Effects of short-term treatment strategies over 4 weeks in Achilles tendinopathy

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Br J Sports Med 2007;41:e6 (http://www.bjsportmed.com/cgi/content/full/41/7/e6). doi: 10.1136/bjsm.2006.031732

Background: The therapeutic efficacy of non-surgical treatment strategies in Achilles tendinopathy (AT) has not been well clarified. Time-consuming and costly combinations of treatment for pain, physiotherapy and biomechanical procedures are often applied.

Objective: To analyse the efficacy of single therapeutic regimens commonly used over a short period of 4 weeks.

Methods: 31 male runners (mileage >32 km/week) with unilateral, untreated AT completed 4 weeks of either physiotherapy (10 treatments: deep-friction, pulsed ultrasound, ice, sensory motor training; (P)), wearing custom fit semirigid insoles (I) or remained without treatment (control group C). Before and after treatment, all patients underwent a treadmill test and a plantar flexion strength exercise. Subjective pain (Pain Disability Index, Pain Experience Scale), as well as strength performance capacity (peak torque), was analysed (mean, 95% CI, repeated measures analysis of variance, $\alpha = 0.05$).

Results: Pain was reduced to <50% of the baseline value after physiotherapy or after wearing insoles (p<0.05). Individual pain reduction was >50% (25%) in 89% (100%) of subjects in I and 55% (73%) in P. Higher eccentric plantar flexion peak torgues after treatment were observed in I and P.

Conclusions: Most patients with AT experience a reduction in pain after only 4 weeks of differentiated, nonsurgical treatment consisting of physiotherapy or semirigid insoles.

Problems of Achilles tendon overuse are cited as one of the major pathologies that reduce physical capacity in everyday living, occupation and sports.¹⁻³ Owing to the long duration of problems and widely varying, individual responses to treatment, the efficacy of single or combined therapeutic measures still remains a matter of debate.^{4 5} Thus, treatment is usually complex, time consuming and costly.^{3 5}

It has often been assumed that Achilles tendon problems are caused by an inflammation.⁶ However, recent histological studies show that inflammatory cells and mediators, usually present after acute mechanical stress, are absent in tendon overuse.^{7–9} On the other hand, high concentrations of glycosa-minoglycans and a loss of the hierarchical collagen structure have been found.^{10 11}

The mechanism of pain development is not well understood. Competing explanatory models describe increased mechanical tendon vulnerability, microruptures and the supplanting of collagen type I by type III. Pain is also ascribed to the mechanical irritation of ingrown nerve endings due to neovascularisation.^{1 9} Reduced perfusion is now considered less important since it was demonstrated that blood supply and oxygen extraction clearly increase during physical exercise.^{12 13} It thus seems certain that tendon tissue must be considered metabolically active to a far greater extent than has been assumed to date.^{8 9}

Pain reduction has traditionally been the main outcome variable of non-surgical treatment in Achilles tendinopathy (AT).³ In daily practice, local physiotherapeutic measures such as deep friction massages, ice and ultrasound are usually applied.^{14–16} However, despite broad acceptance and pain reduction in individual cases, scientific evidence of short-term physiotherapy is still lacking.¹⁴ Currently, adjuvant use of sensory motor training¹⁸ and eccentric exercises^{18–20} are increasingly being discussed. Shalabi *et al*²¹ demonstrated that eccentric exercises led to an improved clinical outcome, reduction in tendon diameter and reduced intratendinous

lesions. Alfredson *et al*^{18 19} had comparably good clinical results after 12 weeks of predominantly eccentric exercise training.

In addition to physiotherapy and training, custom-made insoles are frequently used, but evidence of their efficacy is still lacking.¹⁵ ¹⁶ Recent studies using bone pins have shown that the mechanical effect of insoles, understood as an alignment of the skeleton, is unspecific and only slight during walking and running.²² Currently, sensory motor effects of semirigid insoles are being discussed.²³ ²⁴

The aim of this study was to analyse whether standardised short-term physiotherapy or wearing individually fitted insoles over a period of 4 weeks reduces pain in patients with unilateral AT.

METHODS

Subjects

This randomised controlled trial was performed after approval by the local ethics commission in compliance with good clinical practice (Note for Guidance). In all, 63 patients with AT were initially identified as potential subjects at the university outpatient clinic (Medical Clinic, Department of Sports Medicine, University Clinic of Tübingen, Germany). Since it is known that the frequency of AT increases in male runners with a running distance of >32 km/week,¹⁶ only men (18–50 years, mileage>32 km/week), with unilateral chronic tendon problems during exercise, untreated for at least 6 months, were evaluated. Diagnosis was always made by the same orthopaedic surgeon (palpable nodes within the tendon itself, pressure pain at the mid-substance of the tendon). Patients with local inflammation (swelling, redness) or problems at locations other than the mid-substance of the Achilles tendon were

Abbreviations: AT, Achilles tendinopathy; C, control group; I, group wearing insoles; IAT, individual's anaerobic threshold; P, group undergoing physiotherapy; PDI, Pain Disability Index; PES, Pain Experience Scale

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Accepted 8 January 2007 Published Online First 26 January 2007

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Figure 1 Study flow chart of evaluated patients, participants and dropouts (DO).

excluded. After validation of inclusion/exclusion criteria, 31 subjects remained (fig 1).

Subjects were randomised into one of three groups, according to the planned intervention (control group C, those wearing insoles, I and those who underwent physiotherapy, P). There were three dropouts during the study. One subject (I) did not appear for baseline measurements. One subject (C) had a rupture of the tibialis posterior tendon due to recreational activities. Another subject (C) had a rupture of the Achilles tendon after inclusion. Twenty-eight subjects completed the study (fig 1, table 1).

Study protocol

Initially, a clinical examination was performed and study inclusion criteria were checked (U_1). Personal data, training data and history of pain were anonymously recorded on a case report form. To achieve individually comparable exercise levels during treadmill measurements (80% of the individual's anaerobic threshold (IAT)), patients underwent treadmill ergometry at the end of U_1 (start 8 km/h, increase 2 km/h, increment time 3 min).²⁵

The study protocol consisted of a test–retest procedure (M_1 , M_2), including an intervention phase of 4 weeks. After inclusion and before baseline measurement M_1 , a 2-week training and instruction phase (pain diary, training diary) was completed (fig 2). Both pain recordings and training data were taken at M_1 . M_1 began with another clinical examination to determine possible changes compared with U_1 . Then the subject's pain during activities of daily life (Pain Disability

	Control	Insoles	PT
Age (years)	38 (4.9)	35 (6.7)	41 (5.9)
Height (cm)	177 (3.9)	181 (8.7)	176 (5.9)
Weight (kg)	70.4 (7.3)	76.7 (6.7)	70.7 (7)
Mileage (km/week)	53.1 (10.6)	50 (13.5)	50 (13.6)
IAT (km/h)	13 (1)	13.7 (1.3)	13.3 (0.9)
Pain (months)	7.9 (6.8)	13.8 (16.5)	17.3 (18.7)

Index, PDI²⁶) followed by a first measurement of current pain experience (Pain Experience Scale, PES²⁷) was recorded. Finally, patients underwent a warm-up on the treadmill at 80% of the IAT.

Exercises consisted of a 20 min treadmill run (80% IAT) to elicit their running-specific pain and a subsequent plantar flexion strength test to increase further load on the Achilles tendon. At the end of each exercise, PES was recorded. Strength exercise was carried out concentrically and eccentrically (supine position, 30° knee flexion, Lido-Active, Loredan Biomedical, California USA). After cool-down (treadmill, stretching), the 4-week treatment was explained, a pain and training diary was given to the patients and post-treatment measurement M_2 was scheduled. In M_2 , pain and training diaries were checked for completeness before recording the PDI. Protocols M_1 and M_2 were analogous. Each subject underwent a thorough clinical examination.

Ttreatment and intervention

Standardised physiotherapy consisted of 10 single 30 min sessions (2 or 3/week over 4 weeks). Each included deepfriction massages at the mid-substance of the Achilles tendon, local pulsed ultrasound (1.5 W/cm²), ice application and sensory motor training. This training consisted of three sets of 15 repetitions of balance and stabilisation exercises on a stability pad and eccentric exercises (loading of the calf muscles by lowering the heel standing with the forefoot on stairs, dropjumps and counter-movement jumps). Each patient was informed and instructed by a physiotherapist.

Individually fitted, semirigid insoles with bowl-shaped heels, moulded, longitudinal arch support and detorsion wedge were provided on the basis of a dynamic plantar pressure distribution measurement (EMED SF; Novel, Munich, Germany). Pedography and fitting were always carried out by the same technician. Insoles had to be worn for all physical activities during the treatment phase.

Subjects in C continued their activities without limitations, and no intervention was applied.

Quantities and methods applied

The PDI and PES were used to sum up patients' problems. PDI describes pain during various activities of daily life (family, rest, social activities, sexual activities, independence and lifemaintaining activities) based on visual analogue scales. The sum of pain in all categories was calculated.²⁶ PES was used to quantify the current pain before and after exercises in M₁ and M₂ (fig 2). PES describes the sensory and affective pain qualities, which were recorded using a simple descriptive scale. A summary score of each sensory and affective quality was calculated. Finally, data were t transformed in relation to a reference group with chronic musculoskeletal pain.⁹

Strength performance capacity (ankle plantar flexors) was measured during maximum stress (50° range of motion; concentrically and eccentrically, Lido-Active, Loredan); peak torque (mean of the three highest torques within five repetitions) was calculated.

Statistics

Data were stored in duplicate and compared for plausibility and correctness. After deblinding, data were assessed descriptively (means, SD, 95% CI; Statistical Discovery Software package JMP V.5.0.1.a, SAS Institute). Effects of treatment were analysed by either taking the difference between M_2 and M_1 or calculating the relative change in M_2 compared with M_1 ($(M_2-M_1)\times100\%$). In addition, the number of patients who responded to treatment was calculated.



Figure 2 Study protocol: baseline examination (U₁), 2-week instructional phase (W_1 , W_2), measurement 1 (M_1) and measurement 2 (M_2) separated by a 4-week intervention phase (W_3 - W_6). PDI, Pain Disability Index; PES, Pain Experience Scale. Presentation of methods used and study schedule.

The main quantity for statistical analysis of the effects of therapy on subjective pain was defined as the reduction of the PDI sum score (PDI_{sum}) between M₁ and M₂. Analysis of the equality of groups was performed according to Brown–Forsythe. PDI_{sum} was analysed statistically using a two-factor analysis of variance ($\alpha = 0.05$) for repeated measures with respect to the factors "group" (C, I, P) and "exercise" (M₁, M₂).

RESULTS

Subjective pain

PDI reduction was statistically significant after intervention in I and P (p<0.05, fig 3). The statistical model applied explained 93% of the variances; a significant interaction effect (group× exercise; p<0.05) was obvious.

Regarding individual responses, 21 of 28 patients (9/9 (100%) patients in I, 8/11 (73%) patients in P, and 4/8 (50%) patients in C) reported reduction in pain of >25% compared with baseline level. In all, 14 of 28 patients (8/9 (89%) patients in I, 6/11 (55%) patients in P, and 0/8 (0%) patients in C) reported reduction in pain of >50% compared with baseline level.



Figure 3 Differences in subjective pain (Pain Disability Index (PDI)) after therapy in relation to baseline ($M_2/M_1 \times 100$). Presentation of group means with 95% Cl. C, control group; I, group wearing insoles; M_1 and M_2 measurements 1 and 2; P, group undergoing physiotherapy.

Sensory and affective pain (PES) increased in all groups during M_1 up to a maximum pain following strength stress (fig 4). After treatment (M_2), pain increased only in C (fig 4).

Peak torque

After treatment, no increase in concentric peak torques (mean >10%) could be observed in any group, whereas higher eccentric peak torques (mean >10%) were obvious in I and P (fig 5).

DISCUSSION

Present randomised controlled trial results show pain reduction at rest and during exercise, following a short-term treatment of, 4 weeks of physiotherapy and fitting with insoles.

Patients with AT might benefit from non-surgical treatment.^{3 15} However, about 25–30% of patients were found to be resistant to treatment and had to undergo surgical intervention.^{3 16 28} In an 8-year follow-up, Paavola *et al*²⁸ reported reduction in pain and full functional recovery in 55 of 83



Figure 4 Results of the Pain Experience Scale (PES; t values, mean) pre-exercise, after treadmill run and after strength exercise during measurements 1 and 2 (M₁ and M₂).



Figure 5 Strength performance capacity of plantar flexion after treatment (M₂) in relation to baseline (M₁): isokinetic-eccentric peak torque (group means with 95% CI). C, control group; I, group wearing insoles; P, group undergoing physiotherapy.

patients with subacute or subchronic AT. Nevertheless, the effectiveness of measures applied (modified rest, muscle stretching, local injections, oral non-steroidal anti-inflammatory drugs, orthosis, physical therapy) could not be differentiated, since treatment was performed individually and was not standardised.²⁸

The recent reduction in pain after only 4 weeks of standardised physiotherapy shows that even short-term non-surgical treatment is effective in most patients. Possible mechanisms are nevertheless open to debate. Deep-friction massages can be understood as soft-tissue mobilisation, mechanotherapy (release of adhesions), fibroblast proliferation and stimulation of collagen synthesis.^{29 30} Additional sensory motor and eccentric training seem to be effective as local load management (eg, increase in collagen synthesis) and possibly because of modulation of afferent input to influence neuromuscular regulation.31 Optimisation of dynamic joint stability and postural control is rational, even though the spinal and central neurophysiological mechanisms of training effects remain largely unclear.¹⁷ Furthermore, increased eccentric peak torques indicate treatment-specific sensory motor effects. Modulation of nociception and a reduction in spinal inhibitions might be one explanation.

Training regimens with primarily eccentric stress have been used to treat AT.¹⁸⁻²¹ Alfredson et al¹⁹ prescribed heavy-load eccentric exercises for 12 weeks in 15 patients resistant to treatment. After treatment, all patients resumed their usual running mileage, with considerably reduced pain. Further studies confirmed these results.^{18 32} As a mechanism for reduction in pain, it is postulated that substance P-positive nerve endings,³³ which have grown during neovascularisation, are destroyed mechanically. However, the histological proof is still missing. Using colour Doppler sonography, a reduction in neovascularisation-and thereby a reduction in nerve endings-homogenisation of tendon structures and a decrease in tendon thickness could be demonstrated.^{32 34} By contrast, Khan et al³⁵ pointed out that the 12-month clinical outcome cannot be predicted using ultrasound. Power and colour Doppler sonography did not improve performance.³⁵ Taking the recent study design into account, it remains a matter of debate whether these mechanisms are already available after 4 weeks and whether a persistent, long-term effect could be attained.

Several studies confirm possible reduction in pain using insoles. However, differentiated, pathology-related ascriptions of valid mechanisms have not yet been successful. To date, the dominant opinion of mechanical aligning of the skeleton is

What is already known on this topic

- Achilles tendinopathy is common in athletes.
- Owing to individual responses and the long duration of problems, non-surgical treatment is time consuming (weeks to months).
- Effectiveness of single measures is not clarified.

What this study adds

- Physiotherapy (deep-friction massages, ultrasound, sensory motor and eccentric training) for 4 weeks and individually fitted insoles (moulded, longitudinal arch support, bowl-shaped heel) are effective in runners with Achilles tendinopathy.
- Maintaining running mileage and reduction in pain during activities of daily life as well as during exercise and higher eccentric plantar flexion strength capacity can be observed.

being increasingly criticised.²⁴ ²⁵ The effect varies individually, and, on average, is unspecific on segmental movements.²² Consequently, recent discussions focus on optimisation of muscular-regulated joint stability.²⁵ In this sense, a combination of longitudinal arch support with rear foot stabilisation possibly led directly (compression of the peroneal tendon) or indirectly (stimulation of the proprioceptors by altering joint position) to modulation of the afferent input. However, it must be remembered that within the recent study design it could not be clarified whether these effects are ascribable to reduction in pain itself, or whether an altered movement regulation (and thus load management) leads secondarily to the reduction of problems.

CONCLUSION

A short-term treatment of only 4 weeks of deep friction and ultrasound in combination with sensory motor and eccentric training, as well as insoles with longitudinal arch support might result in reduction in pain in most patients with AT, even without reducing running mileage. As most of the patients had little time to spare, these non-invasive short-term measures could possibly be considered valid treatment options in AT.

ACKNOWLEDGEMENTS

This study was supported by the Fortune-Science Grant Program, University of Tübingen, Germany, IETEC, Fulda, Germany, and Schuhstudio Türk, Freudenstadt, Germany.

We thank our former workgroup at the Medical Clinic, Department of Sports Medicine, University of Tübingen, Germany, for help with measurements.

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Competing interests: None.

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