

Long-term outcomes of muscle volume and Achilles tendon length after Achilles tendon ruptures

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Abstract

Purpose The best treatment for Achilles tendon (AT) ruptures remains controversial. Long-term follow-up with radiological and clinical measurements is needed.

Methods In this retrospective multicentre cohort study, patients ($n = 52$) were assessed at a mean of 91 months follow-up after unilateral AT rupture treated by open, percutaneous or conservative (non-surgical) treatment. Demographic parameters, time off work, maximum calf circumference and clinical scores (ATRS, Hannover, AOFAS) were evaluated. Muscle volume and cross-sectional area of the calf and AT length were measured on MR images and were compared between groups and to each patient's healthy contralateral leg.

Results Reduced muscle volume was found across all groups with a higher muscle volume in the conservative ($729.9 \pm 130.3 \text{ cm}^3$) compared to the percutaneous group

($675.9 \pm 207.4 \text{ cm}^3$, $p = 0.04$). AT length was longer in the affected leg (198.4 ± 24.1 vs. $180.6 \pm 25.0 \text{ mm}$, $p < 0.0001$) without difference in subgroup analysis. Clinically measured ankle dorsiflexion showed poor correlation with AT length ($R^2 = 0.07$, $p = 0.008$). Muscle volume strongly correlated with the cross-sectional area ($R^2 = 0.6$, $p < 0.0001$) but showed a weak correlation with the Hannover score ($R^2 = 0.08$, $p = 0.048$). Maximum calf circumference correlated with muscle volume ($R^2 = 0.42$, $p < 0.0001$).

Conclusions No significant difference between the treatment groups was found in muscle volume, AT length, clinical measures or days off work. Cross-sectional area and maximum calf circumference are cost-effective measurements and a good approximation of muscle volume and can thus be used in a clinical setting while clinical dorsiflexion should not be used.

Level of evidence III.

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Keywords Calf muscle volume · Cross-sectional area · Achilles tendon length · Achilles tendon rupture

Introduction

Achilles tendon ruptures are common in sports medicine and their incidence is increasing [1, 14]. Long-term sequelae of Achilles tendon ruptures can be grave: the injured athlete has to rest, and consequences such as therapy, rehabilitation costs and absence from work pose a substantial socio-economic cost. Thus, it is of pivotal importance, both on the individual and societal level, to identify those therapies with the best outcomes [28].

A common long-term problem in Achilles tendon rupture reconstruction, with substantial influence on overall outcome, has been muscle atrophy [13, 25]. A number of studies emphasized the problem of muscle atrophy and its functional implications on ligament and tendon repair, such as poor outcomes and increased risk of re-tears [2, 22, 32, 35]. About the ankle joint, Valderrabano et al. [35] showed muscle atrophy after reconstruction of posterior tibial tendon rupture. Gladstone et al. [4] and Liem et al. [15] showed that muscle atrophy and fatty infiltration are not reversed by successful tendon repair and play a significant role in determining functional outcome. Thus, it is important to assess muscle atrophy and fatty infiltration of the triceps surae after Achilles tendon rupture treatments.

Experts agree that in active patients, surgical treatment of acute Achilles tendon ruptures is superior to non-operative treatment because it allowed an anatomical reconstruction [1, 17, 20, 26, 29, 36]. However, to date, there is still lack of long-term data concerning muscle atrophy—both in volume and cross-sectional area—and fatty infiltration after different types of Achilles tendon rupture treatment, and it is unclear how such tissue degeneration would affect muscle force and clinical outcomes.

It was therefore the primary objective of this study to systematically assess muscle atrophy across the three treatment groups (open repair, percutaneous repair or conservative treatment) in a retrospective, assessor-blind, multicentre study at a minimum follow-up of 3 years. The second objective of this study was to test for an association between Achilles tendon length, muscular degeneration and clinical outcomes, including subgroup analyses by type of treatment (open, percutaneous, conservative). It was hypothesized that open surgery would perform best by means of muscle atrophy and fatty infiltration across the three treatment groups and that there was a significant negative association between muscle atrophy and fatty infiltration and clinical outcomes. It was also hypothesized that conservative treatment yields excessive Achilles tendons length after healing and that generally Achilles tendon

cannot be restored to their original length, independent of treatment type.

Materials and methods

This study was designed as a retrospective, assessor-blinded, multicentre cohort trial with consecutive patients. An IRB approval was granted for this study by the local institutional review board.

Fifty-two patients (females, 11 and males, 41) were recruited from three orthopaedic centres. Demographic data are shown in Fig. 1.

Patients post-unilateral Achilles tendon rupture were contacted a minimum of 3 years after the injury. Further inclusion criteria were: 20–65 years of age, a healthy contralateral leg with physiological, clinically determined alignment of the knee, foot and ankle, no trauma to the healthy leg, no neuromuscular impairments including muscle dystrophies, no other post-traumatic injuries or osteoarthritis of the knee, foot or ankle of the affected leg.

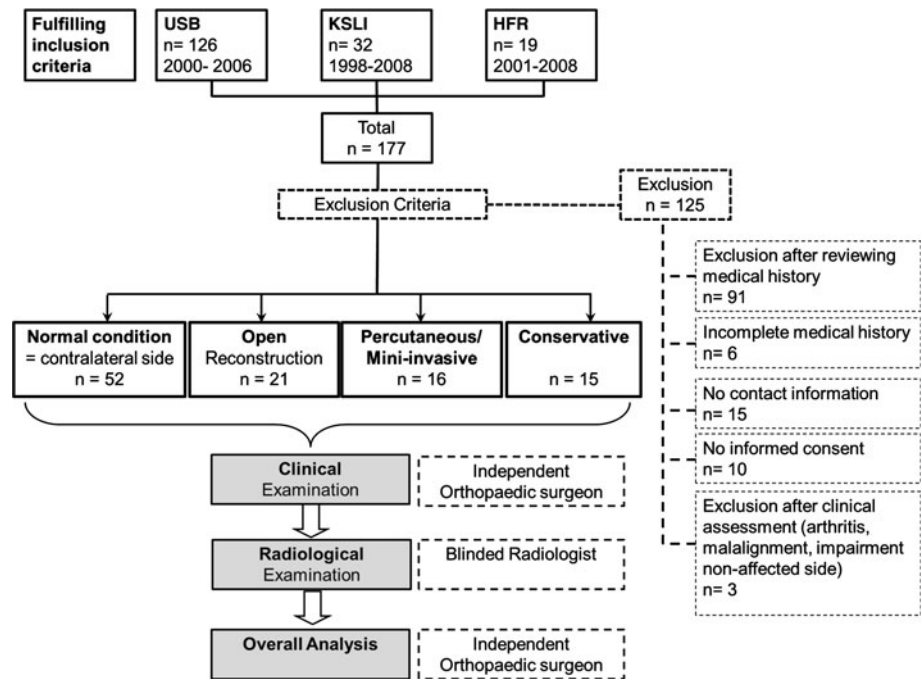
Exclusion criteria were: re-rupture or re-operation of the Achilles tendon, surgical site infection, neuromuscular diseases including muscle dystrophies, ankle valgus of more than 15° or ankle varus of more than 5°, other known pathologies of the non-affected leg, general MRI exclusion criteria (e.g. pacemaker, other metal not qualifying for an MRI, tattoos and claustrophobia) and a body mass index (BMI) of >40 kg/m². Conservatively treated patients were all recruited from one of the study centres.

The post-treatment (operative or conservative) regimens were identical in all study centres. Treatment was early functional rehabilitation in a stability boot and a night splint. After operative treatment or diagnosis (conservative), patients were allowed partial weight bearing for 2 weeks in 20° plantar flexion. For weeks 3 and 4, patients were allowed full weight bearing with 20° of plantar flexion and at week 5 and 6, patients were reduced to a 10° plantar flexing heel pad.

Demographic data and clinical assessment

The type of surgery was noted: open surgery, conservative treatment and percutaneous surgery which consisted of mini-open Kakiuchi [7] repairs, Ma and Griffith [16] repairs and repairs using the Achillon[®] device. The following clinical patient parameters were noted in a clinical assessment of an orthopaedic surgeon (C.R.): gender, age, body height, weight, BMI and activity level according to the Valderrabano sports score (VSS) [35]. Additionally, time off work, maximum calf circumference, thigh circumference 10 and 20 cm above the proximal patella apex, ankle dorsiflexion, SF-36 quality of life, proportion

Fig. 1 Study design. This flow chart depicts patient acquisition at the three hospitals, the number of exclusions as well as the pathway of examinations and analyses. *USB* University Hospital Basel, *KSLI* Canton’s Hospital Liestal, *HFR* Canton’s Hospital Fribourg. All centres are situated in Switzerland



of disabled pensions, Achilles tendon rupture score (ATRS, max. 100 points) [23], Hannover score (max. 100 points) [33] and the AOFAS Hindfoot score (max. 100 points) [11], and adherence of the tendon to the skin were evaluated.

Twenty-one patients were treated with open repair (OPEN), 16 with percutaneous/mini-open repair (PRCT) and 15 with conservative treatment (CONS). The mean follow-up was 91.0 ± 31.3 (mean \pm SD) months. Both sides were ruptured equally often ($n = 26$ in both groups). There was no significant difference in the average follow-up duration between the different treatment groups. Mean age at follow-up was 48.6 ± 8.7 years (range 23–55) with 49.9 ± 6.5 years in females and 48.2 ± 9.3 years in males (n.s.) without any significant difference between the groups. The mean body height was 176.8 ± 8.8 cm (range 157–196) with 166.5 ± 5.8 cm in females and 179.6 ± 7.2 cm in males ($p < 0.0001$) but equally distributed among the treatment groups. The mean BMI was 26.2 ± 3.9 kg/m² (range 19.7–39.4) with 25.7 ± 6.3 kg/m² in females and 26.4 ± 3.0 kg/m² in males (n.s.) and similar in all treatment groups. The SF-36 scores were similar in all groups (Fig. 2).

MRI assessment

MRI assessment was performed on both, the affected leg and the healthy contralateral leg, which served as a normal, matched control.

Imaging was carried out on a 3-TESLA scanner (MAGNETOM Verio, Siemens Medical Solution,

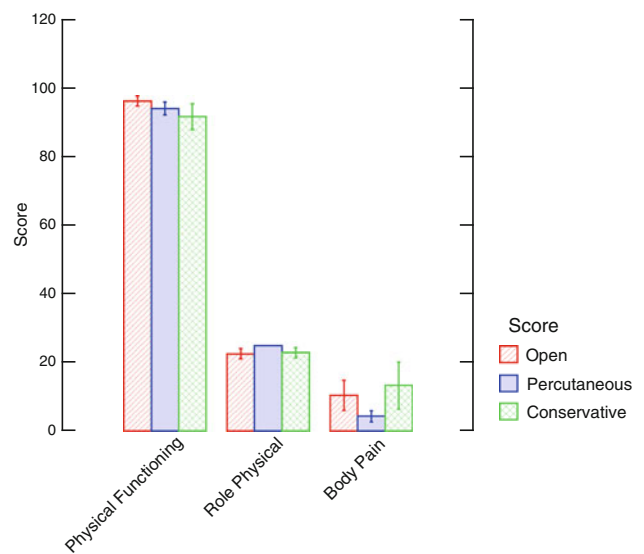


Fig. 2 SF-36 measures across treatment groups. The groups were not significantly different from one another. The box “population average” marks the line of the average score across the population. Selected measures from SF-36, short form 36; no significant differences in any measures were found

Erlangen, Germany). Patients were examined in the supine position with the ankle dorsiflexed at 90° (flexion neutral position) and the knee in 0° flexion and fixed in that position with straps and sandbags to assure reliable side comparability. A peripheral angiography matrix coil with 16 coil elements was used for image acquisition. Three sequences were acquired. First, a single-slab T1-weighted,

magnetization-prepared, rapid acquisition gradient-echo (MPRAGE) sequence (repetition time msec/echo time msec, 1,890/3.37; inversion time msec, 1,000; isotropic resolution, $1 \times 1 \times 1$ mm) was obtained in the coronal plane. The MPRAGE protocol was used to cover the entire lower leg from above the femoral condyles (origin of gastrocnemius muscles) through the bottom of the calcaneus allowing a 3D reconstruction with slices of 1 mm. The accuracy of those measurements was thus 0.1 cm. Second, axial T2-weighted fat-saturated fast spin-echo images (5,000/76; echo train length, 14; number of signals acquired, 1; section thickness, 3 mm; intersection gap, 0 mm; matrix size, 212×512 mm; field of view, 168×400 mm) were acquired of both legs simultaneously from the musculotendinous junction to the calcaneal insertion of the Achilles tendon. Third, axial T1-weighted fast spin-echo images (725/17; echo train length, 2; number of signals acquired, 1; section thickness, 6 mm; intersection gap, 4 mm; matrix size, 384×384 ; field of view, 200×200 mm) were performed separately for both legs resulting in approximately 65 slices per leg. Axial T1-weighted imaging covered the area from above the origin site of the gastrocnemius muscles (femoral condyles) to the calcaneus.

Quantitative and qualitative radiological measurements were noted. Muscle volume (MV) and cross-sectional area (CSA) were measured in the T1-weighted spin-echo images. Both parameters were measured on each calf muscle separately (soleus muscle (SOL), gastrocnemius medialis (GM) and lateralis (GL) muscles). The area of every muscle was measured on each slice from the distal appearance to the femoral condyles and multiplied by 1 cm (6 mm section thickness and 4 mm intersection gap) resulting in a close approximation of the MV (Fig. 3). Cross-sectional area was measured at a defined point at the junction of the proximal third to the middle third of the individually measured tibia length. The length of the Achilles tendon was measured on reconstructed isotropic images obtained by the MPRAGE sequence. The proximal point was defined as the last image before the appearance of the medial gastrocnemius head. The first image showing the AT attached to the calcaneus was defined as the distal point. This site is rarely involved in Achilles tendon rupture and can thus be measured in a ruptured, repaired or impaired tendon [31]. The number of slices between the distal and the proximal point was in turn the amount of millimetres the Achilles tendon travelled between the two points (slices of 1 mm). The resulting length is thus the length of the Achilles tendon.

Semi-quantitative parameters were used to assess the degree of tendon morphology and tendon sheath fluid. The degree of tendon degeneration and tendon sheath fluid were

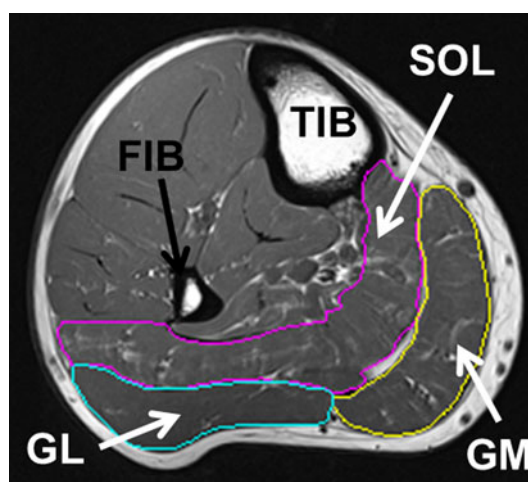


Fig. 3 Muscle volume measurement. The muscular areas of the gastrocnemius and soleus muscle were measured on each axial T1-weighted fast spin-echo image. The area was multiplied by the sum of the slice thickness and the intersection gap in order to obtain the total muscle volume. SOL, soleus muscle; GM, gastrocnemius medialis muscle; GL, gastrocnemius lateralis muscle; TIB, tibia; FIB, fibula

determined according to Valderrabano et al. on a four-point grading scale (none/0, mild/+, moderate/++, severe/+++) on axial T2-weighted fat-saturated fast spin-echo images [34]. Fatty atrophy was defined as originally published on CT images and adapted to MR images for rotator cuff muscle integrity [3, 5]. Radiological parameters were measured using the Santosoft DICOM Editor, Version 3.1.11, Athens, Greece.

Statistical analysis

The sample size was based on the precision of the volume measurements. Based on estimates of MV in three different treatment groups from the literature, an alpha of 5 %, a power of 80 % and a minimum sample size of $n = 43$ patients were required. Thus, after including 20 % ($n = 9$) to account for potential patient attrition or data loss, we included $n = 43 + 9$, or $n = 52$, healthy patients in this study. We used descriptive statistics including mean with standard deviation and frequencies to describe the populations, as appropriate. A one-way analysis of variance (ANOVA) with Bonferroni post hoc analysis was used to compare the parameters noted in Tables 1 and 2 except of skin palpation at the Achilles tendon, where a Chi-square test was used. For all comparisons, R^2 values were calculated in addition to the ANOVA p values. For comparison of the affected with the healthy side, a paired t test was used.

All reported tests were performed at the two-sided 5 % significance level. All analyses were performed with IBM SPSS Statistics Version 20.0[®], IBM Corporation, Armonk, NY, USA.

Results

Clinical assessment

For all treatment groups, days off work were not significantly different; even though it was striking that conservatively treated patients missed twice as many days of work as patients treated with open surgery (Table 1). When comparing surgically treated patients (open and percutaneous) with conservatively treated patients, no difference could be found (surgical, 48.3 ± 47.9 days (range 3–252); conservative, 75.4 ± 177.9 days (range 0–700), n.s.). Additionally, no disability pensions were paid out in any of the treatment groups.

In general, the clinical scores were good to excellent with only few outliers across the subgroups. The clinical scores showed good to excellent results without statistical significant difference (Table 1). Muscle volume correlated only to a small degree with the Hannover score but not with the ATRS or AOFAS hindfoot score (Table 3) as did fatty atrophy (Hannover, $R^2 = 0.03$, n.s.; ATRS, $R^2 = 0.04$,

n.s.; AOFAS, $R^2 = 0.10$, $p = 0.02$, respectively) showing little predictability of MV and fatty atrophy by clinical scores.

Maximum calf circumference (MCC) showed a significant difference between the affected and healthy leg (Table 1). Also, MCC was a good predictor of the degree of MV loss when the ratio of affected/healthy was evaluated ($R^2 = 0.37$, $p < 0.0001$). This was also true when looking at the absolute measures on the affected side and the healthy side (Table 3). The following algorithm gave a good estimate of the MV loss:

$$\text{Muscle volume (cm}^3\text{)} = 35 \\ \times \text{maximum calf circumference (cm)} - 566$$

($R^2 = 0.42$, $p < 0.0001$).

Adherent scars were seen to a higher degree (56 %) in patients treated percutaneously compared to 33 % in the open surgery group ($p = 0.01$). No patient had irritations of the sural nerve.

In a subgroup analysis on gender differences, clinical values did not significantly differ (not depicted).

Table 1 Patient demographics and clinical outcomes

	Treatment group				ANOVA <i>p</i> value	<i>t</i> test vs. healthy <i>p</i> value
	Open (<i>n</i> = 21) Mean (SD)	Percutaneous (<i>n</i> = 16) Mean (SD)	Conservative (<i>n</i> = 15) Mean (SD)	Total (<i>n</i> = 52) Mean (SD)		
Follow-up (months)	97.5 (30.8)	83.4 (34.2)	90.1 (29)	91 (31.30)	n.s.	n.a.
Age at follow-up (years)	48.2 (10.6)	46.4 (7.1)	51.3 (6.9)	48.6 (8.7)	n.s.	n.a.
Age at injury (years)	40.3 (10.1)	39.6 (7.7)	44.2 (7.4)	41.2 (8.7)	n.s.	n.a.
Days off work	38 (23)	62 (67)	75 (175)	56 (100)	n.s.	n.a.
Height (cm)	176.4 (9.2)	175.3 (8)	178.9 (9)	176.8 (8.8)	n.s.	n.a.
Weight (kg)	82.1 (13.7)	78.1 (13.8)	86.2 (9.8)	82.1 (12.9)	n.s.	n.a.
BMI (kg/m ²)	26.4 (4)	25.3 (3.2)	27.1 (4.3)	26.2 (3.9)	n.s.	n.a.
AOFAS score	97 (5.9)	93.2 (7.7)	93.4 (8.3)	94.8 (7.3)	n.s.	n.a.
ATRS score	90.5 (20.1)	88.4 (10.7)	90.6 (12.7)	89.9 (15.4)	n.s.	n.a.
Hannover score	86.7 (6.8)	80.2 (12.2)	79.1 (11.9)	82.5 (10.6)	n.s.	n.a.
Max.calf circumf. affected (cm)	38.1 (3.4)	37.7 (3.1)	37.7 (3.7)	37.9 (3.3)	n.s.	<0.0001
Max.calf circumf. healthy (cm)	39.6 (3.1)	39.1 (2.6)	38.8 (3.4)	39.2 (3)	n.s.	n.a.
Max.calf circumf. %	0.96 (0.04)	0.96 (0.03)	0.97 (0.03)	0.97 (0.03)	n.s.	n.a.
Thigh 20 cm above pat. affected (cm)	52.1 (5.5)	51.1 (4.6)	52.2 (5.2)	51.8 (5.1)	n.s.	n.s.
Thigh 20 cm above pat. healthy (cm)	52.6 (4.7)	51.3 (4.7)	52.7 (4.8)	52.2 (4.7)	n.s.	n.a.
Thigh 20 cm %	0.99 (0.04)	1 (0.01)	0.99 (0.03)	0.99 (0.03)	n.s.	n.a.
SF36 physical funct. functioning	96.2 (6.5)	94.1 (7.1)	91.7 (14.2)	94.2 (9.5)	n.s.	n.a.
SF36 role physical	22.6 (6.7)	25 (0)	22.9 (5.7)	23.4 (5.3)	n.s.	n.a.
SF36 bodily pain	10.5 (19.9)	4.4 (6.3)	13.3 (25.5)	9.4 (18.9)	n.s.	n.a.

% Percent of healthy side, SF36 short form 36, *circumf* circumference, ATRS Achilles tendon rupture score, AOFAS AOFAS hindfoot score, *pat* patella, * Chi square, *n.a.* not applicable, *n.s* not significant

Table 2 MRI assessment

	Treatment group				ANOVA <i>p</i> value	<i>t</i> test vs. healthy <i>p</i> value
	Open (<i>n</i> = 21) Mean (SD)	Percutaneous (<i>n</i> = 16) Mean (SD)	Conservative (<i>n</i> = 15) Mean (SD)	Total (<i>n</i> = 52) Mean (SD)		
<i>Muscle volume</i>						
Total affected (cm ³)	701.1 (141.2)	675.9 (207.4)	729.9 (130.3)	701.6 (159.9)	n.s.	<0.0001
Total healthy (cm ³)	853.7 (164.8)	863 (183.8)	833.6 (121.4)	850.8 (157.4)	n.s.	n.a.
Total %	0.82 (0.07)	0.79 (0.17)	0.88 (0.11)	0.83 (0.12)	n.s.	n.a.
<i>Cross-sectional area</i>						
Total affected (cm ²)	42.4 (8.7)	40.5 (9.8)	43.9 (7.1)	42.3 (8.6)	n.s.	<0.0001
Total healthy (cm ²)	46.8 (12)	44.8 (11.8)	45.8 (6.5)	45.9 (10.5)	n.s.	n.a.
Total %	0.94 (0.22)	0.94 (0.25)	0.96 (0.08)	0.94 (0.2)	n.s.	n.a.
<i>Tendon</i>						
Morphology affected sheath	0.38 (0.59)	0.5 (0.63)	0.43 (0.51)	0.43 (0.57)	n.s.	n.s.
Sheath fluid affected	0.19 (0.51)	0.13 (0.34)	0.2 (0.41)	0.17 (0.43)	n.s.	n.s.
<i>Achilles tendon length</i>						
Affected (mm)	195.6 (27)	199.6 (20)	201.1 (25)	198.4 (24)	n.s.	<0.0001
Healthy (mm)	177.4 (28)	179.4 (23)	186.3 (24)	180.6 (25)	n.s.	n.a.
%	1.11 (0.09)	1.12 (0.08)	1.08 (0.08)	1.1 (0.08)	n.s.	n.a.

%, Percent of healthy side; n.a., not applicable

MRI assessment

Significant MV loss of the affected leg compared to the matched, healthy leg was found across the three treatment subgroups (mean, 17 % reduction). There was a 9.1 % higher MV in the conservatively treated group as compared to the percutaneous treated group (Table 1). When comparing the operative groups (open and percutaneous together) with the conservative group, no difference could be found (n.s.). The subgroup analysis for the different muscles (SOL, GM, GL) did not show any significant differences. When looking at muscle atrophy as defined by fatty infiltration, the soleus and the gastrocnemius were significantly affected ($p < 0.0001$ and 0.006 , respectively). Analysis of the CSA revealed no significant difference in

the treatment subgroups as well as the muscle subgroups (SOL, GM, GL).

The mean Achilles tendon was significantly longer in the affected leg when compared to the healthy leg (198.4 ± 24.1 vs. 180.6 ± 25.0 mm, $p < 0.0001$) but not significantly different between treatment groups. In addition, there was no association between the type of treatment and MV (n.s.), CSA (n.s.), and degree fatty infiltration (n.s.).

Total MV of the healthy or affected side, respectively, did not correlate with the Achilles tendon length on the respective side. Clinically measured ankle dorsiflexion showed only a poor correlation with Achilles tendon length ($R^2 = 0.07$, $p = 0.008$), for this measurement, all legs and Achilles tendon lengths were evaluated ($n = 104$). The

Table 3 Predictors of muscle volume (R square)

	Muscle volume			
	Affected side (<i>n</i> = 52)	Health side (<i>n</i> = 52)	Total (<i>n</i> = 104)	Ratio (affected/healthy)
Cross-sectional area	0.79^a	0.49^a	0.60^a	0.0003
Maximum calf circumference	0.41^a	0.38^a	0.42^a	0.37^a
Hannover	0.08^a			
ATRS	0.049 ^a			
AOFAS	0.001 ^a			

Depicted are R^2 values

Hannover, Hannover Achilles tendon score; ATRS, Achilles tendon rupture score; AOFAS, American Orthopaedic Foot and Ankle Society Hindfoot score; total, $n = 104$

^a Significant

same weak correlation was found for the affected ($R^2 = 0.14$, $p = 0.006$) and the healthy side ($R^2 = 0.01$, n.s.) separately. Clinically measured passive dorsiflexion measurements did not show a difference between the treatment groups (open, $19.7^\circ \pm 3.2^\circ$; percutaneous, $19.9^\circ \pm 3.6^\circ$; conservative, $19.8^\circ \pm 6.7^\circ$; n.s.). In addition, the affected ($19.8^\circ \pm 4.5^\circ$) and the healthy side ($18.4^\circ \pm 5.7^\circ$) did not show a significant difference (n.s.). As expected, MV correlated with CSA (Table 3).

Discussion

This study focused on functional and imaging outcomes after Achilles tendon treatment with comparison of the three treatment mainstreams (open repair, percutaneous suture and non-operative treatment). No significant differences in MRI and clinical scores outcomes were found between the three treatment groups.

Controversy remains as to which treatment algorithm is best followed for acute Achilles tendon rupture. However, there is consensus that open reconstruction is best suitable for athletes [1]. Whichever treatment is chosen, sequelae from surgical site infections, scar adhesions and sural nerve irritation in operative techniques [37], higher re-rupture rates in conservative treatment [10] and deep vein thrombosis, muscle atrophy [6, 13, 25], days off work and loss of function for all treatments exist [25].

Muscle atrophy as a potential consequence of Achilles tendon ruptures has not been correlated with Achilles tendon length. These measurements were now performed on treated and healthy tendons and we found them to be healed in over length with a mean of 110 % compared to the healthy side showing that original tendon length cannot be restored independent of the type of treatment. Until now, Achilles tendon length was measured by implanting metal markers into the tendon and looking at the gapping or increase in ankle dorsiflexion in different post-operative rehabilitation protocols without measuring the actual Achilles tendon length, thus lacking direct comparison to our findings [8, 21, 22]. In a previous study, we evaluated Achilles tendon length [27] and showed that dorsiflexion of the ankle joint, although significant, is only a weak predictor ($R^2 = 0.07$) of Achilles tendon length and should thus not be used to assess Achilles tendon overlength after repairs [12, 21].

Muscle volume was found to be diminished in all treatment groups. Although open surgery is intended to result in better anatomic reconstruction, we found a MV loss of 18 % in the open surgery group. To our knowledge, MV of the posterior calf has never been measured in an accurate manner. When looking at fatty infiltration as an indicator of muscle atrophy, there was no significant

difference between the treatment groups either. Nevertheless, fatty atrophy was still present after more than 7.5 years of follow-up. With the supraspinatus muscle, it is already known that muscle atrophy does not completely reverse [15]. We can support this finding by looking at the posterior calf muscles. A recent study of Olsson et al. [24] revealed that major functional deficits persist even 2 years after Achilles tendon rupture and that patients do not improve significantly between 1 and 2 years after the rupture. Rebeccato et al. retrospectively compared 52 tendon ruptures with a minimal follow-up of 12 months. The results revealed a loss of posterior muscle compartment area of 18 % in the open group, 19 % with Ma-Griffith and 9 % in Kakiuchi repair group, respectively. While Haggmark et al. [6] compared 15 non-operatively with 8 surgically treated patients and showed a significant reduction in calf muscle area (no follow-up stated), Leppilahti et al. [13] showed calf muscle atrophy of 15 % on the affected leg in the analysis of CSA by means of CT scans (mean 3.1 years follow-up). In our findings, the CSA was higher after more than 7.5 years of follow-up although not significantly different. This might be due to the fact that patients might recover to a high degree even between 2.9 years (Rebeccato et al.) and 7.5 years (our cohort) of follow-up. The affected muscles and tendons might recover after such a long follow-up. Another fact that supports the recovery theory is the circumstance that we saw a reduction in MCC of -1.3 cm which was noted to be -2.55 cm in the study of Rebeccato et al. [25].

Although days off work did not significantly differ throughout the treatment groups, it was notable that there were almost twice as many days missed in conservative treatment compared to open surgery, though this might be due to the large standard deviation in the conservatively treated group. This should be considered when cost effectiveness of treatments is discussed. While conservative treatment might be cheaper in direct treatment costs, total societal cost might be equivalent if patients return to work consistently earlier after open treatment. Percutaneous treatment was situated in between the two. When compared to other studies, days off work have been reported to be in the same range [18, 25].

All clinical scores showed good to excellent results. These results were consistent with the literature stating the same ranges even after shorter follow-up periods [9, 30]. Although clinical scores might be a good measure for short- to mid-term outcome, it appeared that the discrimination of results in a long-term follow-up become more difficult in the very good to excellent range. The fact of less muscle atrophy in our long-term follow-up did not add to a better outcome by means of clinical scores.

Of great clinical interest might be two parameters: CSA and MCC both showing strong correlations with MV

(Table 3). The strong correlation between MV and CSA will make it easy to approximate MV by simple measurements on ultrasonography and replace the costly and time-consuming MRI measurements. This supports other studies, which have used the CSA for the approximation of calf MV [6, 13, 19, 25]. Even less costly is the clinical measurement of maximum calf circumference. In contrast to Haggmark et al., who showed a poor correlation, we found a strong correlation of cross-sectional area with muscle volume. This could be due to the fact that Haggmark only measured the CSA and not the MV of the whole posterior calf, giving less accurate results. When looking at correlations between CSA and maximum calf circumference, the latter seemed to be a good predictor of CSA on the affected side as well as for the healthy side.

Strengths of this study were the strict in- and exclusion criteria as well as the long follow-up time of 91 months. Radiologists and radiology assistants were blinded for the study. Additionally, the orthopaedic surgeon performing the follow-up measurements was independent and did not treat the patients before or after the study. For the first time in the literature, MV was measured for the triceps surae and correlated to cross-sectional area, Achilles tendon length and clinical parameters. Fatty atrophy grading according to Goutallier was introduced into calf muscles [5].

There were certain limitations to this study. Firstly, there might be a selection bias and patients that did not have a good outcome might not consent to be followed up. We tried to overcome this burden by contacting all consecutive patients from the database in the years stated in Fig. 1. Operative indications and treatment across centres in a multicentre setup might differ considerably making the operative groups heterogeneous. Additionally, some patients were lost due to the time-consuming examinations at different sites.

Conclusions

No significant difference between the treatment groups was found by means of clinical scores, days off work, clinical measurements, ATL, MV after more than 7.5 years of follow-up. No general recommendation can be given on the generally best treatment option even after examining consecutive patients. Ultrasonography and MCC are easy and cost-effective measurements and a very good approximation of MV of the posterior calf even in the untreated limb and can thus be used in daily clinical work.

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Conflict of interest Each author additionally certifies that he or she has no commercial associations that might pose a conflict of interest in connection with the submitted article. The authors indicate that they have no conflict of interest.

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