Secondary Effects of the Rupture and Reconstruction of the Interosseous Talocalcaneal Ligament on the Peritalar Joints

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Source of support: This research was supported by grants from the Peking University International Hospital Research Funds (YN2016QN10)

Background: The interosseous talocalcaneal ligament (ITCL) is the main soft-tissue contributor to subtalar joint stability. The role of ITCL reconstruction in retaining this stability is minimally reported. Therefore, we conducted this study to investigate the effects of rupture and reconstruction of the ITCL on the subtalar and peritalar joints.

Material/Methods: This experimental study randomly divided 72 rabbits into 3 equal groups of 24 rabbits each. Group I underwent reconstruction surgery, group II underwent resection, and group III was the control group. The cartilages between the talocrural and calcaneocrural joints, and between the subtalar and talonavicular joints on both sides were assessed by gross observation, ink staining, histology, and immunohistochemistry at weeks 4, 8, 16, and 32, postoperatively.

Results: In group II, the quantitative ink staining analysis revealed degeneration of the articular cartilages on the talonavicular joint (T=2.070, P=0.038) and the posterior subtalar joint (T=2.121, P=0.034) compared with the 2 sides of the same rabbit at 4 and 8 postoperative weeks. Comparing the operated sides of all the groups showed the posterior subtalar joints (Hc=9.563, P=0.008) and talonavicular joints (Hc=9.714, P=0.008) had an obvious difference at postoperative week 4; and in the calcaneocrural joints (Hc=6.750, P=0.034), it was noticed at postoperative week 8. Histology and immunohistochemistry findings confirm these observations.

Conclusions: An ITCL resection can lead to the progressive degeneration of the talonavicular and posterior subtalar joints, while an ITCL reconstruction can be beneficial in restoring the stability of these joints, preventing or postponing their degeneration, and protecting the articular cartilages.

MeSH Keywords: Bone Diseases • Finite Element Analysis • Subtalar Joint

Full-text PDF: https://www.medscimonit.com/abstract/index/idArt/925292
Background

The interosseous talocalcaneal ligament (ITCL) is considered the major stabilizer of subtalar joints [1,2]. It is located in the tarsal sinus between the medial and posterior talocalcaneal joints. It originates from the sulcus talus with an insertion at the sulcus calcaneus, which is located anterior and medial to the posterior talocalcaneal joint capsule [3].

A rupture of the ITCL is often caused by severe injury to the subtalar joints. Although this incident is considered infrequent and minimally reported in literature, such issues are sometimes misdiagnosed as the sinus tarsi syndrome (STS) [4–6]. With an increased understanding of the hindfoot structure along with the recent advances in radiologic and arthroscopic diagnostics, the distinguishing characters of ITCL rupture and STS are gradually being recognized [5,7,8]. Despite the fact that researchers have been investigating other novel treatment options for such conditions recently, their results have not provided clear-cut evidence of their efficacy, with a minimal number of studied individuals [5,9,10].

The long-term outcomes of ITCL rupture are not clearly recognized; this is a key factor influencing the treatment strategy. The aim of the current investigation is to identify the degree of secondary degeneration of the subtalar and peritalar joints in ITCL rupture as well as comprehend the protective effects of ITCL reconstruction. This research was conducted to give helpful insights into novel treatment strategies for ITCL rupture. For the purposes of the present study, we wanted to investigate the effects of ITCL resection and reconstruction on subtalar joint stability. We hypothesized that an ITCL resection would negatively impact the stability of this joint, while an ITCL reconstruction could result in beneficial effects and successfully lead to the restoration of joint stability. The null hypothesis was that there would be no difference between the 2 procedures on restoring subtalar joint stability, while the alternative hypothesis was that there would be a statistically significant difference between the 2 procedures in terms of subtalar stability.

Material and Methods

A total of 72 skeletally-mature New Zealand white rabbits were used for this trial. Any rabbits with previous surgery in the studied joints were excluded. The rabbits had an average weight of 2.79 kg (range 2.5–3.5 kg). There were 43 female rabbits (60%) and 29 male rabbits (40%). They were randomly divided into 3 equal groups of 24 rabbits each. Group I un

In group I, a 1-cm incision was made on the medial aspect of the knee joint. The semitendinosus muscle was identified and cut 3 cm in length along the border of the muscle’s tendon to its insertion site on the tibia (Figure 1A). The strand of semitendinosus was woven with a 3-0 polyester thread at both ends, which was long enough to be pulled and fixed (Figure 1B). The graft was stored in normal saline gauze. Then, a transverse incision was made on the dorsum of the foot of the corresponding side and the sinus tarsi were accessed. The ITCL was exposed and resected with a no.11 scalpel under visualization. The articular cartilage and other important structures were carefully protected. A 2 mm-diameter bone tunnel was drilled from the talus superior medially to the calcaneus inferior laterally, which passed through the normal position and direction of the ITCL. The graft was then pulled through the bone tunnel. The talar terminal was fixed with a 1.5 cm-diameter steel-wire button (Figure 1C). A thinner tunnel was drilled along the posterior aspect of the calcaneus. One strand of the thread from the calcaneal terminal was passed through the tunnel and fastened with another thread to form a bone-bridge fixation while the graft maintained tension (Figure 1D). Finally, hemostasis and irrigation of the incision site were performed with 0.02% povidone iodine and normal saline, respectively. The incision was closed with interrupted silk sutures.

Similar to group I, the incision was made on the dorsum of the foot for the rabbits in group II. The ITCL was exposed and resected under visualization [11]. Hemostasis, irrigating, and closure were identical to that described for group I.

Similar to groups I and II, the same incision was made in the control group. The ITCL was exposed and kept intact. Hemostasis, irrigation, and closure were identical to those described for groups I and II. In all the groups, the duration of the surgical procedures was 1 h for each rabbit. Liquid paraffin was embrocated on the prepared area of the skin. The rabbits were returned to their cages after surgery. Penicillin was injected intramuscularly for 3 days to prevent postoperative infection. There was no postoperative leg immobilization.

Six of the 24 animals in each group were sacrificed at postoperative weeks 4, 8, 16, and 32, respectively. One side of the intact foot was chosen randomly for the computed tomography (CT) scan. Meanwhile, both sides of the operated feet from all 3 groups were carefully dissected and examined. The 6 joints along with the 12 articular surfaces of every foot were split and observed. These joints and articular surfaces were the talocrural joint (Cr-Ta is the talar surface of the crus, Ta-Cr is...
the crural surface of the talus), calcaneocrural joint (Cr-Ca is the calcaneal surface of the crus, Ca-Cr is the crural surface of the calcaneus), anterior, medial, and posterior talocalcaneal joints (aTa, mTa, and pTa are the anterior, medial, posterior calcaneal surfaces of the talus; aCa, mCa, and pCa are the anterior, medial, and posterior talar surfaces of the calcaneus, respectively), and the talonavicular joint (Ta-Na is the navicular surface of the talus, Na-Ta is the talar surface of the navicular). The chondral changes in the subtalar and peri-subtalar joints were analyzed by gross observation, ink staining, histology, and immunohistochemistry.

After gross observation, the articular surfaces were stained twice with India ink. A Yoshioka score system [12] was used to quantify the ink-staining results. The sections were fixed with 10% formalin and embedded immediately in paraffin after resection. The specimens embedded in paraffin were cut in 5 μm-thick sections. The staining was hematoxylin and eosin (HE) for routine study as well as a toluidine blue stain to evaluate the loss of protein-polysaccharide concentration in the cartilage ground substance. The histological results were analyzed using the Mankin scoring system [13]. It was originally designed to assess the 4 histological parameters: cartilage structure or erosion, cell distribution or spatial arrangement of chondrocytes, Safranin-O staining, and tidemark intensity, with separate subcategories. The sums of the separate scores range from 0 (normal) to 14 (severe osteoarthritis). In the present study, we used toluidine blue staining instead of Safranin-O staining in the Mankin scoring system because of the higher affinity of toluidine blue to the sulphur content in the cartilages [14]. The changes in collagen formation in the cartilage including types I, II, and III were assessed by immunohistochemical assays.

Statistical analysis

All statistical analyses were conducted using the Statistical Package for Social Sciences (IBM SPSS Statistics Version 23, SPSS US Inc., Chicago, Illinois, US). Mankin scores at each stage
in all the groups were presented as means. We used the Kruskal-Wallis H test to determine the statistically significant differences of continuous variables between 2 or more groups [15]. We used the non-parametric Rank Sum test (Wilcoxon test) to estimate the statistical differences between the 2 paired groups. \( P<0.05 \) was considered statistically significant.

The protocol for this experimental study was reviewed and approved by the Institutional Review Board’s Animal Ethics Committee at our institution (0576).

Results

Gross findings

During the 4-week assessment period, group I (ITCL reconstruction) revealed good tension with no injury to the cartilage. In group II (resection group), obvious hyperemia with some hyperplasia of the synovium was noted; the color was normal on the cartilage surface while fissures and pits were not observed at week 4. Group III (control) revealed no hyperemia or edema of the synovium along with injuries to the cartilage.

During the 8-week assessment period, group I showed similar observations comparable to week 4 while group II showed hyperemia and hyperplasia of the synovium along with injuries to the cartilage. Group III, no hyperemia or edema was observed.

During the 16-week assessment period, group I showed no injury to the cartilage with good tension, while group II revealed slight hyperemia and edema of the synovium, and a spur had formed mainly on the posterior subtalar joint. In group III, no changes were observed.

![Figure 2. Specimen sacrificed at 32 weeks after resection surgery in group II: (A) A specimen of the talus and calcaneus. The black arrow shows the enormous osteophyte at the posterior subtalar joint. (B) A specimen of the crural surface of the navicular (Na-Ta). Ca-Cr – crural surface of the calcaneus; Cr-Ca – calcaneal surface of the crus; Cr-Ta – talar surface of the crus; Na-Ta – talar surface of the navicular; pSJ – posterior subtalar joint; Ta-Cr – crural surface of the talus.](image)
**Table 1.** Results of 2 sides of the 12 articular surfaces compared at every stage in group II by the Wilcoxon test (n=6).

| Articular surface | 4th week | 8th week | 16th week | 32nd week |
|-------------------|----------|----------|-----------|-----------|
|                   | T | P  | T | P  | T | P  | T | P  |
| Cr-Ta             | 0.000 | 1.000 | 1.414 | 0.157 | 1.342 | 0.180 | 1.732 | 0.083 |
| Ta-Cr             | 1.000 | 0.317 | 1.342 | 0.180 | 1.732 | 0.083 | 1.633 | 0.102 |
| Cr-Ca             | 1.000 | 0.317 | 1.414 | 0.157 | 1.732 | 0.083 | 1.342 | 0.180 |
| Ca-Cr             | 1.342 | 0.180 | 1.633 | 0.102 | 1.633 | 0.102 | 1.732 | 0.083 |
| pTa               | 1.000 | 0.317 | 1.342 | 0.180 | 1.857 | 0.063 | 2.070 | 0.038* |
| pCa               | 1.857 | 0.063 | 2.121 | 0.034* | 2.232 | 0.026* | 2.251 | 0.024* |
| mTa               | 0.000 | 1.000 | 0.000 | 1.000 | 1.000 | 0.317 | 1.414 | 0.157 |
| mCa               | 1.000 | 0.317 | 1.000 | 0.317 | 1.414 | 0.157 | 1.000 | 0.317 |
| aTa               | 1.000 | 0.317 | 1.000 | 0.317 | 1.414 | 0.157 | 1.000 | 0.317 |
| aCa               | 1.414 | 0.157 | 1.414 | 0.157 | 1.732 | 0.083 | 2.232 | 0.026* |
| Ta-Na             | 0.000 | 1.000 | 2.000 | 0.046* | 1.414 | 0.157 | 2.000 | 0.046* |
| Na-Ta             | 2.000 | 0.046* | 1.890 | 0.059 | 2.070 | 0.038* | 2.264 | 0.024* |

* Statistically significant difference; aCa – anterior talar surface of the calcaneus; aTa – anterior talocalcaneal joint; Ca-Cr – cranial surface of the calcaneus; Cr-Ca – calcaneal surface of the crus; Cr-Ta – talar surface of the crus; mCa – medial talar surface of the calcaneus; mTa – medial talocalcaneal joint; Na-Ta – talar surface of the navicular; P – P-value; pCa – posterior talar surface of the calcaneus; pTa – posterior talocalcaneal joint; T – T statistic of the Wilcoxin test; Ta-Cr – cranial surface of the talus; Ta-Na – navicular surface of the talus.

During the 32-week assessment period, group I revealed no change while group II revealed some hyperemia/edema of the synovium with obvious hyperplasia of the synovium. Moreover, fissures, fibrosis, and defects were visible on the surfaces of some cartilages, while florid spur formation was mainly observed at the posterior subtalar, talocrural, and talonavicular joints (Figure 2A–2C). In group III, no changes were observed.

### Ink staining

We performed the Wilcoxon test to compare both sides (operated and unoperated) of the same rabbit in group II (resection) and group III (control). The differences in the ink-staining scores presented as Yoshioka scores between the operated side and the contralateral side for all joints did not reach statistical significance in any assessment period (P>0.05). In group II, we noted differences between the talonavicular and posterior subtalar joints during postoperative weeks 4 and 8, respectively. On the other hand, statistically significant differences in the Yoshioka score (ink score) were noted between the talonavicular joint and the anterior subtalar joint (anterior calcaneal surface of talus [aCa]) in the 32nd postoperative week (Table 1). A comparison of the operated sides of all 3 groups by the Kruskal-Wallis test showed results (differences) similar to the ink-staining scores of the examined joints (Table 2).

### Histology and immunohistochemistry

At all assessment periods, the articular cartilages of the 2 sides of the control group and the non-operated side of groups I and II were mostly normal, their Mankin scores were 0–1. The results of the HE staining revealed a smooth surface layer, well-arranged and well-distributed chondrocytes, the lack of a cluster, a complete tidemark, and an even distribution of the toluidine blue stain implying that the degree of degeneration was minimal.

The immunohistochemistry staining showed that type I collagen was localized only in the subchondral bone, type II collagen was evenly distributed, and type III collagen was absent. The histological results showed the cartilage of the operated side in group I was mostly normal. In group II, the chondral injuries became more obvious as the postoperative duration increased. The severity of the chondral injuries varied in the different articular surfaces of the leg, among which the injuries in the posterior subtalar joint and the talonavicular joint were the most concerning (high Mankin degeneration score). The alterations observed in the chondral histology of the calcaneal surface of the posterior subtalar joint at every follow-up time point are presented in Table 3.
In week 4, the result of the chondral histology was close to normal or gently injured (Mankin scores 2–5). The surface layer was not flat and showed a small fissure or a little cluster of chondrocytes. A loss of toluidine blue was seen on the surface layer indicating the failure of dye uptake implying cellular death. We noted positive staining of type I collagen at the surface layer; type II collagen was noted inside the cartilage and around the chondrocyte, respectively.

In week 8, the proportion and degree of chondral injury increased (Mankin scores 6–8). The changes noted were that the surface layer was not flat, a small fissure was present; a little cluster of chondrocytes was found. A loss of toluidine blue was not well-distributed. Type I collagen was detected in the surface layer, type II collagen was distributed in the middle and deep layers, and type III collagen was only found in the injured area.

In week 16, the proportion and degree of chondral injuries increased to a greater degree (Mankin scores 9–11). The surface layer of the cartilage began to show fibrosis and the gap in the middle layer, which was a small fissure in week 4, was

Table 2. Results of the operated side of the 12 articular surfaces compared at every stage in the 3 groups by the Kruskal-Wallis test (n=6).

| Articular surface | 4th week | 8th week | 16th week | 32nd week |
|-------------------|----------|----------|-----------|-----------|
|                   | H<sub>C</sub> | P       | H<sub>C</sub> | P       | H<sub>C</sub> | P       | H<sub>C</sub> | P       |
| Cr-Ta             | 0.000     | 1.000    | 4.250      | 0.119   | 4.235      | 0.120   | 2.414      | 0.299   |
| Ta-Cr             | 2.000     | 0.368    | 4.235      | 0.120   | 4.406      | 0.110   | 7.159      | 0.028*  |
| Cr-Ca             | 2.000     | 0.386    | 4.250      | 0.119   | 4.250      | 0.119   | 2.528      | 0.283   |
| Ca-Cr             | 4.235     | 0.120    | 6.750      | 0.034*  | 4.406      | 0.110   | 2.970      | 0.227   |
| pCa               | 2.000     | 0.368    | 4.235      | 0.120   | 9.563      | 0.008** | 10.187     | 0.006** |
| pCa               | 9.563     | 0.008**  | 10.187     | 0.006** | 12.407     | 0.002** | 9.080      | 0.011** |
| mTa               | 0.000     | 1.000    | 0.000      | 1.000   | 2.000      | 0.368   | 4.250      | 0.119   |
| mCa               | 0.000     | 1.000    | 0.000      | 1.000   | 4.235      | 0.120   | 1.063      | 0.588   |
| aTa               | 2.000     | 0.368    | 2.000      | 0.368   | 2.267      | 0.322   | 2.267      | 0.322   |
| aCa               | 0.607     | 0.738    | 2.267      | 0.322   | 4.406      | 0.110   | 14.201     | 0.001** |
| Ta-Na             | 0.000     | 1.000    | 6.800      | 0.033*  | 1.286      | 0.526   | 4.628      | 0.099   |
| Na-Ta             | 9.714     | 0.008**  | 6.948      | 0.031*  | 6.611      | 0.037*  | 10.127     | 0.006** |

* Statistically significant difference; ** highly significant statistical difference; aCa – anterior talar surface of the calcaneus; aTa – anterior talocalcaneal joint; Ca-Cr – crural surface of the calcaneus; Cr-Ca – calcaneal surface of the crus; Cr-Ta – talar surface of the crus; Hc – H statistic of the Kruskal-Wallis test; mCa – medial talar surface of the calcaneus; mTa – medial talocalcaneal joint; Na-Ta – talar surface of the navicular; P – P-value; pCa – posterior talar surface of the calcaneus; pTa – posterior talocalcaneal joint; T – T statistic of the Wilcoxin test; Ta-Cr – crural surface of the talus; Ta-Na – navicular surface of the talus.

Table 3. Mankin scores for the posterior talar surface of the calcaneus and the talar surface of the navicular at every stage in groups I and II.

| Group  | Articular surface | 4th week | 8th week | 16th week | 32nd week |
|--------|-------------------|----------|----------|-----------|-----------|
| Group I (n=6) | pCa | 0.3 | 1.3 | 2.0 | 3.3 |
| Group I (n=6) | Na-Ta | 1.0 | 1.6 | 2.6 | 4.5 |
| Group II (n=6) | pCa | 3.0 | 6.0 | 9.7 | 12.0 |
| Group II (n=6) | Na-Ta | 4.3 | 6.3 | 10.7 | 12.0 |

Group I – interosseous talocalcaneal ligament reconstruction; group II – interosseous talocalcaneal ligament resection; pCa – posterior talar surface of the calcaneus; Na-Ta – talar surface of the navicular.

In week 4, the result of the chondral histology was close to normal or gently injured (Mankin scores 2–5). The surface layer was not flat and showed a small fissure or a little cluster of chondrocytes. A loss of toluidine blue was seen on the surface layer indicating the failure of dye uptake implying cellular death. We noted positive staining of type I collagen at the surface layer, type II collagen was noted inside the cartilage and around the chondrocyte, respectively.

In week 8, the proportion and degree of chondral injury increased (Mankin scores 6–8). The changes noted were that the surface layer was not flat, a small fissure was present; a little cluster of chondrocytes was found. The staining of toluidine blue was not well-distributed. Type I collagen was detected in the surface layer, type II collagen was distributed in the middle and deep layers, and type III collagen was only found in the injured area.

In week 16, the proportion and degree of chondral injuries increased to a greater degree (Mankin scores 9–11). The surface layer of the cartilage began to show fibrosis and the gap in the middle layer, which was a small fissure in week 4, was
now deep. The arrangement of cells was disturbed, the tide-mark was irregular, and the number of clusters of chondrocytes had increased. The loss of toluidine blue was seen in the surface and middle layers. Type I collagen showed diffuse distribution, type II collagen was mainly distributed in the middle and deep layers, type III collagen was mainly distributed in the moderate and deep layers.

In week 32, in group II, 1 rabbit died due to a rapid intravenous anesthetic injection, and 1 suffered a postoperative joint infection. After eliminating these, another 2 rabbits were selected for supplementation to the study samples. The scores of chondral injuries became higher, and severe chondral injuries were seen (Mankin scores 12–14). In the deep layers of the cartilage, there were large areas of fibrosis. In the same context, the number of cells decreased significantly, the tide-mark was not apparent, and the calcified cartilage layer was undistinguishable (Figure 3A–3C). The loss of toluidine blue was observed in a majority of the cartilage (Figure 3B). The staining of collagen types I and III were strongly and diffusely positive; type II collagen was distributed unevenly in the deep layers (Figure 3C–3E).

Discussion

The ITCL resection resulted in a significant degeneration of the articular cartilages of the study specimens during the early postoperative assessment period. The ITCL reconstruction revealed better outcomes in terms of less cartilage degeneration, good tension, and no gross injuries to the cartilage during all the assessment periods.

As yet, there is no agreement on the mechanism of the movement of subtalar joints, and thus, the effect of the ITCL on the stability of subtalar joints is uncertain [10,16]. In a series of in-vitro studies, Kjaersgaard-Anderssen [17,18] observed a moderate yet significant improvement in subtalar mobility following ITCL resections in the sinus tarsi. This group of researchers studied the 3-plane kinesiology of hindfoot instability among 20 amputation samples of lesions of ligamentous structures in both the sinus and canalis tarsi. The transverse resection of the ITCL improves the rate of the subtalar movement by about 21% in the horizontal plane (internal-external rotation), 16% in the frontal plane (pronation-supination), and up to 57% in the sagittal plane (dorsi- and plantar-flexion) with 43% improvement in dorsiflexion [18]. Tochigi [19] examined the
neutral-zone laxity as a measure of joint flexibility and resistance to applied force; it was examined prior to and following an ITCL resection. The neutral-zone laxity and flexibility were both improved in the axial-distraction test. The ITCL maintained apposition of the subtalar joints; it resulted in the stabilization of the subtalar joints in the face of the drawer forces that were applied to the calcaneus from the lateral side to the medial side. On the other hand, Knudson [20] examined 6 fresh-frozen foot samples with 2 stable zones (supination and pronation) and 2 transition zones. During the assessment period, the authors of the aforementioned study noted little displacement at the stable zones, and noted supination and pronation movements at the transition zones. Therefore, they concluded that the ITCL had significantly contributed to the stability of the subtalar joints, particularly during supination.

In the present study, the changes in the cartilage could be observed microscopically by ink staining while gross changes emerged after the ink staining. The method was validated in the knee joint [21]. We used it to evaluate the cartilages of the subtalar and peritalar joints, which can reveal the area of chondral injury along with the depth of the injury. The HE and toluidine blue staining display the histological changes in the microstructures of the cartilage (ie collagen fibrils and proteoglycans). The Mankin scoring system assesses the structure, cell, matrix, and tidemark at a cellular level. Immunohistochemistry staining was used to investigate collagen changes in the cartilage.

Clinical studies show that cartilage injuries commonly occur with an ITCL injury in a high proportion of the patients [5,7]. The results of our study showed that the talonavicular joint and the posterior subtalar joint were the first joints to be affected after an ITCL resection. Meanwhile, the severity of the degeneration was associated with the postsurgical duration. No obvious enantiomorphous changes were found. The injury on the 2 surfaces of the same articular side showed unequable behavior. The distal surfaces had a tendency to be more severely injured than the proximal surfaces. The underlying reason for this is still unclear.

The talonavicular joint is an ovoid joint, where the acetabulum pedis sockets the talar head. As defined by Klikkian et al. [22], the motion generated by a male ovoid surface moving on a female surface is that of a slide, roll, and spin. The roll is in the opposite direction of the slide [23]. Therefore, when the talus extends and flexes, the roll of the talar head in the acetabulum pedis changes to a slide. At the same time, in the sagittal vector, the talar displacement gradient changes from up to down in a superior-to-inferior direction. When the talus extends and flexes, the impingement occurs. The 2 surfaces are no longer consistent, which is the reason for faster degeneration of the talonavicular.

The subtalar joint also showed rapid degeneration. It bears 50% of the weight transferred from the crus in humans [24]. In rabbits, it also bears a large part of their weight. The changes in the talar displacement grads in the vertical and sagittal vectors lead to disagreement and degeneration of the 2 surfaces. When the ITCL ruptured, the talus descended and the calcaneocural joint bore more weight than an intact foot. That could be the reason that the degree of calcaneocurnal degeneration was faster than that observed in the talocrural joint. The anterior subtalar joint is in the sagittal plane and barely bears any weight in rabbits. A strong ligamentous capsule around the medial subtalar joint on the sustentaculum tali passes horizontally and ends on the medial aspect of the talus. This ligamentous capsule is similar to but much broader than the plantar calcaneonavicular (spring) ligament in humans. The structure, which shows fibrocartilaginous metaplasia in the area articulating the medial subtalar joint of the talus in a normal rabbit foot, plays a fundamental role in maintaining the talocalcaneonavicular connections. This contributes to sharing a great part of the weight transferred from the talus, and prevents its plantar and medial dislocation. This weakens the impact of the ITCL rupture on the anterior and medial subtalar joints; thus, the 2 joints showed little and delayed degeneration.

Reconstruction of the ITCL with a portion of the Achilles’ tendon inserted into a tubed Leeds-Keio artificial ligament was first reported by Kato [25]. Other reports showed satisfactory results of an ITCL reconstruction [5,26]. Due to the diagnostic difficulties (e.g. clinical measures do not reflect isolated subtalar instability, misdiagnosis, and radiographs have high false positive rates of ITCL ruptures), the ITCL rupture was not thoroughly studied before this. Magnetic resonance imaging (MRI) [27–29] and stressed radiography [9,30] provided valuable evidence to reach a diagnosis of ITCL rupture. Arthroscopy [5,7,8] could be the most reliable method in the diagnosis of this disease so far.

This study provided an animal model for a reconstruction of the ITCL with an autograft of the semitendinosus. The semitendinosus is a regular autograft for tendon transplant [31]. One strand of it is similar to the ITCL in diameter and intensity according to the results of our study for ligamentous biomechanics [32]. A steel-wire button referring to the EndoButton (Smith & Nephew, Andover, MA, USA) [33] was used to fix the talar terminal, which provided adequate tension for fixation and a decreased the risk of talar necrosis compared to the bone-bridge fixation of the calcaneal terminal. In group 1, none of the rabbits showed talar necrosis at any postoperative assessment point. The best choice for the autograft is the local tendon. We tried to construct it with 2 strands of the semitendinosus muscle tendon, in order to preserve the distal termination and fix the calcaneal side of the bone bridge. This can reduce the length of the incision on the knee and...
decrease the risk of infection by avoiding an implant button. We selected the semitendinosus muscle, as a dynamic stabilizer would have been destroyed after the posterior tibial tendon was cut, and it could affect the results of this experiment. It is worth mentioning that several factors could have affected the results of this study. During the arthrotomy of the subtalar joints, bleeding and injury to other structures can result in histological changes noted in the articular cartilage. A large number of cytokines in the effused blood could lead to the degeneration of the knee. The bone-tendon interface had no fixation and there was no healing between the interfaces during the early stage after reconstruction [34]. Therefore, when the foot moved, the graft shifted in the bone tunnel (Bungee’s effect and windshield wiper effect) [35], which affected the reconstruction. Perhaps the reconstruction could not completely simulate the biomechanics of the original ITCL.

Conclusions

An ITCL resection can lead to progressive degeneration of the subtalar and peritalar joints. The reconstruction of the ITCL is an effective option in recovering subtalar stability, protecting the articular cartilages, and preventing or delaying the degeneration of joints. Despite the fact that the biomechanics of human joints are not definitively the same as those of rabbits, this study provides a helpful insight into the importance of the ITCL for the stability of the subtalar and peritalar joints. Therefore, more robust studies on human subjects would provide higher-quality evidence about the role of ITCL reconstruction in retaining the subtalar joint stability.

Conflicts of interest

None.

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