CONCISE REPORT

Achilles tendinosis is associated with sprouting of substance P positive nerve fibres

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Patients and Methods

Patients

Ten Achilles tendon samples obtained from patients with Achilles tendinosis and 10 samples from patients with spontaneously ruptured Achilles tendons without a previous history of pain were submitted to the department of pathology between November 2000 and September 2003 for routine histological examination. The diagnostic criteria for Achilles tendinosis were tendon pain for several months and swelling of the tendon in the distal portion. Only spontaneously ruptured tendons which were resected within 48 hours after injury were included in the study. Table 1 shows the characteristics of the patients.

Immunohistochemistry of inflammatory cells and iron staining

Formalin fixed tissue was dehydrated and embedded in paraffin. Paraffin sections 3–5 μm thick were mounted on “Superfrost Plus” slides, heated for 20 minutes at 72°C, deparaffinised, and rehydrated. For antigen retrieval, sections were placed in a microwave for 30 minutes at 240 W in a citrate buffer at pH 7.3, then cooled to room temperature. After rinsing the slides, endogenous peroxidase activity was blocked by methanolic peroxide, the slides were rinsed again, and primary monoclonal antibody against CD3 for the detection of T lymphocytes, CD20 for B lymphocytes, or CD68 for macrophages (DAKO, Hamburg, Germany) in a dilution of 1:100 was applied. All slides were incubated using a Ventana machine and each antibody incubation was performed at 37°C; a labelled streptavidin-biotin-peroxidase method at 37°C was used to visualise positive reaction (Ventana Medical Systems basic DAB detection kit, Ventana Medical Systems Inc, USA). To detect haemosiderophages, paraffin sections were stained by Turnbull’s acid ferrocyanide reaction. Granulocytes were counted in haematoxylin and eosin stained sections prepared according to standard protocols.

The numbers of granulocytes, iron positive cells, CD3, CD20, and CD68 positive cells were evaluated in 10 randomly selected high power fields of view (×400) and expressed per square millimetre.

Immunohistochemistry of nerve fibres

The determination of the substance P (SP) positive sensory nerve fibres and sympathetic tyrosine hydroxylase (TH) positive nerve fibres has been described previously.1 Deparaffinised and rehydrated sections were blocked and incubated overnight with primary antibodies against TH or SP (Chemicon, Temecula, CA, USA). Immunofluorescent staining was achieved using Alexa 546 conjugated secondary antibodies against mouse or rabbit IgG (Molecular Probes, Leiden, The Netherlands). The numbers of TH and SP positive nerve fibres were evaluated in 17 randomly selected high power fields of view (×400) and expressed per square millimetre.

Abbreviations: SP, substance P; TH, tyrosine hydroxylase

Achilles tendinosis, also known as achillodynia, is viewed as a degenerative alteration of the Achilles tendon accompanied by pain and often associated with tendon thickening. It is common in athletes but occurs in non-athletes as well.1 Surgical specimens obtained from affected tendons show a range of degenerative changes such as changes in tendon fibre structure and arrangement as well as an increase in glycosaminoglycans, which may explain the swelling of the tendon.2 The pathophysiology of Achilles tendon pain is still unclear. Nociceptive nerve fibres can increase during inflammatory conditions—for example, in tendon pain is still unclear. Nociceptive nerve fibres can non-athletes as well.1 Surgical specimens obtained from Achilles tendinosis and 10 samples from patients with Achilles tendinosis and 10 samples from patients with Achilles tendinosis were obtained from the department of pathology between November 2000 and September 2003. The diagnostic criteria for Achilles tendinosis were tendon pain for several months and swelling of the tendon in the distal portion. Only spontaneously ruptured tendons which were resected within 48 hours after injury were included in the study. Table 1 shows the characteristics of the patients.

Objective: To identify and characterise nerve fibres and inflammatory alterations in painful Achilles tendinosis and thus gain evidence about the origin of pain in Achilles tendinosis.

Methods: The composition of 10 tendon samples from patients with a prior history of painful Achilles tendinosis and 10 samples from patients with spontaneously ruptured tendons but no previous pain was compared by immunohistochemistry and conventional histology.

Results: The presence of granulation tissue was shown in 8/10 cases of Achilles tendinosis. Nociceptive substance P (SP) positive nerve fibres were significantly increased, and an inflammatory infiltration comprising B and T lymphocytes was found. Additionally, small foci with iron positive haemosiderophages, indicating prior microtraumatic events, were found in 6/10 samples. None of the spontaneously ruptured tendons contained granulation tissue or haemosiderophages. Inflammatory infiltration in these patients consisted almost exclusively of granulocytes and SP positive nerve fibres were decreased. The density of sympathetic nerve fibres did not differ in the two conditions.

Conclusion: Achilles tendinosis is associated with the presence of granulation tissue, haemosiderophages, and SP positive nerve fibres, which may transmit the clinically pertinent pain. Achilles tendinosis may be caused by repeated microtraumata with ensuing organisation that is accompanied by sprouting of nociceptive SP positive nerve fibres.
Statistical analysis
The individual values of both groups were compared by Mann-Whitney test (SPSS V.11.0, SPSS Inc, Chicago, USA). Correlations were analysed using Spearman’s rank correlation analysis (SPSS). A p value <0.05 was the significance level.

Table 1 Characteristics of the patients investigated in this study

<table>
<thead>
<tr>
<th>Variable</th>
<th>Tendon rupture</th>
<th>Tendinosis</th>
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<tbody>
<tr>
<td>Patients (M/F)</td>
<td>8/2</td>
<td>7/3</td>
</tr>
<tr>
<td>Age (years), mean (SD)</td>
<td>48.0 (13.8)</td>
<td>47.3 (13.0)</td>
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<tr>
<td>Duration of pain before operation (months)</td>
<td>0</td>
<td>6–120*</td>
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<tr>
<td>Patients who received paratendinous corticosteroid injections (n)</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Patients with additional pathological conditions (n)</td>
<td>1 t/C1</td>
<td>2 t/C</td>
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*Three patients reported pain for more than 12 months; tthe patient had hypertension; C192 the second patient had calcium pyrophosphate dihydrate deposition disease (there were no calcium pyrophosphate dihydrate deposits visible in the tendon on microscopic examination).

Figure 1 Determination of cell types in painful Achilles tendinosis and ruptured Achilles tendon. The numbers of (A) CD3+ and (B) CD20+ lymphocytes and (C) CD68+ macrophages were determined by immunohistochemistry in 10 tendons from patients with Achilles tendinosis (AchDyn) and 10 tendons from patients with spontaneously ruptured tendons. The number of all cells was averaged from 10 high power fields and expressed as the number of cells/mm². AchDyn, painful Achilles tendinosis.

Figure 2 Comparison of sympathetic and nociceptive nerve fibres in painful tendinosis and ruptured tendon. The numbers of TH positive nerve fibres (A) and SP positive nerve fibres (B) were determined by immunofluorescent histochemistry in 10 tendons from patients with Achilles tendinosis (AchDyn) and 10 tendons from patients with spontaneously ruptured tendons. The number of nerve fibres was averaged from 17 high power fields and expressed as the number of fibres/mm². Box plots give the 10th, 25th, 75th, and 90th centiles. The median is given as a horizontal line within the box. SP, substance P; TH, tyrosine hydroxylase (sympathetic nerve fibres).
RESULTS
Ten Achilles tendon samples obtained from patients with Achilles tendinosis and 10 samples from patients with spontaneously ruptured Achilles tendons without a previous history of pain were studied to characterise the innervation, inflammatory infiltration, presence of granulation tissue, and haemosiderophages. We found that the number of B lymphocytes, T lymphocytes, and macrophages was significantly higher in Achilles tendinosis samples (fig 1A-C), whereas the number of granulocytes tended to be higher in tissue from ruptured tendons (fig 1D). Furthermore, in tendon samples from patients with Achilles tendinosis granulation tissue frequently containing haemosiderophages was found in 8/10 specimens (figs 1E and 3). Granulation tissue consisted of clusters of capillaries embedded in a fibroblast-rich stroma infiltrated by macrophages, B lymphocytes, and T lymphocytes. None of the samples from spontaneously ruptured tendons contained granulation tissue or haemosiderophages (figs 1E, and 3). The mean values for TH positive nerve fibres/mm² for the Achilles tendinosis group and for ruptured tendons were 1.00 and 0.18, respectively.

In a correlation analysis in patients with tendinosis, T cells correlated positively with B cells (Spearman rank correlation coefficient \( r_s = 0.683, p = 0.020 \)) and haemosiderophages \( r_s = 0.710, p = 0.014 \).

DISCUSSION
The origin of Achilles tendinosis has not been elucidated so far. Recently, neovascularisation in the affected tendon section correlating with pain was demonstrated by ultrasonography and magnetic resonance imaging. \( ^8 \) The finding was confirmed by immunohistochemistry, disclosing neovascularisation accompanied by nerve structures, which were not further characterised. \( ^8 \)

We compared the composition of 10 tendon samples from patients with a previous history of achillodynia with 10 samples from patients with spontaneously ruptured tendons without a previous history of pain by immunohistochemistry and conventional histology. Granulation tissue could be demonstrated in 8/10 cases of Achilles tendinosis. The granulation tissue comprised capillary vessels accompanied by an inflammatory infiltrate consisting of macrophages, B lymphocytes, and T lymphocytes. None of the spontaneously ruptured tendons contained granulation tissue or significant amounts of macrophages, B lymphocytes, and T lymphocytes. Instead they were infiltrated by large numbers of granulocytes as expected in the case of an acute traumatic event.

The number of nociceptive SP positive nerve fibres was found to be significantly higher in Achilles tendinosis samples than in tendon samples obtained from spontaneously ruptured tendons, whereas the number of TH positive nerve fibres did not differ. Spontaneously ruptured tendons were used as control group because immunohistochemistry for SP and TH in normal tendons taken from corpses did not deliver reproducible data. There are no published reports on the number of sensory nerve fibres of healthy human Achilles tendon. Both ruptured and Achilles tendinosis tendons contained low numbers of sensory nerve fibres compared with synovial tissue from healthy subjects.
The nerve fibres found in Achilles tendinosis specimens were often located next to small vessels in granulation tissue. Because we often found haemosiderophages within the granulation tissue, we suggest that microtraumatic events might have induced the formation of granulation tissue within the tendon. The degenerative alterations described in Achilles tendinosis may favour the occurrence of microtrauma. The growth of granulation tissue is accompanied by sprouting of sensory nerve fibres as demonstrated in animal models for wound healing. Interestingly, no similar increase of sympathetic nerve fibres was found in our Achilles tendinosis specimens.

The pattern of organisation of experimentally ruptured rat Achilles tendons was studied by Ackermann et al. They found that uncompromised Achilles tendons from rats are devoid of nerve fibres, but after rupture of the tendon, extensive sprouting of nerve fibres into the tendon takes place during the healing phase. SP positive nerve fibres were observed in the emerging granulation tissue next to the rupture site 1 week after rupture and reached a maximum density at week 4 after rupture. In correlation, 1–4 weeks after rupture the nociception to thermal and mechanical stimuli increased. Because we obtained tendon samples very early after tendon rupture, no such sprouting phenomenon was seen in the ruptured Achilles tendons. With the completion of the regenerative process, SP positive nerve fibres withdrew from the tendon tissue 8 weeks after rupture. Assuming a similar time course of organisation, the simultaneous occurrence of granulation tissue, nerve fibres, and haemosiderophages accompanied by longstanding local pain in Achilles tendinosis indicates that repeated microtrauma might have occurred which maintain the presence of granulation tissue and SP positive nerve fibres.

Besides transmitting nociceptive signals, the sprouting sensory nerve fibres may have profound effects on tendon structure because it has been shown that SP stimulates the proliferation of fibroblasts and production of transforming growth factor β by fibroblasts, potentially contributing to the thickening of the tendon seen in Achilles tendinosis.

To summarise, we found granulation tissue, haemosiderophages and, SP positive nerve fibres in tendons affected by Achilles tendinosis. The nociceptive nerve fibres may transmit the clinically pertinent pain. We suggest that Achilles tendinosis may be caused by repeated microtrauma with the ensuing organisation that is accompanied by sprouting of nociceptive SP positive nerve fibres.

ACKNOWLEDGEMENTS

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