

Ultrasonographic features of the intrinsic foot muscles in patients with and without plantar fasciitis: a novel case-control research study

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Abstract

Introduction: The aim of the present study was to compare by ultrasound imaging (USI) the thickness and cross-sectional area (CSA) of the flexor hallucis brevis (FHB), flexor digitorum brevis (FDB), abductor hallucis brevis (AHB) and quadratus plantae (QP) muscles between individuals with and without plantar fasciitis (PF).

Material and methods: A case-control study was performed with 64 participants divided into two groups: A, PF group ($n = 32$) and B, healthy group ($n = 32$).

Results: USI measurements for FHB CSA ($p = 0.035$) decreased, showing statistically significant differences for the PF group, while the QP CSA ($p = 0.40$) increased, showing statistically significant differences for the PF group with respect to the healthy group. The rest of the intrinsic foot muscles (IFM) did not show statistically significant differences; however in FHB, FDB, QP and AHB thicknesses and FDB CSA showed a slightly decrease for the PF group.

Conclusions: USI measurements showed that the CSA of the FHB muscle is reduced in patients with PF while the CSA of the QP muscle is increased in patients with PF.

Key words: ultrasonography, musculoskeletal diseases, plantar fascia.

Introduction

Plantar fasciitis (PF) is the most common cause of heel pain [1]. This condition was described as a degenerative syndrome associated with pain, lack of functionality and stiffness of the plantar fascia [2]. In addition, the calcaneal insertion and mid-foot area were identified as the most common injured locations. Thus, it is frequent to find literature with the terms: neuritis, subcalcaneal bursitis, calcaneal periostitis or heel spur syndrome [3]. Lemont *et al.* reported that PF episodes could be associated with or without inflammation [4]. Several authors have argued that PF develops due to repeated trauma, therefore considering it

an overuse injury [5]. To date, this condition presents an estimated prevalence of 7% in the general population, with bilateral involvement reported in 30% of the affected patients.

The origin of this condition remains unclear despite intrinsic and extrinsic risk factors having been described, such as an excessive body mass index (BMI), reduced ankle dorsiflexion, pes planus, pes cavus, Achilles tendon disturbances, overpronation causing additional stress and weakness of the intrinsic foot muscles and plantar muscles [6] – intrinsic risk factors – and excessive physical activity, low-quality footwear, inadequate surfaces, and walking barefoot [6] – extrinsic factors. Patients with PF experienced severe pain at the first steps in the morning or after non-activity for long periods, such as sleeping or sitting. If these symptoms persist in time, this condition could progress to chronic PF, which leads to an important decrease in quality of life, negative social impact and limited daily activities [7].

McKeon *et al.* proposed a concept drawing a parallelism between the intrinsic foot muscles (IFM) and the trunk core muscles, indicating the importance of these small foot muscles for the biomechanics and foot structures (arches) stabilization [8]. The IFM and extrinsic foot muscles (EFM) work in a coordinated manner to provide movement and stability of the three flexible arch structures. In this line, the PF and the IFM – hallucis brevis (AHB), flexor hallucis brevis (FHB), flexor digitorum brevis (FDB) and quadratus plantae (QP) – modify the foot stiffness in order to dissipate external forces or transmit the internal force of the EFM to the foot [9]. Thus, these muscles have been the subject of study for the diagnosis and treatment of foot disorders, such as PF.

Ultrasound imaging (USI) has been employed to evaluate the muscle architecture, such as thickness and cross-sectional area (CSA), related to fascial and muscular disorders. Regarding the foot and ankle area, Wu *et al.* reported that USI was a reliable method for PF assessment [10]. In the same line, Romero-Morales *et al.* showed that the thickness of the plantar fascia at the insertion was reduced in individuals with Achilles tendinopathy compared with healthy subjects [11]. Calvo-Lobo *et al.* reported thickness alterations of the intrinsic plantar muscles in post-stroke survival [12]. Furthermore, Calvo-Lobo *et al.* observed a reduction in thickness and CSA of the FHB and AHB in patients diagnosed with hallux valgus evaluated by USI [13]. Moreover, Angin *et al.* reported an increase in the PF thickness in patients with pes planus [14]. Franettovich-Smith *et al.* showed that the assessment of the IFM by USI had high reliability within and between test sessions and thus was a reliable and clinically feasible method to assess the IFM morphology in weight bearing [15].

To the authors' knowledge, the thickness and CSA of the IFM have not been evaluated in individuals with PF. Thus, the purpose of the present study was to compare the thickness and CSA of the AHB, FHB and QP in participants with and without PF. Our hypothesis suggests that these muscle structures were altered in patients with PF.

Material and methods

Design

An observational study was performed following the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) specifications from May to August 2021.

Participants

A total of 64 volunteers from 18 to 55 years were recruited for the present study. The sample was divided in two groups: group A, composed of participants diagnosed with PF ($n = 32$), and group B, composed of healthy participants ($n = 32$). The enrollment of the participants was carried out by a specialized medical doctor with more than 10 years of experience in sport medicine. The group A participants were recruited if they met the following inclusion criteria: heel pain at least 1 month, pain with tenderness on palpation located in the middle of the plantar fascia or at the medial calcaneal tubercle, pain during the first steps in the morning or after non-weight bearing activities [16, 17]. Exclusion criteria were as follows: fracture, back or lower limb surgery, participants with any systemic disease, pregnancy, infection, plantar orthoses, length discrepancy greater than 1 cm, corticosteroid interventions, and also lower limb disturbances within the last year (e.g. ankle sprain, tendinopathy or tarsal tunnel syndrome) [18].

Sample size calculation

The sample size calculation was carried out using the G*Power software to measure the difference between the PF and healthy group using the FHB CSA variable of a pilot study ($n = 10$) that was divided into two groups (mean \pm SD): 5 participants with PF (1.46 ± 0.29) and 5 participants for the healthy group (1.65 ± 0.51). For the sample size calculation, a power of 0.80, an α error of 0.05 and an effect size of 0.63 with a one-tailed hypothesis, a power of 0.80, an α error of 0.05 and an effect size of 0.86 with a one-tailed were employed. In conclusion, a sample of 64 was calculated.

Ethics

The research ethics committee of Universidad Europea issued a favorable authorization with

code number CIPI/20/166 for the present study. All the participants signed an informed consent form before study participation. In addition, throughout the study the Helsinki Declaration for human experimentation has been respected.

Ultrasonographic assessment

For the ultrasonographic assessment an ultrasound system with a high-quality Mindray DC-60 with a 6 to 14 MHz linear transducer (L14-6NE) in B mode was employed. All the USI evaluations were performed by the same clinician with more than 3 years of experience in ultrasound imaging. In accordance with the Mickle *et al.* procedure, for the AHB, FDB, QP and FHB evaluations, subjects were laid in a supine position with the knee slightly flexed [19]. Crofts *et al.* reported a high intra- and inter-rater reliability agreement for the aforementioned USI protocol [20]. The FHB thickness was measured longitudinally along the 1st meta-

tarsal bone at the thickest portion of the muscle (Figure 1 A), and after that the transducer was rotated 90° for the CSA at the same location (Figure 1 B) [19]. The FDB thickness was measured longitudinally along a line from the medial calcaneal tubercle to the third toe at the thickest part of the muscle (Figure 1 C) and for the CSA the transducer was also rotated 90° at the same location (Figure 1 D) [19]. In this line, the QP muscle thickness (Figure 1 E) and CSA (Figure 1 F) were evaluated with the transducer in the same place of FDB and located intimately under the FDB muscle. For the AHB assessment the transducer was placed in the middle line between the origin of the muscle on the calcaneal tuberosity and the navicular tuberosity (Figure 2 A). For the AHB CSA the transducer was placed perpendicular to the longitudinal scanning axis of the foot and the anterior plane of the medial malleolus (Figure 2 B) [19]. The final scores were recorded by the mean of

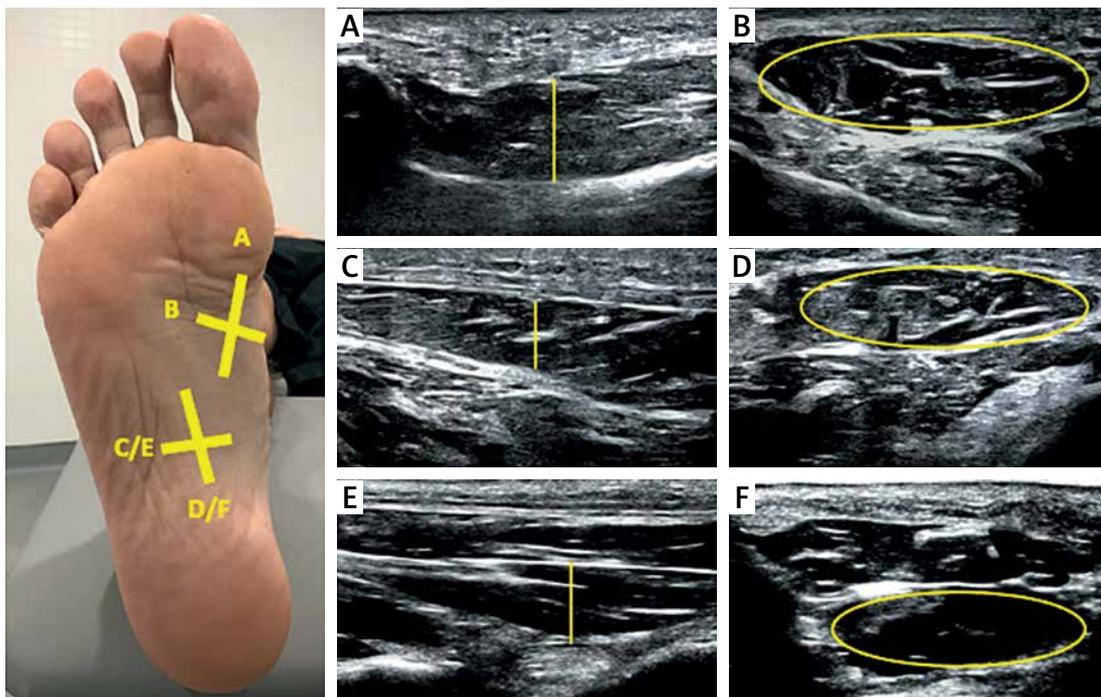


Figure 1. Ultrasound imaging of the FHB, FDB and QP in longitudinal and transversal view



Figure 2. Ultrasound imaging of the AHB in longitudinal and transversal view

3 repeated values for each measurement with the ImageJ software (Bethesda, MD, USA).

Statistical analysis

Statistical analysis was performed using IBM SPSS v.22.0 statistical package (SPSS, Inc. Chicago, IL, USA). To check the normality distribution the Kolmogorov-Smirnov test was employed. A descriptive analysis for the sample was performed. A comparative analysis between the PF and the healthy group was carried out. Mean and standard deviation (SD) and Student's *t* test for independent samples were used for the parametric data. In addition, Levene's test was performed to evaluate the equality of variances. For non-parametric data, median and interquartile range (IR) and the Mann-Whitney *U* test were performed.

A multivariate analysis was employed using linear regression to predict the influence of the descriptive data on the statistically significant outcome measurements. The dependent variables were FHB and QP CSA. The independent variables were group, sex, age, weight, height, and BMI. An α error of 0.05 (95% CI) and a desired power of 80% (β error of 0.2) were used.

Results

According to Table I, sociodemographic data showed significant differences ($p < 0.05$) in age, weight, and body mass index (BMI) between the PF and healthy group. Regarding the IFM, USI measurements for FHB CSA ($p = 0.035$) decreased, showing statistically significant differences for the PF group, while the QP CSA ($p = 0.40$) increased, showing statistically significant differences for the PF group with respect to the healthy group. The rest of the IFM did not show statistically significant differences; however, in FHB, FDB, QP and AHB thicknesses and FDB CSA a slight decrease for the PF group was observed (Table II).

Regarding the linear regression analysis (Table III), the prediction model for FHB CSA ($R^2 = 0.122$) and QP CSA ($R^2 = 0.112$) was not determined by the sociodemographic variables.

Discussion

The results of this research show that individuals with PF have a decrease of FDB CSA and an increase of QP CSA. In addition, for the PF group FHB, FDB, QP and AHB thicknesses showed a decrease for the PF group with respect to the healthy group. To our knowledge, this is the first study to analyze by USI the differences between PF and healthy participants. Plantar muscles were considered of interest for clinicians and researchers due to the influence in the ankle and foot complex and their relationship with lower limb structures and bio-

mechanics, such as the Achilles tendon [14]. Regarding the IFM and lower limb disturbances, such as Achilles tendinopathy, Romero *et al.* reported an increase of the FDB thickness, as well as FDB and FHB CSA in subjects diagnosed with Achilles tendinopathy assessed by USI [21]. However, the results of the present study showed a decrease for the FDB CSA muscle that might be explained due to these participants presenting a functional deficit in the FDB muscles that had to be compensated by the QP muscle (which increases its CSA). Plantar fascia was described as a structure capable of working with efficiency with greater loads. Therefore, in cases when there are large mechanical load fluctuations, the plantar fascia could be affected, for example, showing changes in the thickness of the plantar fascia in subjects with pes planus [22]. In the same line, Romero-Morales *et al.* reported that the thickness of the plantar fascia at the insertion was reduced in participants with Achilles tendinopathy [11]. Considering the FHB muscle, Lobo *et al.* in a previous study observed a decrease in both thickness and CSA variables in participants with hallux valgus [23]. Nevertheless, our results reported similar values with a smaller thickness and CSA in participants with PF. The fact that soft tissue and muscle alterations were described in IFM was closely related to disturbances of the ground reaction forces around the joints of the ankle and foot complex, such PF, hallux valgus, Achilles tendinopathy or pes planus. The present work highlighted the importance of the plantar muscles in the foot and ankle complex; however, it should be considered a very important issue of the foot structure, such as the foot arch, which acts as a passive system which works in a coordinated manner with the IFM muscles to provide stability in weight bearing positions. Based on this criterion, Mickle *et al.* argued that disturbances of the medial longitudinal arch could be related with PF, foot overpronation or navicular drop situations [19]. The results of the present study suggest that a decrease of the CSA of the FHB muscle could lead to an imbalance in the ankle and foot biomechanics which could be directly related to an increase of the QP CSA in order to compensate the lack of functionality and activity of the FHB muscle, expressed as architecture alterations.

The present study might be a starting point to integrate the use of USI inside the physiotherapist battery test for musculoskeletal condition diagnostics and to track the implementation of training and rehabilitation programs. In this line, the assessment by USI of the intrinsic and extrinsic plantar musculature plays an important role in patients with PF. According to the aforementioned concept, in the Huffer *et al.* systematic review, the fact of defining the IFM as a therapeutic target, for example with an exercise program, could be of

Table I. Sociodemographic data of the sample

Data	Plantar fasciitis (n = 32)	Controls (n = 32)	P-value cases vs. controls
Age [years]	43.00 ±11.00 [†]	31.00 ±9.50 [†]	0.001*
Weight [kg]	76.00 ±27.00 [†]	73.00 ±25.50 [†]	0.028 [‡]
Height [m]	1.71 ±6.3*	1.70 ±9.68*	0.465 [‡]
BMI [kg/m ²]	28.02 ±5.56*	24.69 ±5.45*	0.031 [‡]

BMI – body mass index. *Mean ± standard deviation (SD) was applied. **Student's t-test for independent samples was performed. [†]Median ± interquartile range (IR) was used. [‡]Mann-Whitney U test was utilized.

Table II. Ultrasound imaging measurements of intrinsic muscle thickness and CSA

Distance (unit)	Plantar fasciitis (n = 32)	Controls (n = 32)	P-value
FHB CSA [cm ²]	1.43 ±0.65 (1.25–1.72)*	1.76 ±0.56 (1.56–1.96)*	0.035**
FHB Th [cm]	1.13 ±0.30 (1.09–1.26)*	1.17 ±0.15 (1.11–1.22)*	0.552**
FDB CSA [cm ²]	1.57 ±0.56 (1.48–1.76)*	1.66 ±0.34 (1.53–1.78)*	0.416**
FDB Th [cm]	0.68 ±0.21 (0.64–0.77)*	0.71 ±0.14 (0.66–0.76)*	0.457**
QP CSA [cm ²]	1.23 ±0.25 (1.00–1.17)*	1.08 ±0.33 (1.14–1.33)*	0.040**
QP Th [cm]	0.74 ±0.24 (0.69–0.84)*	0.76 ±0.18 (0.70–0.83)*	0.643**
AHB CSA [cm ²]	2.03 ±0.72 (1.91–2.36)*	2.01 ±0.40 (1.86–2.15)*	0.841**
AHB Th [cm]	1.01 ±0.28 (0.97–1.11)*	1.02 ±0.24 (0.96–1.12)*	0.782**

AHB – abductor hallucis brevis, CSA – cross sectional area, FDB – flexor digitorum brevis, FHB – flexor hallucis brevis, QP – quadratus plantae. *Mean ± standard deviation (SD) (minimum- maximum) was applied. **Student's t-test for independent samples was performed. [†]Median ± interquartile range (IR) (minimum- maximum) was used. [‡]Mann-Whitney U test was utilized.

Table III. Multivariate predictive analysis for FHB and QP CSA for patients with plantar fasciitis and controls

Parameter	Model	P-value	Model R ²
FHB CSA [cm ²]	3.430		0.122
	–0.140 × Sex	0.348	
	–0.125 × Age	0.077	
	–0.25 × Weight	0.344	
	–0.010 × Height	0.476	
	0.084 × BMI	0.287	
QP CSA (cm ²)	1.351		0.112
	0.077 × Sex	0.364	
	–0.125 × Age	0.745	
	–0.25 × Weight	0.545	
	–0.010 × Height	0.961	
	0.084 × BMI	0.513	

FHB – flexor hallucis brevis, QP – quadratus plantae. *Multiply: Group (control = 0; tendinopathy = 1); Sex (women = 0; men = 1).

benefit for pain and functionality in patients with PF [24].

The results of the present study are not intended to provide a cause or explanation about the PF etiology or symptomatology. According to the aforementioned PF, several authors consider the PF a multifactorial musculoskeletal condition [6, 7]. Therefore, the measurement of the IFM by USI as a conventional assessment procedure could be beneficial for the diagnosis and management programs in individuals with PF.

This study has some limitations. First, ultrasonographic measurements were not performed by

a researcher blinded to the groups. Second, the BMI variable was measured by Quetelet's formula reporting differences between groups; thus the results of the present study could be influenced by this [25]. In this line, the weight and BMI differences might be explained by Quetelet's index calculation and also the fact that the heaviest participants could load more and increase their connective tissue work (e.g. muscles or tendons) [26].

Additional studies will be needed to gain a better knowledge about the role of the plantar muscles in individuals with PF. Electromyography or dynamometer tools could clarify the muscle acti-

vation and its influence on balance and posture. The assessment with M mode of the muscular activation and inhibition could be useful to extract more information.

In conclusion, USI measurements showed that the CSA of the FHB muscle is reduced in patients with PF while the CSA of the QP muscle is increased in patients with PF. Therefore, the assessment of the IFM by USI could be beneficial for the diagnosis and management programs in individuals with PF.

Conflict of interest

The authors declare no conflict of interest.

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