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Musculoskeletal interventional procedures: With or without imaging guidance?

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A B S T R A C T

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Aspiration and injection of joints and soft tissues is an indispensable skill used in everyday practice by the clinical rheumatologist. Most rheumatologists recognise that performing these procedures using anatomical landmarks is not always successful, particularly in the case of small or infrequently injected joints, bursae or tendon sheaths. Musculoskeletal ultrasound confirms the local pathological-anatomical diagnosis and is the most applicable and feasible imaging method that can be applied in clinical practice in guiding musculoskeletal interventional procedures. From 1993, there has been substantial examination of the accuracy of landmark- and imaging-guided procedures. We have searched the literature and ascertained whether imaging techniques improve the accuracy of musculoskeletal procedures and whether the accuracy of needle placement can be translated into improved clinical outcome (efficacy).

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Introduction

Despite major advances in systemic therapy for the treatment of inflammatory rheumatic diseases, corticosteroid (steroid) injection into the joints, bursae, tendon sheaths or other musculoskeletal soft tissues remains an important tool used in daily clinical practice. These injections are usually guided by the palpation of anatomical landmarks and are termed landmark-guided injection (LMGI) (also known

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as palpation-guided, clinical examination-guided or blind injection techniques). Jones et al., in 1993 were the first to report the success rates of needle placement using a palpation-guided approach [1]. The authors proposed that accurate placement of the needle could diminish the discomfort experienced by the patient and reduce incidental damage to adjacent structures and tissue atrophy within the extra-articular soft tissues.

Ultrasound (US) guidance of aspiration was first reported by Gompels and Darlington in 1981 [2]. The first step of an US-guided procedure is always to confirm and localise the joint or soft tissue pathology. In clinical practice, US has been shown to frequently change the anatomical and pathological diagnosis made on clinical grounds, which alters the decision of whether or not to inject and where to place the needle [3]. The superior diagnostic accuracy of US scanning compared with clinical examination is a major factor that should be examined in future studies that compare the outcomes of LMGI and US-guided injections (USGIs). In addition to aspiration and steroid injections, it is now possible to perform US-guided percutaneous biopsies of the joints, bursae, tendon sheaths, major salivary glands and muscles; to assist insertion of drainage catheters; to detect soft tissue foreign bodies; and to perform nerve blocks. There are two common methods for USGIs: semi-guided or indirect method (skin surface marking) and needle guidance under direct sonographic vision (direct method) [4,5]. Direct visualisation of the needle is considered to be the superior technique because it confirms the correct position of the needle. The needle is observed in real time while it is passing through the tissue. There are no studies that directly compare the performance of these two USGI techniques.

US is the optimal imaging modality for guiding musculoskeletal interventional procedures because it is extensively available at bed side, provides high definition imaging of both needle and joint tissues, incurs relatively low costs and does not use radiation [6]. In this article, we address whether USGIs improve the accuracy of therapeutic musculoskeletal injections compared with LMGI and, further, the accuracy of needle placement can be translated into improved clinical outcome (efficacy) in the different anatomical regions. We searched all English language articles published in the last 35 years using PubMed database and collected clinical trials comparing the accuracy and/or efficacy of LMGI with USGI. We focused on human rheumatic and musculoskeletal conditions and excluded oncology, traumatology and spinal procedures.

Shoulder

Glenohumeral joint

Accuracy

In a study by Cunningham et al., injections of steroid and contrast agent were given in 184 different joints randomised for LMGI and USGI. The accuracy was assessed by radiography of the injected joint, which localised the contrast agent in the injectate. The accuracy of the glenohumeral joint injections reported for USGI was 63% and for LMGI was 40% ($p = NS$) [7]. It should be noted that in this study, USGI was performed using the posterior approach by a relatively inexperienced ultrasonographer with one year of practical US experience, and 40% accuracy was achieved by rheumatologists with a median of 14 years of clinical experience in joint injection. It would be expected that the accuracy of a skilled ultrasonographer interventionist would be much higher, and the sample size in this study was small making statistical analysis difficult.

There are eight clinical studies that investigated the use of LMGI. Three of these used magnetic resonance (MR) arthrography for accurate injection verification, four used contrast medium and fluoroscopy and one used arthroscopy. According to these reports, the mean accuracy of LMGI to the glenohumeral joint was 67% (variation 10–99%) [1,8–14]. In a paper by Tobola et al. [13], three different injection approaches were investigated (anterior, posterior or supraclavicular), and statistically significant differences were absent between the three approaches. However, the anterior route was the most accurate and was independent of the experience level of the provider.

There are five other studies that have investigated the accuracy of USGI before MR arthrography [15–19]. The mean reported accuracy of USGI to the glenohumeral joint were 96% (variation 92–100%). One of these studies compared US- and fluoroscopy-guided techniques and noted that the

first attempt was successful in 72% of the injections in the fluoroscopy-guided group and 94% in the US-guided group [18].

In a cadaveric study with 80 shoulder specimens, the accuracy of USGIs and LMGI were 92.5% and 72.5% ($p = 0.025$), respectively [20]. Two operators injected radiopaque contrast through a posterior approach. After the injections, radiographs of the specimens were obtained to assess the accuracy of the injections.

There are three cadaveric studies that investigated LMGI to the glenohumeral joint [21–23]. The mean accuracy was 94% (variation 91–96%). One study used superior injection approach and noted that caution for the likelihood of penetrating the long head of biceps tendon should be considered [22].

These findings provide evidence that USGIs to the glenohumeral joint are more accurate than LMGI (Table 1).

Efficacy

There is a single comparative study of LMGI and USGI that investigated the efficacy of glenohumeral joint injections using these techniques. In this trial for patients with adhesive capsulitis, the US group had a lower reduction in pain VAS; however, no significant difference was observed between groups for function at 6 weeks post injection [24]. In Cunnington's study, there was no significant difference for outcome variables between USGIs and LMGI when all the joints were grouped together. The results for single joints were not reported. However, there was a greater improvement in the VAS score for function in the accurate injection group [7]. A recent Cochrane review reported no significant improvement in efficacy with USGIs to the shoulder [25]. However, the authors did not perform the analysis on the basis of the exact injection location. In addition, there were fewer side effects in the USGI groups.

Presently, there is not enough data to conclude that USGIs are more efficacious than LMGI in the treatment of glenohumeral joint diseases.

Subacromial space

Accuracy

In assessing the studies on injection into the subacromial space, it should be noted that different studies have used different volumes of injectate, and a larger volume may aid the accuracy but leads to more tissue damage. Two clinical studies have compared LMGI and USGI using MR arthrography for verifying the success of the intervention. Rutten et al. reported that both methods were 100% accurate [26], and Dogu et al. found that accurate injections were performed in 65% in the USGI group and 70% in the LMGI group [27].

Four clinical studies have investigated the accuracy of LMGI into the subacromial space [28–31]. One used MR arthrography and the others used fluoroscopy plus contrast media for verification. The mean accuracy was 61% (variation 29–91%).

Three cadaveric studies have examined the accuracy of LMGI into the subacromial space [21,32,33]. The mean accuracy was 81% (variation 70–91%). Thus, the accuracy of LMGI into the subacromial space is sub-optimal and depends on the operator.

There is not enough data to confirm that USGIs are more accurate than LMGI, and further research is required to clarify this (Table 2).

Table 1

The mean accuracies of glenohumeral joint injection in 18 studies. The number of studies are given in parentheses.

	LMGI (%)	USGI (%)
Comparative clinical study (1)	40	63
Clinical LMGI studies (8)	67	–
Clinical USGI studies (5)	–	96
Comparative cadaveric study (1)	72.5	92.5
Cadaveric LMGI studies (3)	94	–

LMGI, landmark-guided injection; USGI, ultrasound-guided injection.

Efficacy

Five studies have evaluated the efficacy of USGIs and LMGI into the subacromial space, and they all showed better outcomes for the USGI procedures in the short term [34–38]. However, two additional studies found that there is no difference in efficacy between accurate and inaccurate injections [27,29].

According to the literature, USGIs into the subacromial space produce a significantly greater improvement in terms of pain reduction and functional gain than LMGI in chronic shoulder pain, at least in short term.

Biceps tendon sheath, acromioclavicular joint and sternoclavicular joint

Accuracy and efficacy

Injection into the tendon sheath of the long head of the biceps brachii can be more accurately performed by USGI than that by LMGI according to a single study [39]. USGIs showed 87% accuracy, whereas LMGI showed only 27% accuracy. Computed tomography (CT) with contrast agent was used for the verification of the injection; however, the efficacy of this was not reported.

In one clinical study, the accuracy of LMGI to the acromioclavicular joint was 43%. This was verified using radiographic contrast material [40]. There are no papers reporting the efficacy of acromioclavicular injections.

The exact position of USGI and LMGI into the acromioclavicular joint has been examined in three cadaver studies [41–43]. The mean accuracies reported were 96% (variation 90–100%) for USGIs and 62% (variation 40–75%) for LMGI. In a cadaveric study, Partington and Broome found that LMGI were successful in 67% of the acromioclavicular injections [32].

It can be concluded that LMGI to the biceps tendon sheath and acromioclavicular joint are sub-optimal and USGIs are more accurate; however, efficacy studies for this are missing.

The only study evaluating injection accuracy into the sternoclavicular joint used LMGI approach with a reported accuracy of 78% [44]. This was a cadaveric study (Table 3).

Advanced procedures of the shoulder

Some procedures in the shoulder area require exact imaging guidance. Percutaneous US-guided fenestration and aspiration of calcific tendinosis is an example [45]. It was originally done as a fluoroscopic procedure; however, US has largely replaced fluoroscopy because of its excellent safety profile and clinical efficacy. In the shoulder region, nerve blocks can be performed using US guidance (e.g. suprascapular and dorsal scapular nerve blocks). US may be used to diagnose and aspirate a ganglion cyst compressing the nerve in the suprascapular or spinoglenoid notch. Other injectable therapies such as platelet-rich plasma are now available for rotator cuff tendinopathy; however, further studies are required to clarify their efficacy, and there is no comparison of USGI and LMGI for these therapies.

Elbow

Elbow joint: accuracy and efficacy

Two studies have compared the accuracies of LMGI and USGI to the elbow joint. The study by Cunningham [7] reported accuracies of 64% and 91% for LMGI and USGI, respectively ($p = 0.1$). Kim

Table 2

The mean accuracies of subacromial space injection in nine studies. The number of studies are given in parentheses.

	LMGI (%)	USGI (%)
Comparative clinical studies (2)	85	82.5
Clinical LMGI studies (4)	61	–
Cadaveric LMGI studies (3)	95	–

LMGI, landmark-guided injection; USGI, ultrasound-guided injection.

Table 3

The mean accuracies of injection to the biceps tendon sheath, acromioclavicular and sternoclavicular joints in seven studies. The number of studies are given in parentheses.

	LMGI (%)	USGI (%)
Biceps tendon sheath: comparative clinical study (1)	27	87
Acromioclavicular joint: clinical LMGI study (1)	43	–
Acromioclavicular joint: comparative cadaveric studies (3)	62	96
Acromioclavicular joint: cadaveric LMGI study (1)	67	–
Sternoclavicular joint: cadaveric LMGI study (1)	78	–

LMGI, landmark-guided injection; USGI, ultrasound-guided injection.

reported an accuracy of 100% using USGI and 77.5% using LMGI by posterior (olecranon) approach. They verified the injection accuracy using the fluoroscopic contrast method in 80 patients [46].

Two studies have evaluated LMGI accuracy to the elbow joint, and the mean accuracy was 92% (variation 84–100%) [1,10]. Both the studies used radiographic contrast media to verify the success of the intra-articular injection. Of these, only Lopes et al. reported the efficacy and noted significant improvement in the VAS for pain at rest and during movement, VAS for oedema and morning stiffness [10].

It appears that LMGI and USGI to the elbow joint do not differ much in accuracy, although the number of studies is small (Table 4).

Lateral epicondylitis and advanced procedures around the elbow

Lateral epicondylitis is a common condition in which different types of treatments have been investigated with varying results. There are no studies of the accuracy of LMGI and USGI in this area. The definition of accuracy is also complex given that some injection approaches target the surface of the common extensor origin, whereas others aim to penetrate and disrupt the tendon structure. US scanning plays an important role in the differential diagnosis of lateral elbow pain given that the CEO, lateral collateral ligament and humeroradial joint are so closely apposed.

Different types of US-guided procedures such as needle tendon fenestration and platelet-rich plasma injections have been and are to be investigated in the treatment of lateral epicondylitis.

US guidance has an important role in performing nerve blocks at the elbow level such as blocking of the posterior interosseous nerve and radial nerve at the spiral groove. Ganglion cysts compressing nerves can be diagnosed using US and also aspirated under US guidance.

Wrist and hand

Wrist and hand joints: accuracy

Cunnington et al. achieved 79% and 75% accuracy using USGI and LMGI, respectively, in the wrist joint ($p = 0.8$) [7]. In two studies, LMGI to the wrist joint resulted in accuracies of 97% and 50% [10,1]. In these three papers, the radiographic contrast medium method was used for verifying the success of intra-articular injection. Choudur et al. used US guidance in MR arthrography and found that 99% were successfully injected into the wrist [15].

Smith et al. studied the accuracy of USGI and LMGI to the scaphotrapeziotrapezoid joint [47]. This cadaver study showed that USGI were 100% accurate, whereas LMGI were only 80% accurate. There

Table 4

The mean accuracies of elbow joint injection in four studies. The number of studies are given in parentheses.

	LMGI (%)	USGI (%)
Clinical comparative studies (2)	71	96
Clinical LMGI studies (2)	92	–

LMGI, landmark-guided injection.

are two additional cadaver studies examining the accuracies of USGIs to the trapeziometacarpal and distal radioulnar joints, and the accuracies were 94% and 100%, respectively [48,49].

A single study compared the accuracy of LMGI and USGI to PIP and MCP joints [50]. The needle position was intra-articular in 50% of PIP joints and 80% of MCP joints using LMGI, and the corresponding figures for USGI were 92% and 100%. In this study, visualisation of the needle tip within the joint space and distension of the joint capsule following steroid injection defined intra-articular positioning of the needle.

The accuracies of USGIs to the wrist and hand are overall higher than LMGI, but because of the paucity of studies, further research is required to make definite conclusions (Table 5).

Wrist and hand joints: efficacy

Four studies have compared USGI and LMGI to the wrist joint [7,51–53]. These studies report a statistically and clinically significant benefit of USGI over LMGI in the reduction of wrist pain and achievement of minimal clinically important improvement.

High-quality studies of other hand joints on the efficacy of LMGI and USGI are missing.

Soft tissues of the hand: efficacy and accuracy

Kume's randomised prospective study compared USGI and LMGI to 44 wrists with de Quervain's disease (separate extensor pollicis brevis). Pain was significantly reduced more in the USGI group after 4 weeks in this prospective open-label study [54].

In another study, 77 symptomatic carpal tunnels were randomised to be injected by either LMGI or USGI [55]. There were no complications in either treatment group. Compared with LMGI, USGI resulted in a 77% reduction in injection pain, 63% reduction in pain scores, 93% increase in the responder rate, 85% reduction in the non-responder rate, 71% increase in therapeutic duration and 59% reduction in cost/responder/year for a hospital outpatient. All the figures were statistically significant.

In a study by Ustün et al., 46 patients with carpal tunnel syndrome were randomised to receive either LMGI or USGI into the carpal tunnel [56]. The improvement in symptom severity scores in the USGI group at 12 weeks was higher than those in the LMGI group ($p < 0.05$). Average time to symptom relief was shorter in the USGI group ($p < 0.05$).

Lee et al. injected methylene blue dye into 40 flexor tendon sheaths of 5 cadavers using blind or sonographic guidance. Dissection revealed that 70% of USGI were successful, whereas only 15% LMGI were successful [57].

It can be concluded that USGI are more accurate and efficient than LMGI in carpal tunnel and tendon sheath injections of the hand.

Advanced procedures in the hand

All advanced procedures require USGI to be performed, and therefore, there is no comparative study with LMGI. Deeply located ganglion cysts can be aspirated and fenestrated with the help of US. Ulnar and median nerve hydrodissections can be performed using a large volume of injectate to release the nerve from

Table 5

The mean accuracies of injection to the wrist and hand joints in 8 studies. The number of studies are given in parentheses.

	LMGI (%)	USGI (%)
Wrist joint: clinical comparative study (1)	75	79
PIP and MCP joints: clinical comparative study (1)	65	96
Wrist joint: clinical LMGI studies (2)	74	–
Wrist joint: clinical USGI study (1)	–	99
Scaphotrapeziotrapezoid joint: comparative cadaveric study (1)	80	100
Trapeziometacarpal joint: cadaveric USGI study (1)	–	94
Distal radioulnar joint: cadaveric USGI study (1)	–	100

LMGI, landmark-guided injection; USGI, ultrasound-guided injection.

the adjacent soft tissue. US-guided techniques have been developed to release A1 pulleys with a surgical hook knife or needle, and even carpal tunnel flexor retinaculum dissections can be performed under US guidance. Further research is required to determine the efficacy, safety profile and cost-effectiveness of these new procedures compared with traditional open procedures. US imaging plays an important role in foreign body removal in the hand because it can detect all kinds of foreign bodies (metal, wood and glass).

Hip

Hip joint: accuracy and efficacy

There are no studies that directly compare USGIs and LMGI to the hip joint. Five studies have investigated LMGI to the hip joint [58–62]. There is a great variation in the accuracies (51–93%), and the mean figure of accuracy is 71% for LMGI to the hip joint.

Four papers have investigated USGIs using MR arthrography or contrast-enhanced fluoroscopic/CT for verifying the success of the injections [15,63–65]. Accuracies are very high, and the mean value is 99% (variation 97–100%).

Thus, intra-articular injections to the hip joint can be performed with high accuracy using USGIs, whereas the accuracy of LMGI is sub-optimal (Table 6).

There are no high-quality studies comparing the efficacy of USGI and LMGI to hip joints. In clinical work, the diagnostic importance of US is highlighted because the clinical evaluation of hip joint arthritis is very difficult because of its deep-lying position and complex surrounding structures in the deep gluteal and femoral areas.

Soft tissue injections around the hip: efficacy and accuracy

Effectiveness of peritendinous USGI of corticosteroid for the treatment of gluteus medius tendinopathy has been investigated in 54 consecutive patients [66]. There was a 55% average reduction of pain level after treatment ($p < 0.001$), and 1 month after treatment, 72% of the patients showed a clinically significant improvement in pain level.

Finnoff compared the accuracy of US-guided piriformis injections with fluoroscopically guided contrast-controlled piriformis injections in a cadaveric model [67]. Nineteen of the 20 USGIs (95%) were correctly placed, and the liquid latex was found within the piriformis muscle, whereas only 6 of the 20 fluoroscopically guided injections (30%) were accurate ($p = 0.001$).

In another study, a single experienced operator completed 10 sonographically guided obturator internus injections in five unembalmed cadaveric pelvis specimens [68]. The results showed that USGIs to the obturator internus or its bursa are feasible and may play a role in the diagnosis and management of patients presenting with deep gluteal pain syndromes.

USGIs are more accurate than LMGI to and around the hip joint; however, there is a lack of good quality efficacy studies.

Sacroiliac joint

Sacroiliac joint: accuracy and efficacy

There are considerable challenges in assessing the injection accuracy of the sacroiliac joint (SIJ). The definition of accurate injection needs clarification as most techniques are reported for intra-articular

Table 6

The mean accuracies of injection to the hip joint in nine studies. The number of studies are given in parentheses.

	LMGI (%)	USGI (%)
Clinical LMGI studies (5)	71	–
Clinical USGI studies (4)	–	99

LMGI, landmark-guided injection; USGI, ultrasound-guided injection.

injection into the more caudal synovial cavity, although in spondyloarthritis, there may be more marked inflammation in the superior fibrous part of the joint, which requires a periarticular injection because of the lack of a joint cavity. Moreover, in assessing the accuracy, the use of contrast is limited by the small volume capacity of the joint.

There is a lack of high-quality evaluating the accuracy of LMGI. Three studies have investigated the accuracy of USGIs to the SIJ [69–71]. Pekkafehli et al. injected 60 SIJs, and the accuracy of USGIs was 76%, and the efficacy of injection was not reported [69]. Hartung et al. achieved an accuracy of 40% in USGIs to SIJs with a reduction in pain score from 6.9 to 3.9 on a 0–10 scale 1 month later ($p < 0.009$). Verification of the success of the injection (steroid plus gadolinium) was confirmed with MR scanning. They noted that USGIs to the SIJ is challenging. Surprisingly, the pain and symptom control was equal in both intra-articular and periarticular injections [70]. Klauser et al. injected SIJs in cadavers and reported an accuracy of 80% with CT confirming the correct intra-articular needle position. The mean pain relief was 8.6 on a 0–10 scale after 3 months [71]. Thus, more studies are required to confirm the position of LMGI and USGI to the SIJ in terms of accuracy and efficacy.

The difficulty of visualising the needle in the SIJ can be addressed using an image fusion technique where real-time US scanning can be matched to previously obtained CT or MR images. Klauser et al. found this new technique to be reliable in both cadavers and patients, and intra-articular needle positioning was possible [72].

Knee

Knee joint: accuracy

There are five clinical studies that have compared the accuracies of LMGI and USGIs to knee joints [7,73–76]. All the studies used contrast agent with radiography for the verification of the intra-articular injection except one study that used positive aspiration as a sign of right needle position [76]. The mean accuracies for LMGI and USGIs were 70% (variation 32–84%) and 95% (91–97%), respectively.

Four clinical studies have examined the accuracy of LMGI to the knee joint [1,10,77,78]. Contrast agents with radiographs were used for the verification of the injections. The mean accuracy was 84%, but again the variation was wide (66–100%). Toda et al. highlighted the need to change the injection approach according to the severity of knee osteoarthritis (OA) to achieve injection accuracy [78].

Two clinical studies have investigated the accuracy of USGIs to the knee joint. One used US-guided gadolinium for MR arthrography and reported an accuracy of 100% in the knee joints [15]. The other study used contrast agent and radiographs, and the rate of injection accuracy varied depending on the site injected [79]. Injections to the superolateral (100%) and mid-lateral portals (95%) showed significantly higher accuracy than injections to the medial portal (75%).

One cadaver study reported an accuracy of 100% for USGIs and 55–100% for LMGI depending on the experience of the operators [80]. Another cadaver study examined the accuracy for four different intra-articular injection sites in the knee using LMGI [81]. This study included 156 knees from 78 fresh cadavers. Perfect accuracy could not be obtained through any of the portals. The best accuracy rate was in the anterolateral injection portal (85%) and lowest was in the medial mid-patellar portal (56%).

A single study investigated injections to proximal tibiofibular joint in unembalmed cadavers [82]. All 12 US-guided latex injections were placed into the target (100% accuracy), whereas only 7 of 12 injections were placed using LMGI (58%).

According to the literature, USGIs to the knee joint are more accurate than LMGI (Table 7).

Knee joint: efficacy and cost-effectiveness

Sibbitt [83] randomised 94 non-effusive knees with OA to corticosteroid injection by LMGI and USGI. In this study, USGIs resulted in a 48% reduction in procedural pain ($p < 0.001$), 107% increase in the responder rate ($p < 0.001$), 52% reduction in the non-responder rate ($p < 0.001$), 36% increase in therapeutic duration ($p = 0.01$), 13% reduction (\$17) in cost per patient per year and 58% (\$224) reduction in cost per responder per year for a hospital outpatient ($p < 0.001$). The follow-up period was 6 months. In another study [84], 64 knees with palpable effusions were randomised to receive LMGI or

USGIs with corticosteroid. USGI resulted in 48% less procedural pain (VAS; palpation guided: 5.8 ± 3.0 cm, US guided: 3.0 ± 2.8 cm, $p < 0.001$), 183% increase in aspirated synovial fluid volumes ($p < 0.0001$) and improved outcomes at 2 weeks ($p = 0.034$).

In contrast, Toda et al. demonstrated no difference in efficacy between accurate and inaccurate hyaluronic acid injections to the knee in patients with OA [78].

Soft tissues around the knee joint and advanced procedures

A study assessed using US the clinical changes of Baker's cyst of patients with knee OA after steroid injection. In all patients, the cyst was aspirated under US guidance first. Twenty patients received corticosteroid injection into the Baker's cyst and 20 received intra-articular injections [85]. At 4 and 8 weeks, cyst diameters measured by US were lower when the cyst was directly infiltrated than with intra-articular injection ($p < 0.01$).

In a cadaveric model, 12 US-guided and 12 unguided pes anserinus bursa injections using coloured liquid latex were performed on 24 unembalmed adult cadaveric lower extremity specimens. The order of the injection techniques was randomised. After dissection, it was revealed that the accuracy rate was 92% (11 of 12 specimens) for USGI and 17% (2 of 12 specimens) for LMGI [86].

In another cadaveric study [87], a single experienced operator completed 24 USG popliteus tendon sheath injections with diluted coloured latex: 12 using a longitudinal approach and 12 using a transverse approach relative to the tendon. The results showed that USGI can be used to inject the popliteus tendon sheath with a high degree of accuracy. The longitudinal approach was potentially more accurate. Both approaches may result in injectate overflow into the knee joint, most likely through the popliteus hiatus.

There is a current trend towards expanded applications of interventional musculoskeletal USG procedures such as percutaneous patellar tenotomy, patellar tendon-Hoffa's fat pad hydrodissection, meniscal cyst aspiration, pericruciate ganglion cyst aspiration, fibulotibial joint injection and fibular nerve hydrodissection at the knee joint level. There is a paucity of high-quality studies on these techniques, but research is ongoing.

Soft tissue anatomic structures around the knee joint can be more accurately injected using USGIs, but in terms of efficacy, the final conclusion on the role of USGI and LMGI cannot be made.

Ankle and foot

Ankle and foot: clinical studies of accuracy and efficacy

Relatively few clinical studies have been conducted in this area. One study compared LMGIs and USGIs to the ankle joint and found that the accuracies were 58% and 85% ($p = \text{NS}$), respectively [7]. In the paper by Jones et al. [1], the accuracy of ankle joint LMGIs was 66%. A single paper has examined both the accuracy and efficacy of LMGIs to the ankle joint [10] and reported an accuracy of 77%. The authors noted that significant improvement was seen in the VAS for pain, oedema and morning stiffness. The accuracy of the LMGIs to the ankle joint in clinical studies seems sub-optimal, but the benefit of USGIs with respect to accuracy and efficacy requires further investigation.

Table 7

The mean accuracies of injection to knee joint in 13 studies. The number of studies are given in parentheses.

	LMGI (%)	USGI (%)
Comparative clinical studies (5)	70	95
Clinical LMGI studies (4)	84	–
Clinical USGI studies (2)	–	91
Comparative cadaveric study (1)	77.5	100
Cadaveric LMGI study (1)	85	–

LMGI, landmark-guided injection; USGI, ultrasound-guided injection.

Four comparative studies of LMGI and USGI concerning the efficacy of injections to the plantar fasciitis have been performed [88–91]. In these studies, there were altogether 140 patients randomised to USGI or LMGI. The mean follow-up was 26 weeks (12–56 weeks). In all papers, the result showed that LMGI were as efficient as the USGI. In one paper, the recurrence rate of plantar fasciitis was significantly higher when using LMGI [89].

Ankle and foot: cadaveric studies of injection accuracy

There are a considerable number of cadaveric studies of injection accuracies in the foot and ankle. Four studies compared LMGI and USGI to different joints and soft tissues using coloured latex solution and dissection after the injection procedure (the dissecting anatomists were blinded to the injection technique) [92–95]. One paper investigated the accuracy of LMGI and USGI to the ankle joint with reported accuracies of 85% and 100%, respectively [92]. They also noted that the injection accuracy into the sinus tarsi was 90% for USGI and 35% for LMGI. Khosla et al. concluded that intra-articular injections to the subtalar and ankle joints can be successfully performed using palpation alone (100% accuracy for both USGI and LMGI) [93], but USGI was superior for injection to the first and second tarsometatarsal joints (accuracy: LMGI 27% vs USGI 64%; $p = 0.003$). A third paper examined LMGI and USGI to the peroneal tendon sheaths in 20 cadaveric lower limbs [95]. Their accuracies were 60% and 100%, respectively. In the fourth study, LMGI and USGI were performed into the deep and superficial posterior leg compartments by a less experienced and a more experienced clinician [94]. The result showed that these injections were equally accurate with palpation guidance regardless of the level of experience and did not improve with the use of US guidance.

Accuracies of the LMGI in cadaveric models have been investigated in two studies [96,97]. Heidari et al. [97] examined whether an anteromedial or anterolateral approach is better when injecting into the ankle joint. They used 76 ankles from 38 cadavers and noted accuracies of 77.5% and 86%, respectively ($p = \text{NS}$). Another paper investigated anterolateral and postero-lateral approaches to the subtalar joint [96]. Twenty-three of 34 anterolateral injections (68%) and 31 of 34 (91%) postero-lateral injections were successful ($p = 0.016$).

USGI have been investigated in three studies using cadaveric models [98–100]. Wempe et al. [98] injected the first Metatarsophalangeal (MTP) joint in five cadavers and found that the USGI showed 100% accuracy and that the latex injectate penetrated to the articular surfaces of the metatarsosesamoid articulations. Smith et al. [99] examined the accuracies of three injection techniques to the posterior subtalar joint. They found that anterolateral, postero-lateral and postero-medial approach routes can be used equally by USGI with a high degree of accuracy. Finally, the paper by Reach et al. [100] revealed that USGI are 100% accurate when injecting into the first and second MTP joints, ankle, Achilles peritendinous space, posterior tibial tendon sheath and flexor hallucis longus sheath. The subtalar joint injections were 90% accurate.

The cadaveric studies confirm that USGI to the foot and ankle joints and soft tissues are more accurate than LMGI (Table 8).

Synovial biopsies

Synovial biopsy and analysis of synovial tissue can provide valuable insights into the pathophysiologic mechanism, disease status, therapy effect and prognosis of inflammatory joint diseases. The most famous instrument for landmark-guided needle biopsy of synovium was designed by Parker and Pearson [101], although its use is largely confined to the knee joint. The success rate reported for collecting sufficient synovial tissue was 95% in the knee joint, but lower rates (81%) have been published by Schumacher and Kulka in the elbow, wrist and ankle joints using this needle [102]. Moon et al. used a Franklin–Silverman needle in elbow, wrist and ankle biopsies, and the success rate was 68% [103]. Needle biopsy of the synovium can be performed under fluoroscopy control using the Tru-Cut needle. Beaulé et al. obtained samples from the glenohumeral, elbow, wrist, hip, knee and ankle joints using the Tru-Cut needle and reported a success rate of 81% [104]. The difference in the success rates of synovial biopsy is because of technical difficulties, variable intra-articular inflammation of the synovium and the fact that the operator is blind the macroscopic features of the synovium.

Table 8

The mean accuracies of the injections to the ankle and foot joints and soft tissues in 12 studies. The number of studies are given in parentheses.

	LMGI (%)	USGI (%)
Ankle joint: clinical comparative study (1)	58	85
Ankle joint: clinical LMGI study (2)	71.5	–
Ankle joint	85	100
Sinus tarsi	35	90
Cadaveric comparative study (1)		
Ankle and subtalar joints	100	100
I, II tarsometatarsal joint	27	64
Cadaveric comparative study (1)		
Peroneal tendon sheaths	60	100
Cadaveric comparative study (1)		
Posterior leg compartments	90	88
Cadaveric comparative study (1)		
Ankle joint: anteromedial	77.5	–
Ankle joint: anterolateral	86	–
Cadaveric LMGI study (1)		
Subtalar joint: anterolateral	68	–
Anteroposterior	91	–
Cadaveric LMGI study (1)		
MTP 1 joint	–	100
Cadaveric USGI study (1)		
Subtalar joint: different approaches	–	100
Cadaveric USGI study (1)		
I, II MTP, ankle, Achilles peritend, tibialis posterior, flexor hallucis longus tendon sheaths. Subtalar joint	–	100
Cadaveric USGI study (1)	–	90

MTP, Metatarsophalangeal; LMGI, landmark-guided injection; USGI, ultrasound-guided injection.

Koski et al. developed a US-guided biopsy method using portal and forceps, and representative synovial tissue in adequate amounts for histopathological evaluation was obtained from 33 of 37 cases, showing a success rate of 89.2% [105]. Kelly et al. recently demonstrated that an US-guided needle biopsy approach could be successfully used in a large prospective clinical study to harvest good quality synovial tissue and RNA from patients with early inflammatory arthritis [106]. Ninety-three sequential biopsy procedures were assessed from a total of 57 patients. Five different joint sites were biopsied (knee, elbow, wrist, metacarpophalangeal and proximal interphalangeal). No significant complications were reported following the procedure. No difference in pain, swelling and stiffness of the biopsied joint before and after the procedure was demonstrated. A median of 14 biopsy samples was retrieved from each procedure, with 93% of biopsy procedures yielding good quality tissue. The accuracy of US-guided synovial biopsy is somewhere between those of the blind needle biopsy and two-portal system needle arthroscopy. Similar to the needle arthroscopy, US also reveals the synovial hypertrophy and vascularisation (using Doppler mode in US imaging), and the best biopsy site can thus be chosen. The advantage of the US-guided biopsy method of the synovium is that it can be performed not only in the knee but also in most joints of the body and even bursae and tendon sheaths [105,106]. The US-guided synovium biopsy methods are interesting alternatives to other biopsy methods. These methods are well-tolerated techniques that can be performed on an outpatient basis with low running costs. However, there are currently limited data regarding the performance of US-guided synovial biopsy, mainly from observational studies. Thus, it remains critical to evaluate its performance within the clinical trials context against the current gold standard of arthroscopic biopsy [107].

Summary

US is the most applicable and feasible imaging modality for routine clinical use in guiding musculoskeletal procedures. Although many studies have examined the role of imaging guidance for injection, more studies are required regarding the use of US prior to injection that can alter the pathological and anatomical diagnosis. This is a fundamental advantage of US over other imaging

modalities in guiding injection. We have examined the literature on landmark-guided and US-guided procedures in musculoskeletal diseases of the upper and lower limbs and of the SIJ. According to the clinical and cadaveric studies, the US-guided technique is more accurate than the landmark-guided technique in the glenohumeral, acromioclavicular, wrist, hand, hip, knee and foot joints, and in the tendons of the biceps, wrist, hand hip, knee and ankle. Synovial biopsies are more accurate using an US-guided method. In terms of accuracy, both methods work equally and the data are insufficient to conclude for superiority of US-guided techniques in subacromial bursa, sternoclavicular, elbow, ankle and SIJ injections. The superiority of the efficacy of USGI has been shown in a minority of the joint regions including the subacromial bursa, wrist and hand (tendon sheath injections included), and knee joint injections. Plantar fascia injection is equally efficacious using both methods. The data for other anatomical area are presently insufficient to make any further conclusions. There is a trend towards an expanded number of advanced applications of interventional musculoskeletal US that can also be performed by a rheumatologist.

Practice points:

- There is clear evidence that the accuracy of LMGI is sub-optimal, and USGI are more accurate in most anatomical areas
- Much less is known about the efficacy of the USGI.

Research agenda:

- More studies are required to assess how US alters the pathological anatomical diagnosis and the accuracy and efficacy of USGI in different anatomical areas.
- Which US-guided technique (direct or semi-guided) is the most appropriate in different anatomical areas and clinical settings requires further research.

Conflict of interest

Prof Kane, Dr. Koski: No conflict of interest.

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