Assessment of 3-T MRI using susceptibility-weighted imaging to detect and evaluate intra- or periarticular blood metabolites and meniscal tears of the knee

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Abstract

Purpose: The purpose of this study was to assess the suitability of susceptibility-weighted imaging (SWI) sequences using the 3T MRI-unit for assessment of potential intraarticular pathologies in patients with acute and chronic torsion trauma of the knee joint.

Material and methods: Sixty-three patients with subacute and chronic rotary knee joint trauma of either the left or right knee were studied using an Achieva MRI 3T device (Philips, Amsterdam, Netherlands). Ground truth was set by two expert radiologists with seven and 10 years of experience in musculoskeletal imaging. Readings were performed separately for meniscus and joint space including synovia, ligaments, and periarticular soft tissue. Haemorrhage was defined as any lesion that was either T1 or SWI positive, without proton density (PD)-hypointensity (calcification). A lesion was defined as any pathology/variant with any signal positivity of either T1, PD, or SWI.

Results: A total of 63 patients were included (F : M = 22 : 41). The median age of the cohort was 29 years (range 13 to 71 years). Thirty-nine patients showed a meniscal tear, and only three of them (7.7%) demonstrated a meniscal haemorrhage. A total of 18 patients suffered from a periarticular injury, and 16 patients (88.9%) demonstrated a concomitant periarticular haemorrhage.

Conclusions: These data suggest that SWI can be used for the diagnosis of intra- or periarticular blood metabolites because their potential have an impact on mechanical conflict with the surface of the knee joints, in particular the cartilage and their effect on malacic lesions, but it performs poorly in the detection of meniscal pathologies.

Key words: MRI imaging susceptibility, periarticular space, meniscus knee, haemorrhage.
Advantage of SWI sequence in posttraumatic knee MRI

Knee entities include conventional spin-echo T1-weighted image (T1WI), fast spin-echo proton density-weighted, and gradient-echo. The most used sequence for the meniscal inspection is a fast (or turbo) spin-echo, intermediate-weighted fat-suppressed sequence.

Susceptibility-weighted imaging (SWI) is now available on commercial MR scanners and is used widely in brain imaging. This sequence takes advantage of the difference in the magnetic susceptibilities of tissues, achieving a new type of contrast that differs from those obtained with conventional T1- and T2-weighted MR imaging [9]. SWI is extremely sensitive to changes of susceptibility and enhances the contrast between tissues with different susceptibilities. Studies that focus on its use in the knee-joint are lacking. SWI is by definition sensitive to paramagnetic susceptibility effects, in particular blood, calcification, haemosiderin, and slow flow in vascular structures [10]. Therefore, SWI is a potentially useful imaging tool in the demonstration of menisci, cartilage, subchondral bone, ligaments, and muscles. The aim of the present study was to compare SWI and pre-existing MRI techniques for detection of lesions, haemorrhage, and meniscal tear, particularly T1WI and proton density-weighted imaging.

Material and methods

Data acquisition

During the period from January to August 2017 we prospectively collected all posttraumatic cases of routine knee MRI in our institute. Inclusion criteria were the following: all patients who suffered a knee trauma of any kind, regardless of whether the cause was from a direct trauma or a rotary accident. We included all age groups that were sent to us for assessment of post-traumatic consequences such as bone bruise, torn ligament, or meniscal tear. Exclusion criteria were the following: Patients with indication for knee MRI other than trauma, e.g. inflammation, infection, postoperative, rheumatoid arthritis, and bursitis. The MRI was performed on an Achieva MRI 3 Tesla unit (Philips, Amsterdam, Netherlands). For this continuously acquired representative population, we added an additional SWI transversal sequence, as a basic source, to the standard routine knee MRI protocol. VSWI, the 3D reconstructions based on the SWI transversal sequence, allowed calculation of the coronal and sagittal images. The entire MRI study consisted of a 26-minute protocol, with parameters shown in Table 1.

No contrast media was administered. Institutional Review Board (IRB) approval was waived due to the implementation of a well-established SWI clinical sequence into the clinical routine protocol.

Ground truth

Acquired MR images of knee joints were analysed by two radiologists with seven and 10 years of experience in musculoskeletal MRI imaging. For the ground truth, both readers had to agree on the signal intensity independently.

Study

Two additional general radiologists, with seven and three years of experience in musculoskeletal MRI imaging, reviewed the images in consensus for signal intensities.

The readout was performed on a GE-workstation (General Electric, Milwaukee, Wisconsin, USA) for two anatomical regions: 1) meniscus was analysed separately; and 2) the rest of the articular joint space including all ligaments (cruciate, collateral, patellar, and femoropatellar ligaments), synovia, capsule, and periarticular soft tissue.

All readers were blinded to the diagnosis and had to score the signal intensity on T1, proton density (PD), and SWI sequences for both anatomical regions separately. T1 and PD hyperintensity and SWI hypointensity in both regions were recorded and equalled positivity for haemorrhage. Any signal change (hyper- or hypointense) in any sequence was classified as positivity for any pathology (see Statistics below).

Statistics

Table 1. Magnetic resonance imaging examinations protocol

| Parameter               | PD SPAIR coronal | T1 TSE coronal | PD SPAIR sagittal | PD SPAIR transversal | T1 TSE transversal | PD VISTA SPAIR sagittal | SWI transversal |
|-------------------------|------------------|----------------|------------------|----------------------|--------------------|------------------------|----------------|
| TR (ms)                 | 3537             | 633            | Shortest         | 6289                 | 633                | 1300                   | 31             |
| TE (ms)                 | 30               | 20             | 30               | 30                   | 20                 | 32                     | 32 = 7.2/delta TE = 6.2 |
| Slice thickness (mm)    | 3                | 3              | 3                | 3                    | 3                  | 0.7                    | 2              |
| Distance factor (%)     | 0.3 mm           | 0.3            | 0                | 0.3                  | 0.3                | ~0.35                  | ~              |
| FOV (mm)                | 160 × 160        | 160 × 160      | 160 × 361        | 160 × 160            | 170 × 170          | 140 × 159              | 230 × 189      |
| Resolution              | 320 × 277        | 384 × 319      | 160 × 160        | 292 × 224            | 380 × 312          | 200 × 229              | 384 × 316      |
| Voxel size (mm)         | 0.5 × 0.53       | 0.417 × 0.5    | 0.55 × 0.69      | 0.55 × 0.69          | 0.45 × 0.536       | 0.7 × 0.69             | 0.6 × 0.6      |

PD SPAIR – proton density-weighted imaging, T1 TSE – weighted imaging, SWI – susceptibility-weighted imaging
A general lesion was defined as any pathology/variant with either PD, T1, or SWI hyper- or hypointensity. Haemorrhage in the periarticular space was defined as any lesion with either T1 hyperintensity or SWI hypointensity, without conspicuous PD-hypointensity (calcification). Meniscal haemorrhage was defined as T1 hyperintensity and SWI hypointensity, without conspicuous PD-hypointensity (calcification) (Figure 1).

Any signal change in any acquisition plane was defined as hyper- or hypointensity for the whole sequence. For the ground truth, both readers had to agree on positivity independently.

Sensitivity, specificity, and accuracy for haemorrhage and any lesion were calculated for T1, PD, and SWI, respectively, using the reader’s detection rate against the ground truth. To compare significant differences of the sequences, Fisher’s exact test was applied using MedCalc® version 7.6.0.0 (MedCalc Software, Mariakerke, Belgium). A significance level of $p < 0.05$ was utilised.

**Results**

A total of 63 patients were included ($F : M = 22 : 41$). The median age of the cohort was 29 years (range 13 to 71 years). Thirty-nine patients showed a meniscal tear, and only three of them (7.7%) demonstrated a meniscal haemorrhage (Figure 1). A total of 18 patients suffered from a periarticular injury, and 16 patients (88.9%) demonstrated a concomitant periarticular haemorrhage (Figures 2 and 3).

**Periarticular joint space**

SWI demonstrates best sensitivity for detection of haemorrhage in the periarticular joint space: Sensitivity...
Advantage of SWI sequence in posttraumatic knee MRI

for T1, PD, and SWI was 25.0, 62.5, and 100%, respectively (SWI vs. T1: \( p < 0.0001 \) and SWI vs. PD: \( p = 0.018 \)). All three sequences scored specificities > 95%, leading to the following accuracies for T1, PD, and SWI: 81.0, 87.3, and 96.8%, respectively (SWI vs. T1: \( p = 0.009 \)). SWI also showed the best results for any lesion detection in the (peri)articular joint space: accuracy for T1, PD, and SWI was 77.8, 90.2, and 93.7%, respectively (SWI vs. T1: \( p = 0.02 \) and SWI vs. PD: \( p = 0.086 \)). Sensitivity of T1 to detect any lesion in the periarticular joint space was significantly lower for T1 (22.2%) than that for PD (66.7%, \( p = 0.018 \)) and SWI (88.9%, \( p < 0.0001 \)) (Tables 2 and 3).

**Meniscus**

Haemorrhage in the meniscus was best scored on SWI (\( p < 0.0001 \)). Accuracies for T1, PD, and SWI were 42.9, 34.9, and 98.4%, respectively. While all sequences depicted meniscal

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**Table 2.** Sensitivity, specificity, and accuracy for periarticular pathology diagnostic in T1, proton density (PD), and susceptibility-weighted imaging (SWI) sequences

| Pathology (Periarticular) Joint space | T1 | PD | SWI | T1 | PD | SWI |
|--------------------------------------|----|----|-----|----|----|-----|
| **Haemorrhage**                      | %  | %  | %   | %  | %  | %   |
| Sensitivity                          | 25.0 | 62.5 | 100.0 | 22.2 | 66.7 | 88.9 |
| Specificity                          | 100.0 | 95.7 | 95.7 | 100.0 | 100.0 | 95.6 |
| Accuracy                             | 81.0 | 87.3 | 96.8 | 77.8 | 90.5 | 93.7 |

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*Figure 2.* Post-traumatic case with intra- and periarticular haemorrhage. On A) axial susceptibility-weighted imaging (SWI) sequence and C) sagittal SWI excellent visibility of the periarticular haemorrhage with bold hypointensity in the anterior capsule (blue arrow) and intra-articular blood components (hypointense dots, orange arrow), B) corresponding axial proton density (PD)-weighted, and D) sagittal PD-weighted imaging without post-traumatic periarticular blood components. Postoperative changes (red arrows)
Figure 3. Post-traumatic haemorrhage displaying blood on all sequences. A) Axial susceptibility-weighted imaging (SWI) and C) sagittal SWI display intra-articular blood level with typical fat/fluid level (blue arrow) also visible on the B) axial T1-weighted image (hyperintense fat layer on top of fluid [orange arrow]) and corresponding hypointensity on the D) sagittal proton density-weighted image.

Table 3. P-value board comparing sequences for joint space diagnostic in T1, proton density (PD), and susceptibility-weighted imaging (SWI) sequences.

|                      | Haemorrhage vs. T1 | PD | Any lesion vs. T1 | PD |
|----------------------|--------------------|----|------------------|----|
| Sensitivity          | SWI < 0.0001       | 0.018 | SWI < 0.0001     | 0.223 |
|                      | PD 0.073           | PD 0.018 |                       |     |
| Specificity          | SWI 0.490          | 1.000 | SWI 0.490        | 0.490 |
|                      | PD 0.490           | PD 1.000 |                       |     |
| Accuracy             | SWI 0.009          | 0.095 | SWI 0.020        | 0.744 |
|                      | PD 0.470           | PD 0.086 |                       |     |

Furthermore, while SWI demonstrated a remarkable detection rate for haemorrhage and other lesions in the periarticular joint space, it demonstrated the lowest score for any meniscus lesions (Figure 3).

PD demonstrated the best accuracies for diagnosis of any lesion in the meniscus. The accuracy results for T1, PD, and SWI were 82.5, 90.5, and 44.4%, respectively (Figure 2). The SWI accuracy was significantly lower than that of T1 and PD (p < 0.0001) (Tables 4 and 5).

Discussion

The purpose of our research was to apply SWI sequences in post-traumatic patients and to image blood components and their quantity within the structures of articular cavity and periarticular area of the knee joint. The study revealed great specificity and sensitivity of SWI sequences in assessing blood metabolites; however, it did not reveal...
Table 4. Sensitivity, specificity, and accuracy for meniscal pathology diagnostic in T1, proton density (PD), and susceptibility-weighted imaging (SWI) sequences

| Pathology      | Haemorrhage | Any lesion |
|----------------|-------------|------------|
| Sequence       | T1 | PD | SWI | T1 | PD | SWI |
| Sensitivity    | %  | %  | %  | %  | %  | %  |
| Specificity    | 40.0 | 31.7 | 98.3 | 75.0 | 75.0 | 100.0 |
| Accuracy       | 42.9 | 34.9 | 98.4 | 82.5 | 90.5 | 44.4 |

Table 5. Sensitivity, specificity, and accuracy for meniscal pathology diagnostic in T1, proton density (PD), and susceptibility-weighted imaging (SWI) sequences

| P-value board comparing sequences for meniscus | Haemorrhage | Any lesion |
|-----------------------------------------------|-------------|------------|
| vs. T1 | PD | vs. T1 | PD |
| Sensitivity | SWI | 1.000 | 1.000 | SWI | <0.0001 | <0.0001 |
| Specificity | SWI | <0.0001 | <0.0001 | SWI | 0.050 | 0.050 |
| Accuracy | SWI | <0.0001 | <0.0001 | SWI | <0.0001 | <0.0001 |
| PD | 0.477 | 1.000 | 0.271 |
The SWI protocol does not exhibit appropriate sensitivity or specificity in diagnosing ACL injury. In our study, we did not obtain positive images of ACL injury when compared with those from standard MRI sequences [3]. We also did not visualise blood metabolites within injured ACL tendons.

Arthroscopic surgery of the knee joint is a gold standard in assessing intra-articular structures; however, routine examinations to date, such as MRI, constitute a diagnostic routine and may often have an influence on the decision concerning patient’s surgery, which is reflected in the patient’s quality of life and has an economic and diagnostic impact [16, 17]. An experienced physician is able to assess meniscal or ligament apparatus injury based on clinical examination just as well as via diagnostic imaging. However, physicians are unable to determine the quality and quantity of blood metabolites within the knee joint before and after surgery, which may have an influence on microtrauma of articular cartilage of the knee joint.

We are aware that when examining patients’ knee joints using MRI, the SWI sequence is not fundamental. However, the SWI sequence allows us to obtain additional information, which we have previously described, within the cartilage. The sequence also allows the detection of possible venous malformations, intra-articular haemorrhage, early degenerative changes of calcific nature, or venous malformations.

Conclusions

SWI can be used to diagnose intra- and periarticular blood metabolic detection, but its sequence fails to detect meniscal tears according to the present study.

Ethical standards and informed consent

Ethical approval (IRB) and informed consent were waived due to the retrospective nature of the study with irreversible anonymisation of patient identifiers.

Conflict of interest

The authors declare no conflict of interest.

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