

8 'rotator cuff injury'/de OR 'rotator cuff rupture'/de AND [embase]/lim (3561)

9 'rotator cuff'/de AND [embase]/lim (1850)

10 'rotator cuff':ab,ti OR 'musculotendinous cuff':ab,ti OR subscapularis:ab,ti OR supraspinatus:ab,ti OR infraspinatus:ab,ti OR 'teres minor':ab,ti AND [embase]/lim (5679)

11 #9 OR #10 (6120)

12 'rupture'/de AND [embase]/lim (3798)

13 tear*:ab,ti OR torn:ab,ti OR thickness:ab,ti OR lesion*:ab,ti OR ruptur*:ab,ti OR injur*:ab,ti AND [embase]/lim (1001852)

14 #12 OR #13 (1002130)

15 #11 AND #14 (3615)

16 #8 OR #15 (4908)

17 #3 OR #7 (526691)

18 #16 AND #17 (1572)

LILACS (Bireme)

(Mh Ultrasonography OR Tw ultrasound OR Tw ultrasonograph\$ OR Tw Sonograp\$) OR (Mh Magnetic Resonance Imaging OR (Tw magnetic AND Tw resonance AND Tw imag\$) OR Tw MRI) [Words] and Mh Rotator Cuff OR (Tw rotator AND Tw cuff) OR (Tw musculotendinous AND Tw cuff) OR Tw subscapularis OR Tw supraspinatus OR Tw infraspinatus OR (Tw teres AND Tw minor) [Words] and Mh Rupture OR Tw tear \$ OR Tw torn OR Tw thickness OR Tw lesion\$ OR Tw rupture\$ OR Tw injur\$ [Words]

Total references = 30

Appendix 2. Assessment of methodological quality: QUADAS and additional items

Item definition	Item question	Assessment
Representative spec- trum	Was the spectrum of patients representative of the patients who will receive the test in practice?	Yes: (a) the setting was secondary or tertiary care AND (b) the population was patients with shoulder pain suspected of a rotator cuff tear for whom surgery is being considered AND (c) the study was prospective AND (d) recruitment was consecutive
	tice.	Unclear: if insufficient information was given on the setting, selection criteria, or selection procedure to make a judgment
		No: (a) the setting was primary care OR (b) the population was unselected but defined by shoulder pain OR (c) the study was not prospective OR (d) recruitment was not consecutive
Acceptable reference standard	2. Is the reference stan- dard likely to classify the	Yes: if the reference standard was arthroscopy or a combination of arthroscopy and open surgery (including mini-open)



(Continued)	target condition correct- ly?	Unclear: if the target condition was partial thickness rotator cuff tears and the reference standard was open surgery (including mini-open) No: not applicable
Acceptable delay between tests	3. Is the time period be- tween reference stan- dard and index test short enough to be reasonably sure that the target con- dition did not change be- tween the two tests?	Yes: if the average interval between reference standard and index test was one month or less Unclear: if the interval between tests was not clearly reported No: if the average interval between reference standard and index test was longer than one month
Partial verification avoided	4. Did the whole sample or a random selection of the sample, receive verification using the intended reference standard?	Yes: If all patients who received the index test went on to receive verification of their disease status using a reference standard (Score 'Yes' even if different reference tests were used) Unclear: if insufficient information was given on relation of index test and reference standard No: if not all the patients who received the index test underwent a reference standard to verify their true disease status
Differential verifica- tion bias	5. Did patients receive the same reference stan- dard irrespective of the index test result?	Yes: if all patients received the same reference standard, regardless of the result of their index test Unclear: If it is unclear whether different reference standards were used No: if the result of the index test influenced the choice of the reference standard
Incorporation bias	6. Was the reference standard independent of the index test (i.e. the index test did not form part of the reference standard)?	Should be considered 'Yes' for all studies because the index test is not part of the reference standard
Index test results blinded	7. Were the index test results interpreted without knowledge of the results of the reference standard?	Yes: if the person undertaking the index test was blinded to the results of the standard reference Unclear: if insufficient information was given on independent or blind assessment of the index test No: if the results of the reference tests were known to the person undertaking the index tests
Reference standard results blinded	8. Were the reference standard results inter- preted without knowl- edge of the results of the index test?	Yes: if the reference standard results were performed blind to the results of the index test Unclear: if insufficient information was given on independent or blind assessment of the reference standard No: if the results of the index tests were known to the person interpreting the reference tests
Relevant clinical information	9. Were the same clinical data available when test results were interpreted as would be available when the test is used in practice?	Yes: if clinical data would normally be available when the test is interpreted in practice and similar data were available when interpreting the index test in the study Unclear: if insufficient information was given to explain which clinical information was available at the time of assessment



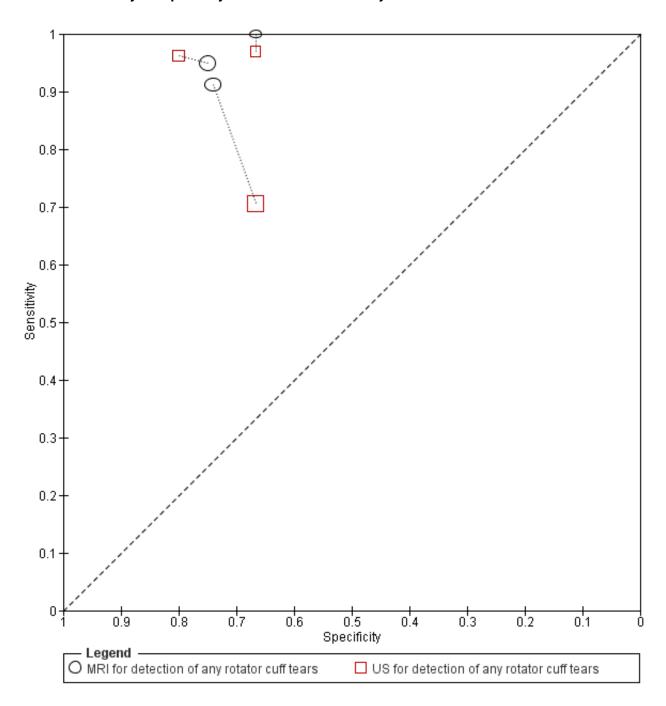
(Continued)		No: if clinical data were not available when index test(s) was(were) interpreted			
Uninterpretable results reported	10. Were uninter- pretable/ intermediate test results reported?	Yes: If the number of uninterpretable test results is stated, or if the number of results reported agrees with the number of patients recruited (indicating no uninterpretable test results)			
		Unclear: if insufficient information was given to permit judgement			
		No: If it states that uninterpretable test results occurred or were excluded and does not report how many			
Withdrawals ex- plained	11. Were withdrawals from the study ex-	Yes: if the number and reasons of all withdrawals from the study were explained (ideally by a flow chart) or if no participants were excluded from the analysis			
	plained?	Unclear: if insufficient information was given on the withdrawals			
		No: if not all withdrawals were explained			
Learning curve / training reported of index test	12. Had index test oper- ators had appropriate training or experience	Yes: (a) if the index test(s) executors were radiologists or shoulder surgeons AND (b) if the tests interpreters had experience in diagnostic of musculoskeletal diseases			
	in musculoskeletal dis- eases?	Unclear: if insufficient information was given to permit judgement			
		No: (a) if the index test(s) executors were not radiologists or shoulder surgeons OR (b) if the tests interpreters had no experience in diagnostic of musculoskeletal diseases			
Learning curve / training reported of	13. Had reference standard test operators had	Yes: (a) if the reference standard(s) executors were shoulder surgeons AND (b) if the results interpreters had experience in shoulder diseases			
reference standard	appropriate training or experience in shoulder	Unclear: if insufficient information was given to permit judgement			
	surgery?	No: (a) if the reference standard(s) executors were not shoulder surgeons OR (b) if the results interpreters had no experience in shoulder diseases			
Index test / criteria	14. Index test criteria for	Yes: (a) if the study provides a clear definition of a positive test result			
for a positive result	a positive result report- ed??	Unclear: if insufficient information was given to permit judgement			
		No: if no definition is given of a positive test result			

Appendix 3. Additional figures

Summary ROC plot of within study comparisons of MRI and US for detection of any rotator cuff tears (Figure 12)



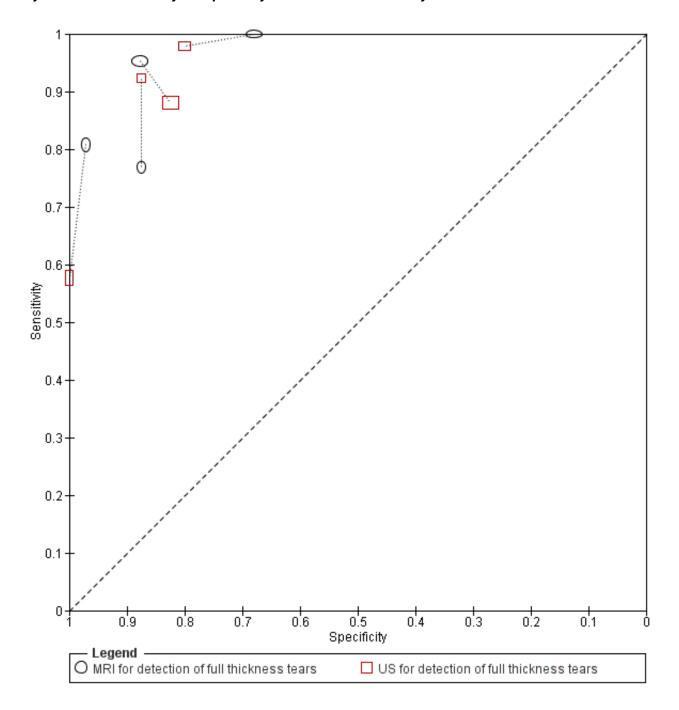
Figure 12. Paired comparison of MRI and US for detection of any rotator cuff tears. Connectling lines link study estimates of sensitivity and specificity for both tests in each study



Summary ROC plot of within study comparisons of MRI and US for detection of full thickness rotator cuff tears (Figure 13)



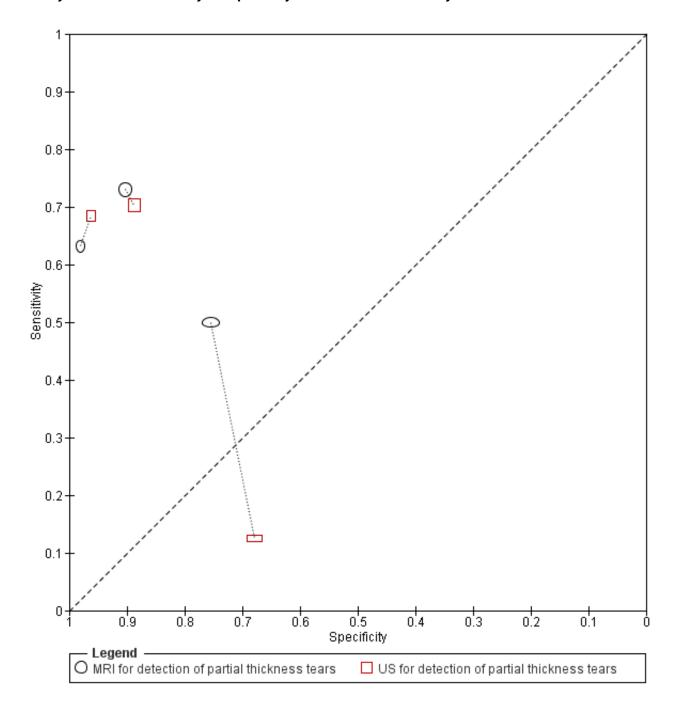
Figure 13. Paired comparison of MRI and US for detection of full thickness rotator cuff tears. Connectling lines link study estimates of sensitivity and specificity for both tests in each study



Summary ROC plot of within study comparisons of MRI and US for detection of partial thickness rotator cuff tears (Figure 14)



Figure 14. Paired comparison of MRI and US for detection of partial thickness rotator cuff tears. Connectling lines link study estimates of sensitivity and specificity for both tests in each study



Appendix 4. Sensitivity analyses for US studies for detection of rotator cuff tears (any, partial or full thickness)

4,11	#
Library	Cochrane

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	Studies	Shoulders	Cases	Summary sensi- tivity (95% CI)	Summary speci- ficity (95% CI)	LR+	LR-
Any rotator cuff tears							
All studies	13	854	626	91 (83, 95)	85 (74, 92)	6 (3, 12)	0.11 (0.05, 0.22)
Acceptable reference standard	5	400	291	94 (88, 97)	91 (82, 95)	10 (5, 22)	0.06 (0.03, 0.14)
Index test results blinded	9	468	348	91 (78, 97)	81 (70, 88)	5 (2, 10)	0.11 (0.04, 0.31)
Full thickness tears						-	
All studies	10	729	386	92 (82, 96)	93 (81, 97)	12 (5, 34)	0.09 (0.04, 0.20)
Acceptable reference standard	6	421	227	95 (86, 98)	91 (85, 95)	11 (6, 20)	0.06 (0.02, 0.16)
Index test results blinded	7	391	201	87 (76, 93)	92 (81, 97)	11 (5, 26)	0.14 (0.08, 0.26)
Partial tears							
All studies	8	660	121	52 (33, 70)	93 (85, 97)	8 (3, 19)	0.52 (0.33, 0.80)
Acceptable reference standard	4	352	56	62 (45, 77)	95 (87, 98)	12 (5, 31)	0.40 (0.26, 0.61)
Index test results blinded	5	322	71	56 (32, 77)	87 (78, 93)	4 (2, 9)	0.51 (0.28, 0.93)



Footnotes

Sensitivity analyses performed by excluding studies that scored 'Unclear' or 'No' for each of the two QUADAS criteria listed in the table.

WHAT'S NEW

Date	Event	Description
3 September 2014	Amended	Republished (September 2014) to include a Plain Language Summary.

CONTRIBUTIONS OF AUTHORS

All authors contributed to the development of the review and commented on and approved the final version. The guarantor of this review is Mario Lenza.

DECLARATIONS OF INTEREST

None known.

SOURCES OF SUPPORT

Internal sources

- Universidade Federal de São Paulo, Brazil.
- The Parker Institute, Denmark.

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- · Teesside University, UK.
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In kind support

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In kind support

External sources

• No sources of support supplied

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

- 1. We only included studies of participants suspected of having rotator cuff tears. Studies that reported results of people who had been previously diagnosed with, or suspected of having, other specific shoulder diagnoses were excluded. If it was unclear whether or not all participants were suspected of having rotator cuff tears, we also excluded these studies
- 2. Inasmuch as there is no set time point beyond which it is known that rotator cuff tears progress, we accepted studies in which the time between the index test and the reference standard test was up to a year (rather than six months as specified in the protocol).
- 3. We included the MEDION database in our search strategy.
- 4. We restricted our analyses to prospective studies and excluded retrospective studies because of the high risk of spectrum and verification biases in these studies.
- 5. We made an amendment in the assessment of methodological quality item seven (index test results blinded). We removed "if the study was retrospective" as a reason to say *No* because we included only prospective studies.
- 6. We made an amendment in the assessment of methodological quality item eight (reference standard results blinded). We excluded "if the study was retrospective" as a reason to say *No* because it was covered by the first part of the sentence.
- 7. We included in the assessment of methodological quality table an additional generic quality item assessing whether or not the criteria for a positive index test result was reported.
- 8. We used the bivariate model for meta-analysis instead of the hierarchical summary ROC (HSROC) model. Given the available information, we assumed a common threshold was applicable but with heterogeneity around this common threshold due to variation



in interpretation in practice. Therefore we consider the bivariate model and the estimation of summary points (with 95% confidence regions) appropriate for summarising the results of the review.

9. We conducted sensitivity analyses to examine the effect of unit of analysis.

INDEX TERMS

Medical Subject Headings (MeSH)

*Arthroscopy; *Magnetic Resonance Imaging; *Rotator Cuff Injuries; *Ultrasonography; Arthrography [*methods]; Prospective Studies; Rotator Cuff [surgery]; Shoulder Pain [*etiology] [surgery]

MeSH check words

Humans