Discussing the potential impact of recent literature in rethinking conventional approaches to recalcitrant plantar fasciitis, these authors explore the use of autologous platelet concentrate injections in treating this common condition.

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Growth Factors For Chronic Plantar Fasciitis?

Plantar fasciitis is the most common condition treated by podiatric foot and ankle specialists in the United States. However, the true etiology of plantar fasciitis is still unknown and has been attributed to many different etiological factors. Even the term “plantar fasciitis” is a misnomer as the plantar fascia is really a tendonous aponeurosis and not a fascial layer.

It is entirely possible that our whole paradigm for treating plantar fasciitis is based on a false foundation, especially in light of the histological findings of Lemont, et. al., regarding specimens of resected plantar fascia. Clearly, these authors’ objective histological evidence must make all of those who treat plantar fasciitis rethink their concept of the true etiology of plantar fasciitis. This proposal that the condition we so commonly refer to as plantar fasciitis be called “plantar fasciosis” is valid and more accurately describes the condition.

These findings are further supported by histological analysis of surgical biopsies of tendons which were affected by “tendonitis,” but had no markers of inflammation. An adoption of this correct nomenclature throughout the profession is likely far into the future, but would perhaps orient our understanding of the condition better, which could possibly lead to better treatment modalities and regimens.

It is widely believed that mechanical plantar fasciitis results from repeated micro–trauma due to overuse, which results in microtears of the tissue substance until a macro injury occurs. The physiological process is then initiated via an inflammatory process – an integral part of the wound healing cascade. This injury of the plantar fascia is not dissimilar to any other musculoskeletal pathological process, such as tendonopathy.

Should We Rethink The Treatment Paradigm For Chronic Plantar Fasciitis?

Does this overwhelming data that recalcitrant plantar fasciitis (fasciosis) is not an inflammatory disorder change our paradigm for treating this condition? These findings cast doubt on two of the mainstays of conservative treatment: non-steroidal antiinflammatory agents and the corticosteroid injections. A controlled study has demonstrated that an NSAID was no better than a simple analgesic or placebo in the treatment of Achilles tendonopathy. Researchers have also demonstrated that corticosteroid injections do not change the pathological process.

Recently, there has been a push to treat recalcitrant cases of plantar fasciitis with shockwave therapy. Results have been mixed, and depending on the cited study, success rates have ranged from 48 percent to 77 percent. The mechanism of this type of therapy is unknown. However, it has been suggested that extracorporeal shockwave therapy may stimulate or reactivate the healing processes in musculoskeletal tissue. This may be due to microdestruction of avascular or minimally vascular tissues, which encourage revascularization, the release of local growth factors and the recruitment of appropriate stem cells.
cells leading to an enhancement of the intrinsic wound healing process.\textsuperscript{14} Drawbacks to this type of treatment include contraindications in patients with neuropathy, destruction of muscle tissue, development of compartment syndromes and cost concerns.\textsuperscript{15, 16}

The efficacy of ESWT may be based on an enhancement of the wound healing cascade – a conversion of a chronic wound to an acute wound which can then go through a normal physiological wound healing process. This may be similar to the phenomena that has been frequently seen among patients who have had spontaneous rupture of the plantar fascia after corticosteroid injection, but eventually have a complete resolution of symptoms after healing.

**What The Literature Reveals About Injecting Autologous Platelet Concentrate**

Taylor, et al., demonstrated in rabbit patellar tendons that injecting autologous blood was safe and that the injected tendons were stronger biomechanically than control tendons at 12 weeks.\textsuperscript{17} Researchers have documented that autologous platelet concentrate (APC+) has four to six times the normal level of growth factors, which results in fibrocyte migration and induction of neurovascular growth.\textsuperscript{18} Another group has demonstrated repair of the avascular area of dog menisci with the use of fibrin clots with a morphologically similar tissue to the resected meniscus.\textsuperscript{19}

Most importantly, researchers have demonstrated that injecting autologous blood into the origin of the extensor carpi radialis brevis is efficacious in treating recalcitrant lateral epicondylitis (tennis elbow).\textsuperscript{20} In their patient series, Edwards et al., found 79 percent of their patients were completely relieved of pain – even after strenuous activity – after one injection of autologous blood. Additionally 89 percent of their patients who required another injection reported complete resolution of symptoms.\textsuperscript{20}

The use of autologous platelet rich plasma is not a new treatment.\textsuperscript{21} The healing cascade, which is the physiological response to any injury or surgical intervention, is well documented and relies on proteins that are delivered to the healing site by platelets and white blood cells in addition to those proteins that are present in the plasma.\textsuperscript{22, 23} Successful tissue healing and regeneration requires a scaffold or matrix, undifferentiated cells and signal proteins and adhesion molecules (growth factors). It is well known that platelets affect mitogenic activity of cells like osteoblasts.\textsuperscript{24}

Bruder and Haynesworth showed platelets at baseline levels act on human mesenchymal stem cells in cell recruitment.\textsuperscript{25} They also determined concentration of platelets above baseline levels has effect on the attraction of these mesenchymal cells to the surgical site as well as proliferation of these cells.\textsuperscript{25}

We believe injecting APC+ into recalcitrant, symptomatic plantar fascia may cause a reparative effect leading to a resolution of symptoms. We have been utilizing this modality for about a year and a half. We call this technique plantar fasciorraphy.

**Taking A Closer Look At Plantar Fasciorraphy**

In order to gauge the effectiveness of this modality, we enrolled nine patients in an office study. While it is a small patient sample, we believe the study results are promising and might facilitate larger studies in the future.

We selected patients for this technique based on several factors, which included:

- a willingness to participate in an investigational technique;
- a willingness to forgo any other concomitant conservative treatment modality like NSAID’s and orthotic devices; and
- not having a cortisone injection within 90 days prior to their plantar fasciopathy.

Each individual had a significantly hypoechoic and thickened plantar fascia as demonstrated by diagnostic ultrasound. High frequency ultrasonography is well documented as a reliable and sensitive method to evaluate human plantar fascia.\textsuperscript{2, 26-29}

Using the SmartPrep\textsuperscript{®} System (Harvest Technologies), we withdrew 20 cc of each patient’s blood. It has been documented that platelets which are prepared via this system are viable and the same as they were when they were in the patient’s bloodstream. The process to harvest the platelets does not damage or change them in any way. After processing, each specimen allowed us to obtain about 3 cc of APC+.

We then anesthetized each patient with a block of posterior tibial peripheral nerve and sural nerve. We subsequently prepared the foot with a skin antiseptic.
Using high-resolution diagnostic ultrasound for guidance, we injected 3 cc of the APC+ into the most hypoechoic areas within the medial and central bands of the affected fascia with a 25-gauge needle. In order to facilitate the actual placement of the APC+, we needled the affected areas of fascia several times and than achieved the infiltration as the needle was retrograded.

After the procedure, we had the patients wear a below the knee cast immobilization boot and instructed them to avoid weightbearing for 48 hours with a subsequent increase in ambulation over the next several days. They were allowed to return to a comfortable shoe after two days. We monitored patients at one week, two weeks, four weeks, three months, six months and one year. At each visit, we documented subjective assessment with a detailed interview in addition to diagnostic imaging to access plantar fascial thickness and signal intensity.

**What The Results Of One Small Study Revealed**

We used several methods to collect information to determine the efficacy of this technique for treating plantar fasciitis (fasciosis).

First, we took ultrasound measurements of the medial, central and lateral bands prior to the injection of APC+ and obtained post-injection measurements as well for each visit. We also measured the medial, central and lateral bands of the plantar fascia of the asymptomatic foot at each visit for comparison and to serve as a control. The average thickness of the symptomatic medial band prior to injection was 7.02 mm as compared to the average thickness of the asymptomatic medial band, which was 4.88 mm. The average thickness of the central band was 6.59 mm for the symptomatic side versus 4.27 mm for the asymptomatic limb. The average thickness for the lateral band was 4.61 mm for the symptomatic limb versus 3.57 mm for the asymptomatic side.

Post-injection, we noted significant changes not only in thickness but also in the signal intensity of the fascial bands. Prior to injection, the bands of the plantar fascia near the point of origin at the calcaneus appeared thicker and more hypoechoic as compared to the asymptomatic side. One can see this in the hard copy images captured at pre-injection.

The post-injection measurements we obtained at one week, four weeks, two months, and three months revealed a significant change in thickness as well as signal intensity. At one week post-injection, the medial band showed an average reduction in thickness of 1.45 mm. By four weeks, the average reduction in thickness was 1.99 mm. At two months, it was 2.30 mm. At three months, it was 2.29 mm. The central band also decreased in thickness. There was little change in the lateral band but note these patients demonstrated the typical type of plantar fascial pain, which is located more medially at the medial tubercle than laterally. At the end of three months, the average thickness of the previously symptomatic medial and central bands was 5.03 mm and 5.39 mm respectively. The average thickness of the asymptomatic medial and central bands was 4.63 mm and 4.20 mm respectively.

Nine patients were enrolled in this study. Six patients achieved complete resolution of symptoms after two months. One patient had complete resolutions of pain after being dropped from the study due to an injection of corticosteroid. One of the remaining two patients had...
complete resolution after a second injection of APC+, and the last patient had only occasional pain when walking barefoot. Interestingly, all patients in this study had improvement that was noted on diagnostic ultrasound.

At one year, seven of nine patients had complete resolution of their plantar fascial pain for a 77.8 percent success rate, which is very comparable to the results from the lateral epicondylitis study. One patient was considered a failure and the final patient had all pain resolved, but was dismissed form the study because of a subsequent steroid injection.

Final Thoughts
The advent of new diagnostic technology in addition to the recent histological findings documented in the current literature casts serious doubt upon our current understanding of the true etiological nature and subsequent treatment of plantar fasciitis (fasciosis). Our successful early findings with the injecting APC+ indicate this may become a very commonly used modality in treating this difficult condition.

Certainly, the injection of autologous blood is safe and none of our patients experienced any complications from their plantar fasciorrhapsy. This technique cannot impair the biomechanical function of the foot, unlike other invasive procedures that transect or resect part or the entire human plantar fascia.

While these early results are encouraging, the overall efficacy of injecting APC+ into human plantar fascia for treating recalcitrant plantar fasciitis is likely to be greater when combined with a complete, multi-factorial treatment regimen versus a single isolated modality as we used in this study.

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References