Immunohistochemical Analysis of Sensory Nerve Endings in Ankle Ligaments: A Cadaver Study

Susanne Rein\textsuperscript{a}, Elisabet Hagert\textsuperscript{d}, Uwe Hanisch\textsuperscript{c}, Sophie Lwowski\textsuperscript{a}, Armin Fieguth\textsuperscript{b}, Hans Zwipp\textsuperscript{a}

\textsuperscript{a}Department of Trauma and Reconstructive Surgery, University Hospital ‘Carl Gustav Carus’, Dresden, 
\textsuperscript{b}Institute of Pathology, ‘Carl Thiem’ Hospital, Cottbus, and \textsuperscript{c}Institute of Legal Medicine, University Hospital of Hannover, Hannover, Germany; \textsuperscript{d}Department of Clinical Science and Education, Hand and Foot Surgery Center, Karolinska Institutet, Stockholm, Sweden

Key Words
Ankle · Immunohistochemistry · Ligaments · Mechanoreceptors · Proprioception

Abstract

\textbf{Background:} The aim of this study was to analyze the pattern and types of sensory nerve endings in ankle ligaments using immunohistochemical techniques, in order to gain more insight into functional ankle stability. \textbf{Methods:} One hundred forty ligaments from 10 cadaver feet were included: the calcaneofibular and anterior/posterior talofibular ligaments from the lateral complex; inferior extensor retinaculum complex, talocalcaneal oblique and canalis tarsi ligaments from the sinus tarsi; deltoid ligament with its individual portions from the medial complex, and anterior tibiofibular ligament (ATiFL) from the syndesmosis. Mechanoreceptors were classified according to Freeman and Wyke [Acta Anat (Basel) 1967; 68:321–333] after staining with hematoxylin-eosin, low-afﬁnity neurotrophin receptor p75, protein gene product 9.5, and S-100 protein. \textbf{Results:} Free nerve endings were the predominant sensory endings in all four complexes, with the greatest density in the lateral and medial complexes; followed by Ruffini endings, unclassifiable corpuscles, Pacini corpuscles, and Golgi-like endings. Ruffini endings were significantly more prevalent in the ATiFL than in the medial complex, and more common than Pacini corpuscles and Golgi-like endings in the lateral, medial, and sinus tarsi complexes. A greater number of blood vessels correlated with a greater number of free nerve endings. There was a negative correlation between the number of Ruffini endings, unclassifiable corpuscles, and age. \textbf{Conclusions:} Free nerve endings are the dominant mechanoreceptor type in the ankle ligaments, followed by Ruffini endings. The ligaments of the lateral and medial ankle complexes are more innervated than the sinus tarsi ligaments.

Introduction

Joint stability relies upon fine interactions of both static and dynamic elements. While static joint stability is constituted by the anatomical congruity of joint surfaces and the ligamentous restraints acting to limit joint translations, the dynamic aspects of joint stability chiefly concern proprioceptive control of the compressive and directional muscular forces acting on a joint [Frank, 2004].
Ligaments are important structures with a role not only in the static stability of joints but also in the sensorimotor control of joint movements. As such, ligaments are recognized as sensory organs, capable of monitoring and supplying relevant kinesthetic and proprioceptive data [Solomonow, 2006].

Proprioception is a critical part of ankle stability. The stability of the ankle joint is provided by a lateral and medial complex of ligaments as well as the distal tibiofibular syndesmosis [Zwipp, 1994]. The subtalar joint plays a key role in adapting the foot to the ground [Stagni et al., 2003], which is stabilized by the sinus tarsi ligaments [Schmidt, 1978; Zwipp, 1994].

Several studies have investigated sensory nerve endings of ankle ligaments using the gold chloride technique [Michelson and Hutchins, 1995; Moraes et al., 2008]. However, gold chloride impregnates not only nerve tissue but also elastic fibers in blood vessels and reticular fibers, thus providing nonspecific imaging of neural elements in tissue [Soule, 1962; Koch et al., 1995; Gómez-Barrena et al., 1999].

Recently, immunohistochemical analysis precisely determined sensory nerve endings in wrist ligaments [Hagert et al., 2004, 2005, 2007]. This technique utilized specific neural and perineural markers: glial cell S-100 protein (S100), low-affinity p75 neurotrophic receptor (p75), and protein gene product 9.5 (PGP 9.5). The innervation of ligaments is characterized by specific nerve endings, so-called mechanoreceptors, which can be classified due to their typical shape and neurophysiological traits according to Freeman and Wyke [Freeman and Wyke, 1967a]. As different mechanoreceptors have been attributed different functions, a profound analysis of the distribution of sensory nerve endings in ankle-stabilizing ligaments will give more insight into functional ankle stability.

The aim of this study was to analyze the pattern and types of mechanoreceptors in the different anatomical complexes of ankle ligaments using designated immunohistochemical markers, as well as the overall vascularity of the ligaments.

Materials and Methods

Cadaver Specimens

All protocols in this study were approved by the local ethics committee review board. Ten feet from 5 subjects (2 women and 3 men) with a mean age of 57 ± 20 years (range 36–86 years) were included in this study. Five left and 5 right feet were analyzed. The cadavers were refrigerated (4 °C) pending ligament harvest, and the mean time between death and harvest was 3.6 ± 2.4 days (range 1–7 days). All feet were assessed macroscopically and showed no signs of ligament injury or structural abnormality.

Ligament Specimens

A lateral and medial semicircular skin incision was made over the ankle. The lateral and medial ligaments as well as the ligaments of the sinus tarsi were exposed.

Sinus tarsi ligaments were defined according to Schmidt [1978]. Dissection of the different portions of the deltoid ligament was based on the description of Pankovich and Shivaram [1979]. The other ligaments were defined according to the description of Zwipp [1994].

The anterior (ATFL) and posterior talofibular ligaments (PTFL) as well as the calcaneofibular ligament (CFL) were obtained from the lateral complex. The lateral (IERI), intermediate (IER), and medial (IERM) roots of the inferior extensor retinaculum (IER) and the talocalcaneal oblique ligament (TCOL) and the canalis tarsi ligament (CTL) were resected in the sinus tarsi. The tibiocalcaneal (TLN), tibiocalcaneal (TCL), and superficial tibiotalar (STTL) ligaments from the superficial layer, as well as the anterior (ATTL) and posterior tibiotalar (PTTL) portions from the

Abbreviations used in this paper

| Ligaments
| Lateral complex |
| ATFL | anterior talofibular ligament |
| CFL | calcaneofibular ligament |
| PTFL | posterior talofibular ligament |

| Medial complex |
| (a) Superficial layer |
| TNL | tibionaviculare ligament |
| TCL | tibiocalcaneal ligament |
| STTL | superficial tibiotalar ligament |
| (b) Deep layer |
| ATTL | anterior tibiotalar ligament |
| PTTL | posterior tibiotalar ligament |

Sinus tarsi |
| IER | inferior extensor retinaculum |
| IERI | inferior extensor retinaculum, intermediate root |
| IERL | inferior extensor retinaculum, lateral root |
| IERM | inferior extensor retinaculum, medial root |
| TCOL | talocalcaneal oblique ligament; synonym: cervical ligament |
| CTL | canalis tarsi ligament; synonym: interosseous talocalcaneal ligament |

Distal tibiofibular syndesmosis |
| ATFL | anterior tibiotalar ligament |

Further abbreviations |
| H&E | hematoxylin-eosin |
| HRP | horseradish peroxidase |
| IR | immunoreactivity |
| p75 | low-affinity nerve growth factor receptor p75 |
| PBS | phosphate-buffered saline |
| PGP 9.5 | protein gene product 9.5 |
| S100 | S-100 protein |
deep layer of the deltoid, were harvested from the medial complex. The anterior interosseous tibiofibular ligament (ATiFL) from the distal tibiofibular syndesmosis was also resected. All 140 ligaments were completely dissected from their insertion into bone.

**Immunohistochemistry**

Specimens were immediately fixed in 4% buffered formaldehyde solution (pH = 7.4) for 24 h at 4°C, decalcified with diaminonethane-tetraacetic acid (EDTA), and embedded in paraffin. Sections of 4 μm were cut and mounted onto silane-coated slides for conventional staining and immunohistochemistry. All ligaments were cut at 5 levels, with a 50-μm cutting interval between each level.

The mounted sections were dehydrated beginning with xylol in decreasing concentrations. Sections were then rehydrated with distilled water. Slides were incubated in a 1% H2O2 blocked endogenous peroxidase activity for 5 min at room temperature, and rinsed for 3 × 5 min in phosphate-buffered saline (PBS; pH = 7.4). Subsequently, the slides were treated with an ultra vision horse-radish peroxidase-polymer kit (HRP-polymer kit, code TL-060HL; Thermo Scientific, Schwerte, Germany) for 5 min at 37°C, followed by incubation with primary antibodies for 60 min at 37°C. After rinsing with PBS 3 × 5 min, the secondary antibody with enhancer of the ultra vision HRP-polymer kit was applied for 10 min at room temperature. The sections were washed in PBS 3 × 5 min again before the HRP-polymer kit was used for 15 min at room temperature. Afterwards the sections were rinsed in PBS 3 × 5 min once again and detected with chromogene 3-amino-9-ethylcarbazol (AEC; Romulin, code REEC810 L; Zytomed Systems; Berlin, Germany) for 15 min at room temperature.

Thereafter followed rinsing in distilled water, and counterstaining with hematoxylin was performed. Finally, sections were dehydrated and covered with Entellan (Merck, Darmstadt, Germany). As a negative control, identical staining without addition of primary antibodies was performed in parallel followed by counterstaining with hematoxylin. When staining for PGP 9.5 was performed, the slides were initially pretreated with 1% Triton X-100 solution for 20 min at room temperature and then rinsed in PBS 3 × 5 min before treatment with 1% H2O2, as described above.

**Antibodies**

Polyclonal rabbit antisera against p75 (working dilution: 1:200; code N-3908; Sigma, Saint Louis, Mo., USA) was used. The antibody specifically reacts with the low-affinity neurotrophic receptor, which binds nerve growth factor and all other neurotrophins, including brain-derived neurotrophic factor, neurotrophin-3, and neurotrophin-4.

Polyclonal rabbit antisera against PGP 9.5 (working dilution 1:500; code: 7863-0504; AbD Serotec, Düsseldorf, Germany) and polyclonal antisera against S100 (working dilution 1:500; code Z 0311; DakoCytomation, Glostrup, Denmark) were also used. The antibody against PGP 9.5 is a panneuronal marker reacting with PGP 9.5 in all mammalian species tested, including humans. The antibody against S100 specifically stains the S100 protein, including Schwann cells of the peripheral nervous system.

The monoclonal mouse anti-human smooth muscle–actin 1A4 antibody (sm-actin; working dilution 1:750; code M 0851; DakoCytomation) labels smooth muscle cells in blood vessels, whereas epithelia, lymphocytes, cardiac and skeletal muscle cells, endothelial cells, fat cells, Schwann cells, and fibroblasts are negative.

**Morphological Analysis and Cell Counting**

Histological examination of the stained tissue sections was performed using a Leica light microscope (Leitz DMRBE, Wetzlar, Germany) with a Leica camera (Leica DC 300; Leica Microsystems CMS GmbH, Heerbrugg, Switzerland).

Hematoxylin-eosin (H&E)-stained slices were used for determination of tissue morphology, and all ligaments were evaluated to exclude signs of ligament lesions before starting the mechanoreceptor analysis, because ligamentous lesions lead to a gradually decreased number of sensory nerve endings over time [Denti et al., 1994]. In order to exclude acute lesions, histological analysis was focused on determining possible necrosis, bleeding, fibrin exudation, and granulation tissue. For exclusion of old lesions, bleeding residuals, focal intraligamentous vascularization inside the connective collagen tissue, fibroblast proliferation, hyalinization of collagen fibroblasts, scar tissue, chondroid metaplasia outside the insertion, calcification, and ossification were assessed.

Mechanoreceptors were analyzed according to the classification of Freeman and Wyke, modified by Hagert [Freeman and Wyke, 1967b; Hagert, 2008]. Ruffini, Pacini, Golgi-like, and free nerve endings as well as unclassifiable corpuscles were counted in the S100, p75, and PGP 9.5 stainings in all 5 levels with respect to the total cell count per section at an original magnification of 400× (high-power field). A standard 10 × 10 grid was used for determination of mechanoreceptor size. Blood vessels were counted at two representative levels in the sm-actin stain, identified by specific immunoreactivity (IR) of sm-actin of the smooth muscle cells in the wall of the vessels. All specimens were blinded for cell counts.

Sensory corpuscles, which could neither clearly be defined as Ruffini, Pacini, or Golgi-like corpuscles nor as nerve fascicles, were deemed unclassifiable according to Hagert et al. [2005; Hagert, 2008, 2010].

**Statistical Analysis**

Means with standard deviations were used for descriptive statistics throughout the article except for figure 4 and table 1, which are reported as absolute values.

The first purpose was to investigate the general distribution of mechanoreceptors and blood vessel in the ligaments. Contingency tables with Fisher’s exact test were used for this analysis. The level of significance was considered high with p < 0.05.

The second purpose was to analyze the general distribution of sensory nerve endings in all ligaments (n = 140), which was performed with the Friedman test followed by the Wilcoxon test with post hoc Bonferroni adjustments. The five different types of mechanoreceptors allowed ten possible test comparisons, which resulted in a significance level of p < 0.005.

The third purpose was to analyze the mechanoreceptors between the anatomical complexes. The ATiFL as a part of the distal tibiofibular syndesmosis, and the lateral, medial, and sinus tarsi complexes were compared. The Kolmogorov-Smirnov test was performed to investigate data distribution. Since all groups were found to not have normal distributions, the subsequent statistical analysis was performed using the Kruskal-Wallis test followed by the Mann-Whitney test with post hoc Bonferroni adjustments, with a final level of significance of p ≤ 0.0083, due to six tests of significance between the four groups at the Bonferroni adjustment.
The fourth purpose was to compare the quantity of the different types of mechanoreceptors within each anatomical complex. The Friedman test, followed by the Wilcoxon test with post hoc Bonferroni adjustments, was performed. The final level of significance was $p^0.005$, since ten tests of significance between the five mechanoreceptor groups were performed at the Bonferroni adjustment.

The fifth purpose was to test correlations with regard to age, sensory nerve endings, and blood vessel numbers. Correlation analysis was performed with Spearman's rho coefficient with a two-sided significance level of $p^0.05$. All 140 ligaments were examined together for the correlation analysis.

**Results**

Acute and chronic lesions were excluded in all ligaments following analysis using H&E (table 1).

**Ruffini Endings (Type I)**

The central axon was clearly demarcated via staining for PGP 9.5 and S100, whereas it showed no specific IR in staining for p75 (fig. 1). The thin and incomplete capsule was IR for p75. Occasionally, identification of smaller corpuscles located within the capsule of corpuscles was also noted. The dendritic nerve terminals within the corpuscle showed IR for S100, p75, and PGP 9.5. Sometimes Ruffini endings were grouped as two endings together. The size of a single Ruffini ending varied between 50 and 120 μm. One hundred eight of the 140 ligaments (77%) contained Ruffini endings.

**Pacini Corpuscles (Type II)**

The characteristic lamellar and thick capsule of the Pacini corpuscles displayed intense p75 IR. There was a strong intracorpuscular reaction after processing for S100. Nerve fibers of the afferent nerve fascicle, the so-called parent axon, as well as thin nerve fibers within the corpuscles, were demarcated after staining for PGP 9.5 (fig. 2). Pacini corpuscles were generally smaller than Ruffini endings. They were observed in 49 of 140 ligaments (35%).

**Golgi-Like Endings (Type III)**

Golgi-like endings were rarely found. They were only present in 15 ligaments (11%), with a total of 28 Golgi-like endings in all 140 investigated ligaments.

They had a large fusiform body with a thin capsule and smaller grouped corpuscles seen within the large body. The smaller corpuscles contained terminal nerve endings with S100, p75, and PGP 9.5 IR. Thick parent axons were seen to enter the corpuscle at one side (fig. 3). The longitudinal axis of the Golgi-like endings was arranged in parallel to the ligamentous collagen bundle composition. Golgi-like endings were generally remarkable due to their size, which considerably varied between 150 and 800 μm for the true corpuscle.

**Table 1. Exclusion of posttraumatic ligamentous lesions**

<table>
<thead>
<tr>
<th>Histological feature</th>
<th>ATiFL (n = 10)</th>
<th>Lateral complex (n = 30)</th>
<th>Medial complex (n = 50)</th>
<th>Sinus tarsi (n = 50)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Autolysis</td>
<td>yes</td>
<td>yes</td>
<td>yes</td>
<td>yes</td>
</tr>
<tr>
<td>Signs of acute injury</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Bleeding residuals</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Focal intraligamentous vascularization (inside connective collagen tissue)</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Fibroblast proliferation</td>
<td>–</td>
<td>–</td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td>Hyalinization of collagen fibers</td>
<td>–</td>
<td>2</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>Scar tissue</td>
<td>1</td>
<td>2</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>Chondroid metaplasia outside ligament insertion area</td>
<td>1</td>
<td>2</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>Calcification</td>
<td>1</td>
<td>2</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>Ossification</td>
<td>1</td>
<td>2</td>
<td>4</td>
<td>3</td>
</tr>
</tbody>
</table>

Results of the tissue morphology analysis in H&E staining are presented. In terms of histological signs of old ligamentous lesions, only few focal and small herd (not larger than one high-resolution visual field) reparative or scar tissue changes were observed, which can be interpreted as an expression of degeneration or state after microtrauma. There, the general/overall ligamentous structure was intact. No morphological correlate for an old complete or partial ligament rupture was seen.
Free Nerve Endings (Type IV)

Nerve fascicles were generally observed in the vicinity of vessels and in the connective tissue sheaths within the ligaments as well as in the epiligament. There was marked p75 and S100 IR in the nerve fascicle, whereas the axons within the nerve fascicles were stained for PGP 9.5. The size of the free nerve endings varied depending on the degree of myelination, the number of axons, and the cutting level. All investigated ligaments (100%) contained free nerve endings.
Unclassifiable Corpuscles (Type V)

Certain sensory corpuscles could neither be clearly defined as Ruffini, Pacini, Golgi-like nor as nerve fascicles and were therefore deemed unclassifiable. A variation of size and appearance has been observed. An incomplete capsule was often seen after staining for p75. Unclassifiable corpuscles were documented in 95 of 140 ligaments (68%).

General Distribution (n = 140)

Sensory nerve endings and blood vessels were equally distributed and significantly more common in the epiligamentous regions compared to the interstitial area \((p < 0.0001, \text{ respectively})\) (fig. 4). Furthermore, sensory nerve endings were significantly more common close to the insertion into bone rather than the central, or midportion, of a ligament \((p = 0.001)\) or equal distribution. Interestingly, blood vessels were significantly more often interstitial \((p = 0.043)\) and at the insertion into the bone \((p < 0.0001)\) than equally dispersed in the ligament (fig. 4).

The free nerve endings were the predominant sensory ending, followed by Ruffini endings, unclassifiable corpuscles, Pacini corpuscles, and ultimately Golgi-like endings (fig. 5). Statistical analysis showed significant differences between all sensory nerve endings \((p < 0.0001, \text{ respectively})\), except between Ruffini endings and unclassifiable corpuscles. The calculated relationship between the sensory nerve endings is shown in table 2.

No Ruffini, Pacini, or Golgi-like corpuscles were found in 24 of 140 ligaments (17%). Only free nerve endings were observed in 12 ligaments (8.5%) (lateral complex: 1 ATFL and 3 CFL; medial complex: 1 TNL, 1 STTL, 1 ATTL, and 1 PTTL; sinus tarsi: 1 IERL, 2 IERM, and 1 CTL). Only free nerve endings and unclassifiable corpuscles were seen in 12 ligaments (8.5%) (lateral complex: 1 ATFL and 1 PTFL; medial complex: 3 TCL and 2 PTTL; sinus tarsi: 1 IERL, 1 IERM, 2 TCOL, and 1 CTL).

Distribution of Mechatoreceptors between the Anatomical Complexes

There were no discernible differences in the general morphology of corpuscles between the anatomical complexes. The ATiFL ligament had significantly more Ruffini corpuscles than the medial complex \((p = 0.005; \text{ fig. 6})\). Significantly more free nerve endings were counted in the lateral complex as compared to the sinus tarsi \((p < 0.0001; \text{ fig. 7})\). Also, significantly more blood vessels were found in the lateral complex as compared to the sinus tarsi \((p = 0.006)\) and the ATiFL \((p = 0.002; \text{ fig. 7})\). Furthermore, the medial complex had significantly more free nerve endings \((p < 0.0001)\) and blood vessels \((p = 0.002)\) than the sinus tarsi (fig. 7).
Distribution of Mechanoreceptors within the
Anatomical Complexes

Significantly more Ruffini endings than Pacini corpuscles were observed in the lateral, medial, and sinus tarsi complexes (p < 0.0001, respectively) (fig. 6). Similarly, significantly more Ruffini endings than Golgi-like endings were found in the lateral, medial, and sinus tarsi complexes (p < 0.0001, respectively) (fig. 6). Significantly more unclassifiable corpuscles than Pacini corpuscles were observed in the lateral (p = 0.002), medial (p = 0.004), and sinus tarsi complexes (p < 0.0001) (fig. 6).

Significantly more unclassifiable corpuscles than Golgi-like endings were observed in the lateral, medial, and sinus tarsi complexes (p < 0.0001, respectively) (fig. 6).
Finally, significantly more free nerve endings than Ruffini, Pacini, Golgi-like, and unclassifiable corpuscles were seen in the lateral, medial, and sinus tarsi complexes (p < 0.0001, respectively). Significantly fewer Golgi-like endings (#) than Ruffini endings and unclassifiable corpuscles were observed in the lateral, medial, and sinus tarsi complexes (p < 0.0001, respectively). Significantly fewer Pacini (lateral: p = 0.002; medial: p = 0.004; sinus tarsi: p < 0.0001) and Golgi-like endings (p < 0.0001, respectively) than unclassifiable corpuscles (+) were seen in the lateral, medial, and sinus tarsi complexes.

**Correlation Analysis**

A greater number of blood vessels correlated significantly with a greater number of free nerve endings (p < 0.0001; r = 0.57; fig. 8). There was also a significant positive correlation between the number of Ruffini endings and the number of free nerve endings (p = 0.006; r = 0.23) as well as unclassifiable corpuscles (p < 0.0001; r = 0.43). In addition, there was a significant positive correlation between the number of Pacini corpuscles and Golgi-like endings (p = 0.05; r = 0.17).

There was a significant negative correlation between Ruffini endings (r = -0.33; fig. 9), as well as unclassifiable corpuscles (r = -0.44; fig. 10) and age, with a decrease in innervation in older specimens (p < 0.0001, respectively).

**Discussion**

**General Distribution**

In our study, sensory corpuscles and blood vessels were found mainly close to ligament insertions into bone as well as in the epiligamentous region of ligaments. This is in accordance with previous publications, which also describe the majority of mecanoreceptors at the insertion of ligaments in the shoulder [Morisawa, 1998], wrist [Hagert et al., 2007; Tomita et al. 2007], knee [Kennedy et al., 1982; Schultz et al., 1984; Schutte et al., 1987; Del Valle et al., 1998], and ankle joints [Moraes et al., 2008]. Similarly, 93% of the mecanoreceptors in cat lateral ankle ligaments have been found near the attachment to the fibula and calcaneus [Takebayashi et al., 1997]. Krauspe
**Fig. 8.** A higher number of blood vessels significantly correlated with a higher number of free nerve endings ($p < 0.0001; r = 0.57$).

**Fig. 9.** The number of Ruffini endings significantly decreased with older age ($p < 0.0001; r = -0.33$).

**Fig. 10.** The number of unclassifiable corpuscles significantly decreased with older age ($p < 0.0001; r = -0.44$).
et al. [1992] also reported that a majority of mechanoreceptors in the cat anterior cruciate ligament were found near its femoral attachment. In human knees mechanoreceptors are located near the surface of the ligament, equivalent to the epiligamentous region described above [Schultz et al., 1984]. This epiligamentous area has been proposed to be of importance for ligament nutrition and protection, as well as for providing support to the neurovascular bundles in the ligament [Chowdhury et al., 1991]. A polar distribution of mechanoreceptors allows them to act more sensitively as monitors of tension applied to the ligament as compared to those situated in the center of a ligament [Takebayashi et al., 1997]. Since ligaments are more resistant to strains close to their insertions, this serves to ensure triggering of sensory nerve endings only by potentially noxious motions, while they remain silent during ordinary joint activity [Solomonow, 2006].

Disruption of lateral ankle ligaments mostly occurs intraligamentously [Zwipp, 1986], indicating a preservation of vascularity and sensory nerve endings at the bony insertions which is known to have a positive effect on ligament healing [Ivie et al., 2002]. Immobilization of the ankle, both for conservative and for postoperative treatment, generally leads to healing of a disrupted ligament. If the ligament tensile strength is restored, sensory reeducation with proprioceptive exercises can be initiated to stimulate the remaining sensory nerve endings at the bony insertion and enhance proprioceptive function of the ankle ligaments [Freeman et al., 1965; Cooper and Farr, 1978; Bernier and Perrin, 1998]. The ligamentous insertion regions should thus be conserved during surgery to avoid iatrogenic lesions of important sensory nerve endings [De Avila et al., 1989].

Free nerve endings were the predominant mechanoreceptor type seen in the ankle ligaments, followed by Ruffini endings, unclassifiable corpuscles, Pacini corpuscles, and Golgi-like endings. In relation to the proprioception of the ankle joint it indicates an importance of reacting to noxious, chemical, mechanical, and inflammatory stimuli. The subsequent predominance of Ruffini endings serves to maintain control of joint position and kinesthesia. Joint acceleration/deceleration, as detected by Pacini corpuscles, appears to play only a minor role in ankle joint proprioception. Analogous results were found in recent immunohistological analysis of wrist ligaments, where Ruffini endings were seen more often than Pacini corpuscles [Hagert et al., 2004, 2005, 2007]. Only few Golgi-like corpuscles have been found in the whole analysis. This is in contrast with previous literature, where a predominance of Pacini corpuscles, followed by Golgi-like corpuscles, and lastly Ruffini corpuscles, were described in the deltoid ligament and/or lateral ankle ligaments using the gold chloride technique [Michelson and Hutchins, 1995; Moraes et al., 2008]. In addition, as Golgi-like endings detect extreme ranges of joint movement [Newton, 1982], they appear more often in ligaments of big joints, e.g. the cruciate ligament of the knee [Johansson et al., 1991], than in ligaments of small joints [Lin et al., 2006; Hagert, 2008; Chikenji et al., 2010].

The innate function and physiological properties of unclassifiable corpuscles is presently unknown. A pure morphological analysis of unclassifiable corpuscles can never deduce a physiological function – rather, a microneurographic investigation would be of importance to further our knowledge of this enigmatic sensory nerve ending. The high number of unclassifiable corpuscles seen in our study is an expression of the limitations of an applied descriptive analysis using 2D light-microscopy, which does not permit a complete representation of the whole mechanoreceptor. The primary difference between our study and those published previously is that we have used a combination of immunohistochemical markers to target specific mechanoreceptor structures, and we performed the total analysis on a large number of sections at five different levels in each ligament. We propose that our analysis offers a greater distinction of both receptor type and frequency based on this. However, immunofluorescence analysis offers a 3D analysis of sensory nerve endings and is frequently used by researchers [Hagert et al., 2012; Lee et al., 2012]. Nevertheless, even 3D reconstructions of multiple levels of ligaments have revealed a great portion of unclassifiable corpuscles in wrist ligaments [Tomita et al., 2007], which points to a possible limitation of the existing gold-standard classification of Freeman and Wyke, modified by Hagert [Freeman and Wyke, 1967a, b; Hagert, 2008].

Few ligaments have only had free nerve endings without Ruffini, Pacini, or Golgi-like corpuscles. However, no general pattern of particular ligaments of this could be observed, because ligaments from all anatomical complexes were involved.

**Anatomical Complexes**

In our study, Ruffini endings were the second most common receptor overall, and they were particularly frequent in the ATiFL. Studies on the biomechanical function of the ATiFL has shown that once the ATiFL is cut, the tibia and lateral malleolus can move separately [Taramoto et al., 2008]. The distal tibiofibular syndesmosis at-
taches the fibula dynamically at the tibia, which gives the fibula the possibility to perform a three-dimensional movement during physiological dorsalextension and plantarflexion of the ankle joint [Close, 1956; Reimann and Anderhuber, 1980; Peter et al., 1994; Beumer et al., 2003; Rammelt et al., 2008]. The function of the Ruffini ending in the ATiFL is thus to provide constant monitoring of the joint position and motion of this important articulation, and corresponds well with our frequent findings of Ruffini endings in this region.

The lateral and medial complexes are richly innervated by free nerve endings, whereas the sinus tarsi is less so. It seems, therefore, that the ligaments of the lateral and medial complexes as the primary stabilizers of the ankle joint have more capacity for the detection of proprioceptive information than the sinus tarsi ligaments.

In 1965, Freeman et al. [1965] were the first to introduce the term ‘functional instability’ as a patient’s subjective feeling that the ankle is giving way, meaning that a patient has the feeling of instability of the ankle, particularly when walking or running on uneven ground [Freeman et al., 1965]. It has been hypothesized that this proprioceptive deficit may result from a partial deafferentation caused by damage of sensory nerve endings of the joint capsule and ligaments, resulting in functional instability of the ankle [Freeman et al., 1965]. The latter could lead to loss of position sense [Glencross and Thornton, 1981], delayed peroneal reaction time [Konradsen and Ravn, 1991; Karlsson and Andreason, 1992], strength deficits, and impaired postural control [Tropp and Odenrick, 1988; Richie, 2001; Hertel, 2002; Arnold et al., 2009; Munn et al., 2010]. This innervation study shows that no-circence, mediated by free nerve endings, has enormous importance in ankle proprioception, followed by joint position sense due to a predominant occurrence of Ruffini endings.

**Correlation Analysis**

We found a positive correlation between the degree of vascularility and innervation, corresponding with previous studies on human ligaments [Chowdhury et al., 1991; Hagert et al., 2004, 2005, 2007]. Similarly, there was a positive correlation between Ruffini endings and free nerve endings or unclassifiable corpuscles, while the presence of Pacini corpuscles was positively correlated with that of Golgi-like endings. As discussed above, mechanoreceptors are mainly located in the epiligament and the ligamentous insertion into bone. The insertion region of a ligament is particularly exposed to a complex combination of both compressive and tensile forces [Spalazzi et al., 2006]. The presence of a multitude of receptor types in this region thus permits sensorimotor monitoring of all joint activities, where different mechanoreceptor types complement each other due to their different physiological properties, guaranteeing sensorimotor control of all complex joint movements. This ligament-bone interface thus has ideal anatomical and physiological properties for joint protective sensorimotor functions.

The presence of Ruffini endings had a negative correlation with age, with a decrease in Ruffini endings in older specimens. Although these results have a limitation in that they are based on a small sample size, the finding is interesting as it is in agreement with previous studies on animals where a general decrease in neurotrophins has been correlated with the reduction of sensory nerves that occurs in senescence [Ulfhake et al., 2002], and the total number of Ruffini endings in the rabbit anterior cruciate ligament has been found to significantly decrease with aging [Aydog et al., 2006]. These results also correlate with clinical observations on functional ankle stability, where older age has been associated with increased peroneal reaction time and impaired balance control [Rein et al., 2010]. A decrease in sensory nerve endings with older age may thus contribute to the lack of balance and coordination observed in elderly individuals, resulting in an increased risk of fall and injury [Runge, 2002; Kwan et al., 2011].

In conclusion, sensory nerve endings are mainly located closed to the ligament insertion into bone and in the epiligamementous region. Free nerve endings were the predominant receptor type, followed by Ruffini endings, indicating that nociception and joint position have enormous importance in ankle proprioception. The ligaments of the lateral and medial ankle complexes are more innervated than the sinus tarsi ligaments, which implies more capacity for detection of proprioceptive information in these ligaments. A decrease in sensory nerve endings with age has been observed.

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Sensory Nerve Endings in Ankle Ligaments

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