

Systematic Review of MRI, Ultrasound, and Scintigraphy as Outcome Measures for Structural Pathology in Interventional Therapeutic Studies of Knee Arthritis: Focus on Responsiveness

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ABSTRACT. Objective. Validated imaging outcome tools to assess response to therapies in a single joint are required. Our aim was to review the published literature to ascertain the responsiveness of novel imaging techniques as outcome measures in interventional therapeutic studies of knee arthritis.

Methods. An Ovid Medline search was performed for original articles in English that used imaging techniques to assess response at the knee joint to therapy in osteoarthritis, rheumatoid arthritis, and psoriatic arthritis. Changes in response to therapy were assessed with regard to both internal and external responsiveness.

Results. In the studies that presented appropriate statistical data to allow responsiveness to be assessed, MRI was generally found to be internally responsive to pathologies imaged, and externally responsive, referenced against both other imaging modalities and biochemical biomarkers of arthritis. Ultrasonography was found to demonstrate internal responsiveness with regard to synovial thickness, effusion size, and popliteal cyst size. External responsiveness was demonstrated against several referenced health status measures. Scintigraphy was found to be externally responsive in the majority of studies, with internal responsiveness demonstrated in 1 study.

Conclusion. While the imaging techniques appear to be responsive from the data we present, further inspection reveals that interpreting the responsiveness of imaging techniques was difficult, largely because of a lack of standardization of image acquisition, definitions of pathology, and scoring systems. Refined pathological definitions and scoring systems are required to enable the development of valid and responsive tools for interventional clinical trials. (J Rheumatol First Release Oct 1 2010; doi:10.3899/jrheum.100377)

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ULTRASONOGRAPHY MAGNETIC RESONANCE IMAGING SCINTIGRAPHY
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Assessment of the efficacy of therapies in the clinical trial setting relies on the use of outcome tools to define improvement. Additionally in the clinical setting, application of objective measurement tools to guide management results in

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better outcomes¹. However, the development and application of such tools in clinical trials in rheumatology is relatively recent, and for many rheumatic conditions, appropriately validated tools are not currently available. Improvement of outcome measurement in rheumatology clinical trials is the focus of the Outcome Measures in Rheumatology Clinical Trials group (OMERACT), an informal network of working groups. OMERACT's agenda is to establish validated, objective, and feasible measurement tools that demonstrate truth, discrimination, and feasibility (the OMERACT filter)². Application of this filter allows an outcome tool to be assessed regarding metric properties such as face, content, construct, and criterion validity, reproducibility, and responsiveness.

Many of the validated outcome measures in rheumatology clinical trials [such as the Disease Activity Score (DAS) and the American College of Rheumatology (ACR) responder criteria] consist of a composite of domains assessing systemic response to therapies. These outcome tools have

been invaluable in establishing the efficacy of pharmacotherapeutics in rheumatology, and aided the advancement of therapies and treatment paradigms, such that the treatment and outcome of many rheumatic conditions is now much improved. However, it has been recognized that these outcome tools may have limited use in evaluating outcomes at the single-joint level, for example when assessing localized intraarticular (IA) therapy³ in the setting of systemic arthritis with only a few refractory joints, and in pauciarticular processes such as monoarthritis. Validated, objective outcome tools are required in order to move the field of rheumatology forward in these situations.

Recently, the OMERACT Single Joint Working Group reviewed publications evaluating clinical, radiographic, and functional assessments of single joints, focusing on the knee³. Our aim was to examine novel, nonradiographic imaging techniques and to systematically assess the published literature for evidence of their responsiveness as outcome tools for arthritis of the knee joint.

MATERIALS AND METHODS

A systematic literature review was undertaken with the objective of identifying studies that used imaging outcome tools to measure response to interventional therapies in arthritis of the knee joint, limited to cohorts with a diagnosis of rheumatoid arthritis (RA), osteoarthritis (OA), or psoriatic arthritis (PsA). Therefore the inclusion criteria were published studies in English, assessing humans *in vivo*, comparing imaging of structural tissue pathology of the knee joint before and after a specific therapeutic intervention. Studies were excluded if they measured only biomechanical pathology, such as abnormal joint alignment. Studies were also excluded if they imaged multiple joints, and data regarding the knee specifically could not be extracted, or they were not the first report. The quality of the trial methodology or reporting was not an inclusion or exclusion criterion. An Ovid Medline search was conducted to identify articles published between 1950 and May 2010 using such search terms as rheumatoid arthritis; arthritis; osteoarthritis, hip; osteoarthritis; osteoarthritis, knee; psoriatic arthritis; ultrasonography, Doppler; magnetic resonance imaging; whole body imaging; diagnostic imaging; radionuclide imaging; three-dimensional; scintigraphy; tomography; x-ray computed or computer tomography; and optical coherence tomography. The titles and abstracts were reviewed to exclude duplicates and identify those articles meeting inclusion criteria. The articles were reviewed and data extracted and inserted into a template based on the previous work of the OMERACT Single Joint Assessment Working Group, but modified for our review³. The extracted data addressed aspects of study design and size, imaging modality and specifications, and pathology imaged, including definitions, scoring system, and change in response to interventional therapies to assess responsiveness, including the statistical method used.

For the purpose of this review, we focused on internal responsiveness and external responsiveness. Internal responsiveness was defined as the ability of an outcome tool to demonstrate temporal changes in response to therapy. Statistical tests considered appropriate for determining responsiveness included the paired t test, standardized response mean (SRM), standardized effect size (SES), and Gyatt's responsiveness index (GRI)⁴. For our review, if studies used SRM, SEM, or GRI to assess responsiveness, the tool was considered responsive if the result was ≥ 0.2 (considered a small response)⁴. External responsiveness is the extent to which changes in an outcome tool correlate with other referenced measurements. Appropriate statistical tests to assess external responsiveness include the receiver-operating characteristic method, correlation coefficients, and regression analysis⁴.

RESULTS

The Medline search identified 5824 published articles, 77 of which were duplicates. Articles having the word "knee" in the title, abstract, or notes numbered 1148, and these were reviewed to determine whether they met the inclusion criteria. A further 6 articles were identified by consulting experts and examining reference lists of relevant review articles. We included 49 articles in our review. Reasons for the exclusion of articles appear in Figure 1.

Individual study designs, including the disease studied, cohort size, and imaging techniques used, are summarized in Table 1. Therapies studied were predominantly IA, including corticosteroids, radiosynovectomy, or hyaluronic acid; oral therapies included methotrexate, leflunomide, corticosteroids, or nonsteroidal antiinflammatory drugs; parenteral therapies included infliximab, etanercept, and corticosteroids; surgery ranged from synovectomy to wedge osteotomy; and physical therapy included repetitive short-wave diathermy.

Scoring systems for knee arthritis were examined in magnetic resonance imaging (MRI) studies (Table 2), ultrasound studies (Table 3), scintigraphic studies (Table 4), and 1 thermography study; no studies were included that used computed tomography or optical coherence tomography.

MRI studies generally provided adequate details of the machine settings, including slice thickness (Table 2). Changes in machine settings and the use of contrast agents occurred in relation to when the study was undertaken. Joint positioning was rarely described in MRI studies. Descriptions of the imaging appearance of pathology in MRI studies were rarely given. Scoring systems varied greatly; some studies scored individual pathologies. Additionally, some studies used software to calculate scores based on cross-sectional area or volume of the imaged tissue or pathology, tissue thickness, or tissue interface, while others used physician readers to score according to dichotomous or semiquantitative systems. Internal responsiveness of MRI to changes in the following pathologies was found in at least 1 article: synovial thickness, enhancement, effusion, bone marrow lesions, cartilage volume and morphology, osteophytes, and quadriceps volume. External responsiveness was rarely studied, but MRI demonstrated external responsiveness in 2 of the articles, referenced against both other imaging modalities and biochemical biomarkers of arthritis.

The ultrasound studies (Table 3) demonstrated a large amount of heterogeneity. While most studies adequately described the machine and machine settings, there was variability in the settings, for example the pulse repetition frequency, wall filter, and Doppler settings (when used) were not uniform. The position of the knee during image acquisition was specified in more than half the studies, although the knee position again was not uniform. Knee positioning included flexion, neutral, and extension, with 1 author using

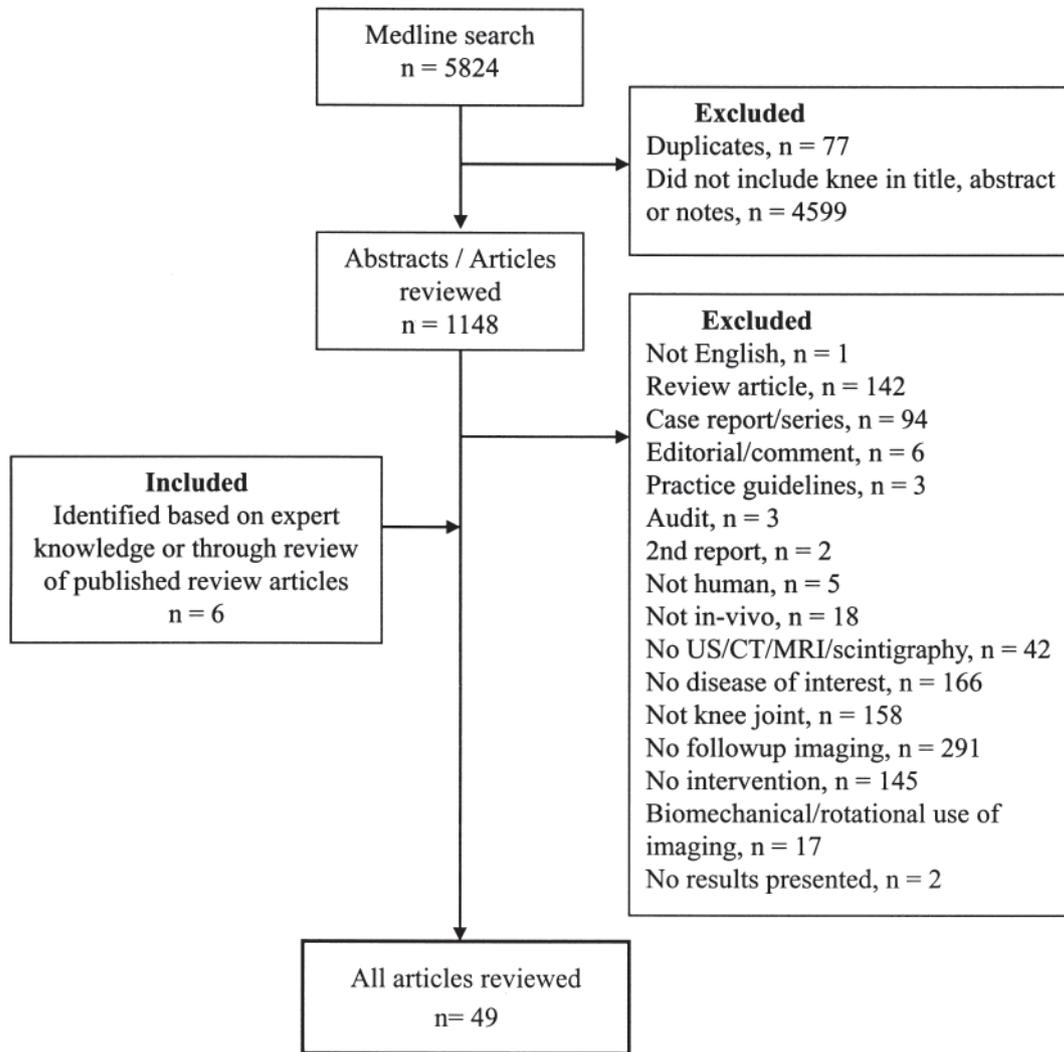


Figure 1. Search strategy and reasons for excluding trials from the review — assesses external responsiveness only.

an extended knee with contracted quadriceps in 1 article, and relaxed quadriceps in another. The definition of the imaging appearance of pathology was described in fewer than half the articles, and was seemingly unrelated to the year of publication. Definitions were not uniform, nor were they in keeping with OMERACT definitions of ultrasound-defined pathology⁵. Scoring systems varied between being descriptive of morphology, dichotomous, semiquantitative (0–3 scale), quantitative measurements (such as tissue thickness), or summative quantitative systems (adding regional tissue thickness measurements). Ultrasonography was found to demonstrate internal responsiveness with regard to synovial thickness, effusion size, and popliteal cyst size. External responsiveness was demonstrated against several referenced health status measures.

Scintigraphic studies are presented in Table 4. Technetium was the most common labeling agent, but other agents were used. Almost a third of articles did not describe

the scoring systems, and when specified, they included semiquantitative systems or computer-generated scoring systems. Scintigraphy was found to be externally responsive in the majority of articles, but internal responsiveness was examined in only 1 article.

DISCUSSION

In our systematic review we identified only 49 articles using MRI, ultrasound, and scintigraphy to assess change in response to therapies in arthritis of the knee joint. Studies assessing the responsiveness of MRI were the most prevalent, and they reported MRI to be both internally and externally responsive to many pathological features of knee arthritis, including synovial thickness, enhancement, effusion, bone marrow lesions, cartilage volume and morphology, osteophytes, and quadriceps volume. Three reports assessed the internal responsiveness of ultrasound, finding it responsive to changes in synovial thickness, effusion, and

Table 1. Summary of studies identified meeting inclusion criteria, including study design, disease studied, imaging modality, and route of therapy.

Study	Study Design				No. Patients Disease			Imaging			Th	Therapy
	Powered	Controlled	Randomized	Blinded	RA	PsA	OA	MRI	US	Sc		
Acebes ⁸ 2006	—	—	—	—			30		Y			IA ST
Al-Janabi ⁹ 1988	—	—	—	—	7					Y		IA ST
Al-Janabi ¹⁰ 1992	—	—	—	Single	11					Y		IA ST
Alonso-Ruiz ¹¹ 1998	—	—	—	—	10			Y				IA HA or placebo
Bagge ¹² 1996	—	Y	Y	Single	11			Y				IA IgG or IA saline
Batalov ¹³ 1999	—	—	—	—	24				Y			Arthroscopic synovectomy
Beckers ¹⁴ 2006	—	—	—	—	16			Y	Y	Y		Steroids or NSAID
Brandt ¹⁵ 2006	—	—	—	Single			30	Y				NSAID or paracetamol
Creamer ¹⁶ 1994	—	Y	Y	Single			12	Y		Y		IA HA
Creamer ¹⁷ 1997	—	Y	—	Single	16			Y				Aspiration and IA ST ± yttrium synovectomy
Cubukcu ¹⁸ 2005	—	Y	Y	—			30	Y				IA HA or placebo
De Bois ¹⁹ 1993	—	—	—	Single	7					Y		IA ST
Fiocco ²⁰ 1996	—	Y	—	Single	12	11			Y			Surgical synovectomy
Fiocco ²¹ 2005	—	—	—	—	12	8			Y			SC etanercept 25 mg twice weekly
Forslind ²² 2004	—	—	—	Single	20			Y				Oral prednisolone and/or various DMARD*
Arzu Gencoglu ²³ 2003	—	—	—	—	23					Y		Y-90 silicate radionuclide synovectomy
Arzu Gencoglu ²⁴ 2002	—	—	—	—	15					Y		Sulindac
Glaser ²⁵ 2007	—	—	—	—			21	Y				Autologous chondrocyte implant
Hunter ²⁶ 2006	Y	Y	Y	Double			150	Y				Oral experimental therapy (NI-15713)
Iagnocco ²⁷ 2006	—	—	—	Single	10	13			Y			IA MTX weekly for 8 wks, then oral MTX
Jan ²⁸ 2006	—	Y	—	—			36		Y			Repetitive shortwave diathermy
Jones ²⁹ 1991	—	—	—	Single	13		9			Y		IA ST
Kroner ³⁰ 2007	—	—	—	Single			20	Y				Lateral closing wedge osteotomy
Lee ³¹ 2003	—	—	—	—	14			Y				IA holium-166-chitosan complex
Leitch ³² 1996	—	—	—	Single	6			Y				Arthrocentesis and IA ST
Marzo-Ortega ³³ 2007	—	—	—	Single	6			Y				Infliximab
Mizner ³⁴ 2005	—	—	—	—	20			Y				Tricompartmental knee arthroplasty
Moore ³⁵ 1975	—	—	—	—	17				Y			IA steroid and local anesthetic or local anesthetic or synovectomy [†]
Newman ³⁶ 1996	—	—	—	—	5	1			Y			Arthrocentesis and IA ST
Ostergaard ³⁷ 1996	—	—	—	—	15	2	1	Y				Aspirate to dryness and IA methylprednisolone 80 mg*
Ozturk ³⁸ 2006	—	Y	Y	Single	47			Y				IA HA and IA steroid or IA HA
Reece ³⁹ 2002	—	Y	Y	Double	39			Y				MTX or leflunomide
Rubaltelli ⁴⁰ 1994	—	—	—	—	12	13			Y			Arthroscopic synovectomy
Salaffi ⁴¹ 2004	—	—	—	Single	18				Y			IA ST
Sharma ⁴² 1999	—	Y	Y	Double			49	Y				Nimesulide or piroxicam
Shin ⁴³ 2007	—	Y	—	Single	16			Y				IA 188 Re-tin colloid 3 doses
Shio ⁴⁴ 2006	—	—	—	—	10				Y			IV Infliximab
Soroo ⁴⁵ 2005	—	Y	—	Single	12					Y		IA steroid or IA P-32 Colloid
Tannenbaum ⁴⁶ 1987	—	—	—	Single	15					Y		Sulindac
Uematsu ⁴⁷ 1979	—	—	—	—	14					Y		High tibial osteotomy
Van Holsbeeck ⁴⁸ 1988	—	—	—	—	20				Y	Y	Y	IA ST
Veale ⁴⁹ 1999	Y	Y	Y	Single	13			Y				Arthroscopy and biopsy followed by IA anti-CD4

Table 1. Continued.

Study	Study Design				No. Patients Disease			Imaging			Th	Therapy
	Powered	Controlled	Randomized	Blinded	RA	PsA	OA	MRI	US	Sc		
Wluka ⁵⁰ 2002	Y	Y	Y	Double			136	Y				Vitamin E
Youssef ⁵¹ 1996	—	—	—	—	9					Y		IV methylprednisolone
Song ⁵² 2009	—	Y	Y	Double			41	Y	Y			IA bradykinin receptor 2 antagonist
Raynaud ⁵³ 2009	Y	Y	Y	Double			355	Y				Licofelone or naproxen
Raynaud ⁵⁴ 2008	U	U	U	U			107	Y				Bisphosphonate or NSAID or COX-2 inhibitor
Raynaud ⁵⁵ 2008	Y	Y	Y	Double			355	Y				Licofelone or naproxen
Anandacoomarasamy ⁵⁶ 2008	Y	—	—	—			19	Y				IA Hylan

* Variety of oral therapies also used. † One subject had surgical therapy. RA: rheumatoid arthritis; PsA: psoriatic arthritis; OA: osteoarthritis; MRI: magnetic resonance imaging; US: ultrasonography; Sc: scintigraphy; Th: thermography; IA: intraarticular; IV: intravenous; HA: hyaluronic acid; ST: corticosteroid; P: parenteral; PH: physical; SC: subcutaneous; DMARD: disease-modifying antirheumatic drug; MTX: methotrexate; NSAID: nonsteroidal antiinflammatory drug; COX-2: cyclooxygenase-2; U: unknown; Y: Yes.

popliteal cysts size, with marked variability in the definitions of pathology, image acquisition, and scoring system used. The external responsiveness of ultrasonography was also assessed in 4 reports, finding correlations between ultrasound and externally referenced health status measures. Scintigraphy was found to be externally responsive in the majority of articles, with little investigation into its internal responsiveness.

While the imaging techniques appear to be responsive from the data, further inspection reveals that interpreting the responsiveness of imaging techniques was difficult for many reasons. Challenges in assessing the responsiveness of these imaging techniques focused on the image acquisition and analysis. These included a lack of standardization in acquiring images, a lack of standard definitions of pathology, and a lack of standardized scoring systems. It is inevitable that variability will be associated with imaging studies, as advances in technology result in improvements in machine capabilities and techniques, such as the addition of contrast agents. Additionally, scientific progress, such as advances by OMERACT in creating standard definitions of ultrasound imaged pathology, will influence study methodology and publication style. However, based on the current published literature, it is difficult to identify which imaging techniques fulfill the OMERACT filter with regard to discrimination; and the variability demonstrated in machine settings, image acquisition, definitions of pathology, and scoring systems makes comparisons between articles difficult.

This systematic review has limitations. We focused on 1 joint only (the knee), and importantly we excluded observational studies in favor of those looking at interventional effects. Other groups have looked at responsiveness for MRI cartilage measures in OA knee using longitudinal studies^{6,7}. Adequately assessing the responsiveness of a tool is dependent on using effective therapies. If a study uses a therapy

that is unable to bring about change in health status, then the responsiveness of the tool may be underestimated⁴. Similarly, if a study examines a subgroup that is likely to respond excessively well to an outcome, then the responsiveness of the tool is likely to be overestimated. In our review, no analysis of the effectiveness of the intervention, or the likelihood of the treatment cohort to respond, was made. We were only able to assess the responsiveness of MRI, ultrasound, and scintigraphy regarding studies that reported appropriate measures and it was not possible to calculate SRM from many studies.

The imaging modalities we studied are well placed to be used as objective outcome tools in knee arthritis, but more work is required before these modalities can be considered to fulfill the OMERACT filter of truth, discrimination, and feasibility as outcomes tools in both inflammatory arthritis and OA of the knee. Our review has identified insufficiencies in the principles of discrimination. Our review highlights greater issues that need further consideration before outcome tools for clinical trials evaluating therapies at the knee joint can be developed. First, whether a generic (as distinct from disease-specific) outcome tool is needed must be carefully considered. Second, the pathologies to be imaged need to be identified and their imaging appearances defined. Image acquisition must be standardized for each imaging modality, including the machine setting, fields of view, and joint positions, although it is recognized that this will change with emerging technology. We recommend that research focus on these issues, and on developing international consensus guidelines for the use of modern imaging techniques in rheumatology clinical trials. Once these basic considerations have been addressed, further work assessing the performance metrics of the tool can be undertaken. Until validated responsive imaging outcome tools are developed, the ability to rigorously examine the mode of action and efficacy of therapies on single joints *in vivo* is limited.

Table 2. Summary of the magnetic resonance imaging (MRI) studies, including pathology scoring system, and responsiveness.

Study	RA	PsA	OA	Gd-DTPA	Pathology	Scoring System	Response	Statistical Assessment Used to Assess Internal Responsiveness?	Internally Responsive?	Statistical Assessment Used to Assess External Responsiveness?	Externally Responsive?
Alonso-Ruiz ¹¹	Y			Y	Synovium	Synovial thickness (mm) in supra patella pouch Synovial enhancement (%) software	MRI demonstrated a reduction in synovial thickness, at all timepoints Enhancement significantly reduced at followup from baseline	Y	Y	N	U/A
Anandaco-omarasamy ⁵⁶			Y		Cartilage volume Cartilage defects	Manual segmentation Semiquantitative score 0–4	No change in cartilage volume of cartilage defect scores	N	N	N	U/A
Bagge ¹²	Y			Y	Synovitis/pannus Hydrops Baker cyst	Unclear	MRI improved in 2/6 receiving IgG, worsened in 1/6. MRI findings improved in 3/5 receiving saline, and worsened in none	N	U/A	N	U/A
Beckers ¹⁴	Y			Y	Synovitis	Relative enhancement, and rate of early enhancement and static enhancement	At 4 weeks, significant correlation between changes in PET, MRI, CRP, MMP-3, but not ultrasound			Y	Y
Brandt ¹⁵			Y	Y	Synovitis effusion	Computer-aided assessment of volume	Both treatment groups showed a reduction in mean total effusion after therapy (p < 0.009 and p < 0.01)	Y	Y	N	U/A
Creamer ¹⁶			Y		Pannus, effusion, enhancement	Pannus, effusion and enhancement score with both semiquantitative scoring system (0–2) and computer-aided quantification	No MRI difference between baseline and followup	N	U/A	N	U/A
Creamer ⁵⁷			Y		Effusion Synovial thickening	Change in effusion volume Synovial thickening unspecified	IA ST cohort: change in fluid volume calculated by computer strongly correlated with aspirated volume. Significant reduction in MRI synovial volume and enhancement by both methods at 1 week. No change in pannus by either method. Synovectomy cohort: there was no change in pannus, effusion, or enhancement by MRI by either scoring system	N	U/A	N	U/A
Cubukcu ¹⁸			Y	Y	Patella cartilage	Modified Shahriaree classification (0–4)	Significant improvement in cartilage grade in the treatment group, but not the placebo group. No change in other MRI features.	N	U/A	N	U/A
Forslind ²²	Y			Y	Synovitis Erosions Edema	Sum of 0–3 in 4 compartments Number and location Presence	All but 1 subject with baseline synovitis had synovitis at followup. No. of subjects with erosions increased from 5 to 11 at followup. No. of subjects with edema progressed with time from 1 to 4	N	U/A	N	U/A
Glaser ²⁵			Y		Cartilage	Software to determine volume, mean and local cartilage thickness, cartilage bone interface, and volume normalized to size of cartilage	A mean increase in cartilage volume and thickness of 6% was observed in the treatment group, p < 0.001	Y	Y	N	U/A

Table 2. Continued.

Study	RA	PsA	OA	Gd-DTPA	Pathology	Scoring System	Response	Statistical Assessment Used to Assess Internal Responsiveness?	Internally Responsive?	Statistical Assessment Used to Assess External Responsiveness?	Externally Responsive?
Hunter ²⁶			Y		Cartilage volume Bone edema, attrition, cysts Osteophytes Synovial cavity distension	WORMS	Small changes in scores for cartilage morphology, synovitis, and osteophytes, with a trend toward differences between treatment and placebo groups	Y	Y	N	U/A
Kroner ³⁰			Y		Bone marrow edema	Bone marrow edema volume calculated, and and regions in medial tibial and femoral compartments summed	Mean size of bone marrow edema lesion decreased from 4.21 cm ³ to 1.2 cm ³ at 1 year, and was maintained at 7 years. JOA score improved at 1 year but decreased slightly at 7 years	Y	Y	N	U/A
Lee ³¹	Y			Y	Synovial enhancement Synovial thickness Baker cyst Erosions	Software (scion image) calculated volume Synovial thickness measured in SPP Effusion calculated using scion image volume in mm ³ Baker cyst volume Number of erosions	Synovial enhancement and thickness and Baker cysts volume did not significantly decrease, although effusion did. Erosion scores did not change	Y	Y (effusion only)	Y	N
Leitch ³²	Y			Y	Effusion Synovium	Effusions scored 0–4 Synovial thickness measured in mm	Reduction in effusion volume by semiquantitative score in 4 patients. Reduction in synovial thickness, most striking in those with thickest synovium at baseline, the rest had minimal changes in fluid and synovium	N	U/A	N	U/A
Marzo-Ontega ³³		Y		Y	Bone marrow edema Synovium	Semiquantitative score 0–3 for bone marrow edema Synovial enhancement volume calculated using software	Bone edema resolved in 1, was unchanged in 1. Reduction in synovial enhancement volumes in 4/6	N	U/A	N	U/A
Mizner ³⁴			Y		Quads volume	Cross-sectional area	Quads volume decreased as did strength and voluntary activation	Internal	Y	Y	Y
Ostergaard ³⁷	Y	Y	Y	Y	Synovitis	Relative enhancement, rate of early enhancement	All knees demonstrated a decrease in rate of early enhancement at 7 days. All knees that relapsed demonstrated an increase in rate of early enhancement at relapse. Knees in clinical remission at day 30 have a decrease in rate of early enhancement compared to baseline, but most of these had increased since day 7.	N	U/A	N	U/A
Ozturk ³⁸			Y		Marrow edema Cartilage thinning Baker cyst Effusion Subchondral cysts Erosions	Global score 1–8 (w/edema) Cartilage thinning Baker cyst Effusion Subchondral cysts Erosions	No significant progression in MRI scores for either group	N	U/N	N	U/A

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Table 2. Continued.

Study	RA	PsA	OA	Gd-DTPA	Pathology	Scoring System	Response	Statistical Assessment Used to Assess Internal Responsiveness?	Internally Responsive?	Statistical Assessment Used to Assess External Responsiveness?	Externally Responsive?
Raynauld ⁵⁴			Y		Subchondral bone changes Meniscal lesions	Identified as edema or cyst, then measured in mm Described elsewhere	No change in bone lesions Unclear	Y (bone lesions only)	N	N	U/A
Raynauld ⁵⁵			Y		Cartilage volume and mean thickness Meniscal extrusion	3-D coordinate system and Euclidean distance between bone to cartilage interface and cartilage to soft tissue interface Extrusion scored on 0–2 semiquantitative scale	Less cartilage loss in the licofelone group	N	U/A	N	U/A
Raynauld ⁵³			Y		Cartilage volume	Unclear	Significantly less cartilage loss in the licofelone group	Y	Y	N	U/A
Reece ³⁹	Y			Y	Synovial inflammation	The rate of enhancement and maximal enhancement	Significant improvement in rate of enhancement in leflunomide group compared to MTX group. No difference between the groups with regard to a decrease in maximal enhancement that improved with therapy	N	U/A	N	U/A
Sharma ⁴²			Y		Cartilage thickness	Cartilage thickness (0–4)	No change over 24 weeks in cartilage grades. OA Severity Index, tenderness and swelling significantly improved at 8 and 24 weeks in the patients that completed the timepoints (n = 49 and 11), with no difference between treatment groups	U/C	U/A	N	U/A
Shin ⁴³	Y			Y	Synovitis Effusion	Change in synovitis scored from –2 to 2, Sum of mm in SPP and IPP in the sagittal plane Change in effusions scored from –2 to 2	Synovial thickening improved in treated knee in 87.5% and effusion decreased in 43.7%. Synovial thickness decreased at 6 months (by 1.68 mm), but slightly increased in control knees (–0.18 mm)	N	U/A	N	U/A
Song ⁵²			Y	Y	Contrast medium enhancement Effusion	Score 0–3 based on measurement in mm (< 5 mm, < 8 mm, < 11 mm, ≥ 11 mm) Semiquantitative score 0–3	Reduction in effusion in the high-dose treatment group. At followup, MRI correlated with ultrasound	U/C	U/A	N	U/A
Veale ⁴⁹	Y			Y	Synovitis	Synovial maximal rate of enhancement, and maximal enhancement	MRI demonstrated a trend toward a dose effect for improvement in synovial measurements with therapy (nonsignificant)	N	U/A	N	U/A
Wluka ⁵⁰			Y		Cartilage volume	Unclear	Loss of cartilage from baseline (similar volume loss in both cohorts)	N	U/A	N	U/A

RA: rheumatoid arthritis; PsA: psoriatic arthritis; OA: osteoarthritis; U/A: unable to assess; PET: positron emission tomography; CRP: C-reactive protein; MMP: matrix metalloproteinase; IA ST: intraarticular corticosteroid; WOMBS: whole-organ magnetic resonance imaging score; JOA: Japanese Orthopaedic Association; SPP: suprapatellar pouch; IPP: intrapatellar pouch. Y: yes; N: no.

Table 3. Summary of the ultrasound studies, including pathology studied, scoring system used, and responsiveness.

Study	Disease			Knee Position	Regions Imaged	Definition of Pathology Imaged	Definition of Scoring System Used	Description of Response	Statistical Assessment Used to Assess Internal Responsiveness?	Internally Responsive?	Statistical Assessment Used to Assess External Responsiveness?	Externally Responsive?
	RA	PsA	OA									
Acebes ⁸			Y		Popliteal fossa		Popliteal cyst synovial wall thickness in mm, popliteal cyst wall area using built-in software	Popliteal cyst size and wall thickness decreased after therapy	Y	Y	Y	Y
Batalov ¹³	Y				SPP		Maximal synovial thickening in the SPP (anterior and posterior wall) in the AP diameter in mm. Effusion depth assessed as the maximal AP thickness of the suprapatella sac in mm	Mean ultrasound synovitis thickness and effusion significantly reduced at 3 and 12 months posttherapy	N	U/A	N	U/A
Song ⁵²			Y		SPP, LR		Effusion 0–3 based on measurements in mm, synovial hypertrophy 0–3 based on measurements in mm;	No changes in the treatment group. At followup, ultrasound demonstrated correlations with symptoms and MRI	U/C	U/A	N	U/A
Beckers ¹⁴	Y			Extended	SPP, MR LR (a)	Y**	PD semiquantitative scale 0–3 Contrast enhancement semiquantitative score 0–3 using slope values synovial thickening present when the sum of the anterior and posterior walls of the pouch measured > 1 mm. Power Doppler signal assessed on semi-quantitative scale 0–3	At 4 weeks, significant correlation between changes in PET, MRI, CRP, and MMP-3, but not ultrasound	N	N	Y	N
Fiocco ²⁰	Y			Extended (d)	SPP, MR LR (b)	Y*	Synovial thickening assessed, as highest score of the SPP or MR or LR as assessed in mm. Pouch thickness measured in mm, then graded 0–3, morphology noted	Ultrasound synovial thickness and effusion decreased significantly at 2 and 12 months. Morphological pattern did not change after synovectomy	Y	Y	N	U/A
Fiocco ²¹	Y	Y		Extended (e)	SPP, MR LR (b)	Y*	Synovial thickening assessed, as highest score of the SPP or MR or LR assessed in mm. Power Doppler graded from 0–3	Ultrasound variables decreased at 12 months, as did all clinical measures of disease activity (except global health score)	N	U/A	N	U/A
Iagnocco ²⁷	Y	Y		Neutral	SPP	Y	Synovium of posterior wall of SPP present or absent (> 3 mm considered present) Popliteal cyst present or absent, effusion present or absent	At 3 months, only the PD score was significantly decreased. Synovial thickness decreased by week and further by week 17 (5.5 mm, 4.83 mm, 4.65 mm) Number with effusion decreased from 12 to 4 at week 9, Baker cysts decreased from 5 to 1 (although not significant)	Y	Y	N	U/A
Jan ²⁸			Y	30° flexion	SPP, LR MR		Sac thickness, a summation of measurements in mm in the SPP, LR, MR	Decrease in sac thickness in the treatment group, but not in control group	N	U/A	N	U/A

Table 3. Continued.

Study	Disease			Knee Position	Regions Imaged	Definition of Pathology Imaged	Definition of Scoring System Used	Description of Response	Statistical Assessment Used to Assess Internal Responsiveness?	Internally Responsive?	Statistical Assessment Used to Assess External Responsiveness?	Externally Responsive?
	RA	PsA	OA									
Moore ³⁵	Y				Popliteal fossa		Popliteal cyst presence	Effusion and cysts persisted despite clinical improvement in the majority. IA therapy cohort: effusions and cysts remained at 2–6 weeks. Synovectomy cohort: regression of effusion and cyst	N	U/A	N	U/A
Newman ³⁶	Y	Y			SPP, LR MR		Effusion presence; Power Doppler signal assessed 1–4 in the SPP	Effusion reduced in all subjects posttherapy. Doppler signal reduced in all knees, at least 2 grades	N	U/A	N	U/A
Rubaltelli ⁴⁰	Y	Y	Extension (e)		SPP***, MR, LR	Y	Synovial sac thickness in mm, then graded 0–3	Significant reduction in ultrasound detected synovitis postsurgery in all sites (SPP, $p < 0.005$, MR, $p < 0.05$, LR, $p < 0.02$)	Y	Y	N	U/A
Salaffi ⁴¹	Y		30° flexion		SPP, LR MR		Vascularity of SPP calculated through time-intensity curves using Doppler and contrast	16 of 18 knees had qualitative reduction in Doppler signal changes in ultrasound measurements correlated with changes in clinical measurements	N	U/A	Y	Y
Shio ⁴⁴	Y				SPP		Semiquantification of power Doppler signal (0 = 0 signals, 1 = 1–4 signals, 2 = 5–8 signals, 3 > 8 signals). Resistive index (RI) calculated	Significant improvement in the Doppler grade after therapy, but no improvement in RI after therapy	N	U/A	Y	Y
van Holsbeeck ⁴⁸	Y		30° flexion		SPP		Maximal synovial thickening in the SPP (anterior and posterior walls) in the AP diameter in mm	Decrease in the synovial thickness at 10 and 14 days. Synovial fluid was present in fewer subjects after therapy	N	U/A	N	U/A

* Synovium defined as space between femoral cartilage and prefemoral fat pad. ** Synovium defined as matter between femoral cortex and quadriceps tendon. *** Synovium defined as zone between femoral cartilage and prefemoral fat pad. (a) MR and LR defined as regions at the middle third of the patella with the knee extended. (b) MR and LR defined as the vertical plane of the lateral and medial borders of the patella with the biceps femoris contracted and knee extended. (c) MR and LR defined as the regions imaged when the tail of the probe is placed with the tail lying in the middle transaction of the patella. (d) Biceps femoris contracted. (e) Biceps femoris at rest. RA: rheumatoid arthritis; PsA: psoriatic arthritis; OA: osteoarthritis; U/A: unable to assess; SPP: skin perfusion pressure; PD: Power Doppler; PET: positron emission tomography; MRI: magnetic resonance imaging; CRP: C-reactive protein; MMP: matrix metalloproteinase; IA: intraarticular; AP: anterior/posterior; U/C: unclear. Y: yes; N: no.

REFERENCES

- Grigor C, Capell H, Stirling A, McMahon AD, Lock P, Vallance R, et al. Effect of a treatment strategy of tight control for rheumatoid arthritis (the TICORA study): a single-blind randomised controlled trial. *Lancet* 2004;364:263-9.
- Boers M, Brooks P, Strand CV, Tugwell P. The OMERACT filter for Outcome Measures in Rheumatology. *J Rheumatol* 1998;25:198-9.
- Giles JT, Mease P, Boers M, Bresnihan B, Conaghan PG, Heald A, et al. Assessing single joints in arthritis clinical trials. *J Rheumatol* 2007;34:641-7.
- Husted JA, Cook RJ, Farewell VT, Gladman DD. Methods for assessing responsiveness: a critical review and recommendations. *J Clin Epidemiol* 2000;53:459-68.
- Wakefield RJ, Balint PV, Szkudlarek M, Filippucci E, Backhaus M, D'Agostino M-A, et al. Musculoskeletal ultrasound including definitions for ultrasonographic pathology. *J Rheumatol* 2005;32:2485-7. [erratum appears in *J Rheumatol* 2006;33:440]
- Eckstein F, Cicuttini F, Raynauld J, Waterton J, Peterfy C. Magnetic resonance imaging (MRI) of articular cartilage in knee

Table 4. Summary of the scintigraphy, including scoring system used, and responsiveness.

Study	RA	PsA	OA	Label	Scoring System	Description of Response	Statistical Assessment Used to Assess Internal Responsiveness?	Internally Responsive?	Statistical Assessment Used to Assess External Responsiveness?	Externally Responsive?
Al-Janabi ⁹	Y			99Tc m -HMPAO, Leukocyte imaging	Unclear	Mean 68% reduction in neutrophil migration post- IA ST	N	U/A	N	U/A
Al-Janabi ¹⁰	Y			99Tc m- MDP and 99Tc m HMPOA labeled white cells	Unclear	Linear correlation between change in neutrophil uptake and change in pain (R = 0.87, p < 0.001)	N	U/A	Y	Y
Beckers ¹⁴	Y			18 F-FDG, PET scan	Standardized value uptakes (SVU). Considered positive when uptake corresponded with site of joint synovium	At 4 weeks, significant correlation between changes in PET, MRI < CRP, MMP-3, but not ultrasound	N	U/A	Y	Y
Creamer ¹⁶			Y	99m Tc bone scan	Unclear	99m Tc demonstrated no difference over time in either group	N	U/A	N	U/A
De Bois ¹⁹	Y			99Tc m pertechnetate labeled IgG scintigraphy	0 not increased, 1 = increased faint, 2 = increased moderate, 3 = increased marked	Scintigraphy scores decreased in all knees after therapy	N	U/A	Y	Y
Arzu Gencoglu ²³	Y			Tc 99m HIG and Tc 99m MDP scans	Quantitative analysis by dividing mean count per pixel in knee by mean count per pixel in adjacent normal bone	Tc 99m HIG index values significantly reduced at 3 months (p < 0.001) in the 13 knees that showed clinical response. No significant change in those with a fair or poor clinical response	Y	Y	N	U/A
Arzu Gencoglu ²⁴	Y			Tc 99m-labeled polyclonal HIG	Dividing mean pixel count/pixel region in knee by mean counts/pixel in the adjacent normal bone	No significant change in the imaging results at the knee joint over time	N	U/A	N	U/A
Jones ²⁹	Y			99m Tc HMPAO-labeled neutrophils	Interpolative background subtraction method, expressed as a percentage uptake of dose	Linear reduction in pain decrease and reduction in neutrophil migration 5/6 with P32 demonstrated a decrease in swelling and pain	N	U/A	Y	Y
Soroa ⁴⁵	Y			MDP scan	Computer analysis 0-3.	Data presentation not complete	N	U/A	N	U/A
Tennenbaum ⁴⁶	Y			Tc-99m-MDP and Ga-67 citrate scans	Calculated qualitative ratio of joint to bone and joint to soft tissue	Data presentation not complete	N	U/A	N	U/A
Uematsu ⁴⁷			Y	99m TC labeled MDP or pyrophosphate	Unclear	7 had decreased uptake at followup, and 7 had no change	N	U/A	N	U/A
Youssef ⁵¹	Y			111 in study and 99m Tc HMPAO study	Time activity curves over the region of interest superimposed on a graph with SE bars, and visually inspected for differences. 2 compartment model based on blood pool activity and uptake into synovium used to derive a curve	Significant decrease in neutrophil ingress in 13/16 knees, occurring within 1.5 hours of therapy. In 2 of the 3 with no decrease, the baseline neutrophil ingress was low	N	U/A	N	U/A

RA: rheumatoid arthritis; PsA: psoriatic arthritis; OA: osteoarthritis; HMPAO: hexa-methyl-propylene-amine-oxime; IA ST: intraarticular corticosteroid; U/A: unable to assess; MDP: methylene diphosphonate; FDG: fluoro-deoxy-glucose; PET: positron emission tomography; MRI: magnetic resonance imaging; CRP: C-reactive protein; MMP: matrix metalloproteinase; HIG: human immunoglobulin.

- osteoarthritis (OA): morphological assessment. *Osteoarthritis Cartilage* 2006;14 Suppl:A46-75.
7. Hunter DJ, Le Graverand M-PH, Eckstein F. Radiologic markers of osteoarthritis progression. *Curr Opin Rheumatol* 2009;21:110-7.
 8. Acebes JC, Sanchez-Pernaute O, Diaz-Oca A, Herrero-Beaumont G. Ultrasonographic assessment of Baker's cysts after intra-articular corticosteroid injection in knee osteoarthritis. *J Clin Ultrasound* 2006;34:113-7.
 9. Al-Janabi MA, Jones AK, Solanki K, Sobnack R, Bomanji J, Al-Nahhas AA, et al. 99Tcm-labelled leucocyte imaging in active rheumatoid arthritis. *Nucl Med Commun* 1988;9:987-91.
 10. Al-Janabi MA, Sobnack R, Solanki KK, Jones AK, Critchley M, Smith ML, et al. The response of 99Tcm-methylene diphosphonate and 99Tcm-hexametazime-labelled neutrophils to intra-articular steroid injection in rheumatoid arthritis. *Nucl Med Commun* 1992;13:528-34.
 11. Alonso-Ruiz A, Perez-Ruiz F, Calabozo M, Saez F, Ignacio Pijoan J, Ruiz-Lucea E, et al. Efficacy of radiosynovectomy of the knee in rheumatoid arthritis: evaluation with magnetic resonance imaging. *Clin Rheumatol* 1998;17:277-81.
 12. Bagge E, Geijer M, Tarkowski A. Intra-articular administration of polyclonal immunoglobulin G in rheumatoid arthritis. A double-blind, placebo-controlled pilot study. *Scand J Rheumatol* 1996;25:174-6.
 13. Batalov A, Kuzmanova S, Atanasov A. Ultrasound follow-up study of arthroscoped patients with gonitis. *Folia Med (Plovdiv)* 1999;41:63-70.
 14. Beckers C, Jeukens X, Ribbens C, Andre B, Marcelis S, Leclercq P, et al. (18)F-FDG PET imaging of rheumatoid knee synovitis correlates with dynamic magnetic resonance and sonographic assessments as well as with the serum level of metalloproteinase-3. *Eur J Nucl Med Mol Imaging* 2006;33:275-80.
 15. Brandt KD, Mazza SA, Buckwalter KA. Acetaminophen, like conventional NSAIDs, may reduce synovitis in osteoarthritic knees. *Rheumatology* 2006;45:1389-94.
 16. Creamer P, Sharif M, George E, Meadows K, Cushnaghan J, Shinmei M, et al. Intra-articular hyaluronic acid in osteoarthritis of the knee: an investigation into mechanisms of action. *Osteoarthritis Cartilage* 1994;2:133-40.
 17. Creamer P, Hochberg MC. Why does osteoarthritis of the knee hurt — sometimes? *Br J Rheumatol* 1997;36:726-8.
 18. Cubukcu D, Ardic F, Karabulut N, Topuz O, Hylan G-F 20 efficacy on articular cartilage quality in patients with knee osteoarthritis: clinical and MRI assessment. *Clin Rheumatol* 2005;24:336-41.
 19. de Bois MH, Arndt JW, Tak PP, Kluin PM, van der Velde EA, Pauwels EK, et al. 99Tcm-labelled polyclonal human immunoglobulin G scintigraphy before and after intra-articular knee injection of triamcinolone hexacetonide in patients with rheumatoid arthritis. *Nucl Med Commun* 1993;14:883-7.
 20. Fiocco U, Cozzi L, Rubaltelli L, Rigon C, De Candia A, Tregnaghi A, et al. Long-term sonographic follow-up of rheumatoid and psoriatic proliferative knee joint synovitis. *Br J Rheumatol* 1996;35:155-63.
 21. Fiocco U, Ferro F, Vezzu M, Cozzi L, Checchetto C, Sfriso P, et al. Rheumatoid and psoriatic knee synovitis: clinical, grey scale, and power Doppler ultrasound assessment of the response to etanercept. *Ann Rheum Dis* 2005;64:899-905.
 22. Forslind K, Larsson EM, Eberhardt K, Johansson A, Svensson B. Magnetic resonance imaging of the knee: a tool for prediction of joint damage in early rheumatoid arthritis? *Scand J Rheumatol* 2004;33:154-61.
 23. Arzu Gencoglu E, Aras G, Kucuk O, Atay G, Tutak I, Ataman S, et al. Comparison of Tc-99m HIG and three phase Tc-99m MDP bone scintigraphy for evaluating the efficacy of Yttrium-90 silicate radionuclide scintigraphy. *Clin Nucl Med* 2003;28:277-85.
 24. Gencoglu EA, Aras G, Kucuk O, Atay G, Tutak I, Ataman S, et al. Utility of Tc-99m human polyclonal immunoglobulin G scintigraphy for assessing the efficacy of yttrium-90 silicate therapy in rheumatoid knee synovitis. *Clin Nucl Med* 2002;27:395-400.
 25. Glaser C, Tins BJ, Trumm CG, Richardson JB, Reiser MF, McCall IW. Quantitative 3D MR evaluation of autologous chondrocyte implantation in the knee: feasibility and initial results. *Osteoarthritis Cartilage* 2007;15:798-807.
 26. Hunter DJ, Conaghan PG, Peterfy CG, Bloch D, Guermazi A, Woodworth T, et al. Responsiveness, effect size, and smallest detectable difference of magnetic resonance imaging in knee osteoarthritis. *Osteoarthritis Cartilage* 2006;14 Suppl:A112-5.
 27. Iagnocco A, Cerioni A, Coari G, Ossandon A, Masciangelo R, Valesini G. Intra-articular methotrexate in the treatment of rheumatoid arthritis and psoriatic arthritis: a clinical and sonographic study. *Clin Rheumatol* 2006;25:159-63.
 28. Jan MH, Chai HM, Wang CL, Lin YF, Tsai LY. Effects of repetitive shortwave diathermy for reducing synovitis in patients with knee osteoarthritis: an ultrasonographic study. *Phys Ther* 2006;86:236-44.
 29. Jones AK, al-Janabi MA, Solanki K, Sobnack R, Greenwood A, Doyle DV, et al. In vivo leukocyte migration in arthritis. *Arthritis Rheum* 1991;34:270-5.
 30. Kroner A, Berger C, Kluger R, Oberhauser G, Bock P, Engel A. Influence of high tibial osteotomy on bone marrow edema in the knee. *Clin Orthop Relat Res* 2007;454:155-62.
 31. Lee SH, Suh JS, Kim HS, Lee JD, Song J, Lee SK. MR evaluation of radiation synovectomy of the knee by means of intra-articular injection of holmium-166-chitosan complex in patients with rheumatoid arthritis: results at 4-month follow-up. *Korean J Radiol* 2003;4:170-8.
 32. Leitch R, Walker SE, Hillard AE. The rheumatoid knee before and after arthrocentesis and prednisolone injection: evaluation by Gd-enhanced MRI. *Clin Rheumatol* 1996;15:358-66.
 33. Marzo-Ortega H, McGonagle R, Rhodes LA, Tan AL, Conaghan PG, O'Connor P, et al. Efficacy of infliximab on MRI-determined bone oedema in psoriatic arthritis. *Ann Rheum Dis* 2007;66:778-81.
 34. Mizner RL, Petterson SC, Stevens JE, Vandenberg K, Snyder-Mackler L. Early quadriceps strength loss after total knee arthroplasty. The contributions of muscle atrophy and failure of voluntary muscle activation. *J Bone Joint Surg Am* 2005;87:1047-53.
 35. Moore CP, Sarti DA, Louie JS. Ultrasonographic demonstration of popliteal cysts in rheumatoid arthritis. A noninvasive technique. *Arthritis Rheum* 1975;18:577-80.
 36. Newman J, Laing T, McCarthy C, Adler R. Power Doppler sonography of synovitis: assessment of therapeutic response — preliminary observations. *Radiology* 1996;198:582-4.
 37. Ostergaard M, Stoltenberg M, Henriksen O, Lorenzen I. Quantitative assessment of synovial inflammation by dynamic gadolinium-enhanced magnetic resonance imaging. A study of the effect of intra-articular methylprednisolone on the rate of early synovial enhancement. *Br J Rheumatol* 1996;35:50-9.
 38. Ozturk C, Atamaz F, Hepgul S, Argin M, Arkun R. The safety and efficacy of intraarticular hyaluronan with/without corticosteroid in knee osteoarthritis: 1-year, single-blind, randomized study. *Rheumatol Int* 2006;26:314-9.
 39. Reece RJ, Kraan MC, Radjenovic A, Veale DJ, O'Connor P, Ridgway JP, et al. Comparative assessment of leflunomide and methotrexate for the treatment of rheumatoid arthritis, by dynamic enhanced magnetic resonance imaging. *Arthritis Rheum* 2002;46:366-72.
 40. Rubaltelli L, Fiocco U, Cozzi L, Baldovin M, Rigon C, Bortoletto P, et al. Prospective sonographic and arthroscopic evaluation of proliferative knee joint synovitis. *J Ultrasound Med*

- 1994;13:855-62.
41. Salaffi F, Carotti M, Manganelli P, Filippucci E, Giuseppetti GM, Grassi W. Contrast-enhanced power Doppler sonography of knee synovitis in rheumatoid arthritis: assessment of therapeutic response. *Clin Rheumatol* 2004;23:285-90.
 42. Sharma S, Rastogi S, Gupta V, Rohtagi D, Gulati P. Comparative efficacy and safety of nimesulide versus piroxicam in osteoarthritis with special reference to chondroprotection. *Am J Ther* 1999;6:191-7.
 43. Shin K, Lee JC, Choi HJ, Jeong JM, Son M, Lee YJ, et al. Radiation synovectomy using ¹⁸⁸Re-tin colloid improves knee synovitis as shown by MRI in refractory rheumatoid arthritis. *Nucl Med Commun* 2007;28:239-44.
 44. Shio K, Homma F, Kanno Y, Yamadera Y, Ohguchi Y, Nishimaki T, et al. Doppler sonographic comparative study on usefulness of synovial vascularity between knee and metacarpophalangeal joints for evaluation of articular inflammation in patients with rheumatoid arthritis treated by infliximab. *Mod Rheumatol* 2006;16:220-5.
 45. Soroa VE, del Huerto Velazquez Espeche M, Giannone C, Caviglia H, Galatros G, Fernandez D, et al. Effects of radiosynovectomy with p-32 colloid therapy in hemophilia and rheumatoid arthritis. *Cancer Biother Radiopharm* 2005;20:344-8.
 46. Tannenbaum H, Rosenthal L, Arzoumanian A. Quantitative scintigraphy using radiophosphate and radiogallium in patients with rheumatoid arthritis. *Clin Exp Rheumatol* 1987;5:17-21.
 47. Uematsu A, Kim EE. Role of radionuclide joint imaging in high tibial osteotomy. *Clin Orthop Relat Res* 1979;144:220-5.
 48. van Holsbeeck M, van Holsbeeck K, Gevers G, Marchal G, van Steen A, Favril A, et al. Staging and follow-up of rheumatoid arthritis of the knee. Comparison of sonography, thermography, and clinical assessment. *J Ultrasound Med* 1988;7:561-6.
 49. Veale DJ, Reece RJ, Parsons W, Radjenovic A, O'Connor PJ, Orgles CS, et al. Intra-articular primatised anti-CD4: efficacy in resistant rheumatoid knees. A study of combined arthroscopy, magnetic resonance imaging, and histology. *Ann Rheum Dis* 1999;58:342-9.
 50. Wluka AE, Stuckey S, Brand C, Cicuttini FM. Supplementary vitamin E does not affect the loss of cartilage volume in knee osteoarthritis: a 2 year double blind randomized placebo controlled study. *J Rheumatol* 2002;29:2585-91.
 51. Youssef PP, Cormack J, Evill CA, Peter DT, Roberts-Thomson PJ, Ahern MJ, et al. Neutrophil trafficking into inflamed joints in patients with rheumatoid arthritis, and the effects of methylprednisolone. *Arthritis Rheum* 1996;39:216-25.
 52. Song IH, Althoff CE, Hermann KG, Scheel AK, Knetsch T, Burmester GR, et al. Contrast-enhanced ultrasound in monitoring the efficacy of a bradykinin receptor 2 antagonist in painful knee osteoarthritis compared with MRI. *Ann Rheum Dis* 2009;68:75-83.
 53. Raynauld JP, Martel-Pelletier J, Bias P, Laufer S, Haraoui B, Choquette D, et al. Protective effects of licofelone, a 5-lipoxygenase and cyclo-oxygenase inhibitor, versus naproxen on cartilage loss in knee osteoarthritis: a first multicentre clinical trial using quantitative MRI. *Ann Rheum Dis* 2009;68:938-47.
 54. Raynauld JP, Martel-Pelletier J, Berthiaume MJ, Abram F, Choquette D, Haraoui B, et al. Correlation between bone lesion changes and cartilage volume loss in patients with osteoarthritis of the knee as assessed by quantitative magnetic resonance imaging over a 24-month period. *Ann Rheum Dis* 2008;67:683-8.
 55. Raynauld JP, Martel-Pelletier J, Abram F, Dorais M, Haraoui B, Choquette D, et al. Analysis of the precision and sensitivity to change of different approaches to assess cartilage loss by quantitative MRI in a longitudinal multicentre clinical trial in patients with knee osteoarthritis. *Arthritis Res Ther* 2008;10:R129.
 56. Anandacoomarasamy A, Bagga H, Ding C, Burkhardt D, Sambrook PN, March LM. Predictors of clinical response to intraarticular Hylan injections — a prospective study using synovial fluid measures, clinical outcomes, and magnetic resonance imaging. *J Rheumatol* 2008;35:685-90.
 57. Creamer P, Keen M, Zananiri F, Waterton JC, Maciewicz RA, Oliver C, et al. Quantitative magnetic resonance imaging of the knee: a method of measuring response to intra-articular treatments. *Ann Rheum Dis* 1997;56:378-81.