

# Efficacy of Platelet-rich Plasma vs Corticosteroid Injection in Chronic Plantar Fasciitis: A Comparative Study

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## ABSTRACT

**Introduction:** Chronic plantar fasciitis (PF) is the most common cause of foot complaints making up to 11–15% of the foot symptoms requiring professional care among adults. Also, it is a common problem that affects sport participants as well as inactive middle-aged individuals. The purpose of this study was to compare the effect of steroid and platelet-rich plasma (PRP) in chronic PF. The results were assessed by comparing American orthopedic foot and ankle scale (AOFAS) and visual analogue scale (VAS) before injection, 1, 2, and 6 months after injection in both groups comprising 25 patients in each group.

**Results:** In our study of 50 patients, in steroid group-A, there is significant change in mean VAS from  $6.28 \pm 0.86$  before follow-up injection to  $2.8 \pm 0.76$  in first follow-up visit (1 month after injection) and to  $2.92 \pm 0.75$  in second follow-up visit (2 months after 1st injection) and remained constant at  $2.92 \pm 0.75$  at third follow-up (6 months post 1st injection) and significant change in mean AOFAS from  $67 \pm 10$  before injection to  $85.76 \pm 5.44$  in first follow-up visit (1 month after injection) and to  $84.16 \pm 5.94$  in second follow-up visit (2 months after first injection) and deteriorated further to  $83.92 \pm 5.84$  at third follow-up (6 months after first injection). In 25 patients in group B, there is significant change in mean VAS from  $5.8 \pm 80.78$  before follow-up injection to  $1.96 \pm 0.45$  in first follow-up visit (1 month after injection) and to  $1.96 \pm 0.45$  in second follow-up visit (2 months after first injection) and remained constant at  $1.96 \pm 0.45$  at third follow-up (6 months post first injection) and significant change in AOFAS from  $68.44 \pm 17.78$  before injection to  $89.56 \pm 0.91$  in first follow-up visit (1 month after injection) and to  $89.84 \pm 0.55$  in second follow-up visit (2 months after first injection) and improved further to  $89.92 \pm 0.40$  at third follow-up (6 months after first injection).

**Conclusion:** PRP injection is more effective than corticosteroid injection in the treatment of chronic PF in the long run.

**Keywords:** Chronic plantar fasciitis, Corticosteroid, Platelet-rich plasma.

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## INTRODUCTION

Chronic plantar fasciitis is the most common cause of foot complaints, making up 11–15% of the foot symptoms requiring medical care among adults.<sup>1</sup>

As well, it is a common problem that affects sport participants as well as inactive middle-aged individuals.<sup>2</sup> The diagnosis is grounded on typical history and the finding of localized tenderness in the middle calcaneal tubercle.<sup>3</sup> Overpulling and stretching of plantar fascia either from excessive exercise or overuse, recurrent trauma, ageing, obesity, poor fitting foot wear, or poor foot alignment while running or extended standing produce microscopic tears of collagen or cystic degeneration in the origin of plantar fascia causing pain and inflammation. The classic presentation of PF is pain on the sole of foot at the inferior region of the heel which is particularly worse with the first step taken on rising in the morning. Increasing knowledge of the pathology has led to the widespread application of a large number of conservative treatments for recalcitrant PF including physiotherapy, plantar fascia stretching exercises, ice packs, night splints, prefabricated and custom-made insert shoe modification, and nonsteroidal antiinflammatory drugs (NSAIDs).<sup>4</sup>

Local injection modalities are commonly used secondary to conservative therapies in the treatment of patients who have resistant PF. Corticosteroid injections have been used to treat plantar heel pain since the 1950s.<sup>5</sup> The advantages of corticosteroid injections include low cost, low complexity, and rapid pain relief. However, many are concerned about the potential complications associated with this treatment modality, which may offset its benefits. PRP, which is a natural concentrate of autologous growth

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factors, is now being widely tested in different fields of medicine for its possibilities in aiding the regeneration of tissue with low healing potential.<sup>6</sup> In Europe and the United States, there is an increasing prevalence of the use of autologous blood products to facilitate healing in a variety of applications. New data exist about specific growth factors, which play a crucial role in the healing process. With that knowledge, there is abundant enthusiasm in the application of concentrated platelets, which release a supramaximal quantity of these growth factors to stimulate recovery in nonhealing injuries.<sup>7</sup>

Autologous PRP was proved to improve the early neotendon properties and improve tissue healing by enhancing cellular chemotaxis, proliferation and differentiation, removal of tissue debris, angiogenesis, and the laying down of extracellular matrix. Various growth factors released from PRP are shown in Table 1.<sup>8</sup>

**Table 1:** Various growth factors released from PRP

PDGF
• Stimulates cell replication
• Promotes angiogenesis
• Promotes epithelialization
• Promotes granulation tissue formation
TGF
• Promotes formation of extracellular matrix
• Regulates bone cell metabolism
VEGF
• Promotes angiogenesis
EGF
• Promotes cell differentiation and stimulates
• Reepithelialization, angiogenesis, and collagenase activity
FGF
• Promotes proliferation of endothelial cells and fibroblasts
• Stimulates angiogenesis

Abbreviations: PDGF, platelet-derived growth factor; TGF, transforming growth factor; VEGF, vascular endothelial growth factor; EGF, epidermal growth factor; FGF, fibroblast growth factor

## MATERIALS AND METHODS

### Inclusion Criteria

Fifty patients who visited our hospital outpatient department (OPD) from 2016 to 2018 were included in our study. All patients gave informed consent to participate in the study. Patients were included if they were above 18 years of age and experienced heel pain felt maximally over the planter aspect for at least 6 months continuously. In patients with bilateral heel pain, right foot was included in our study.

### Exclusion Criteria

Patients not willing to give consent, and diabetic patients and patients with age less than 18 years were excluded in our study.

Patients were randomly divided into two groups, group A to be administered corticosteroid and group B to be administered PRP injection, consisting of 25 patients each.

Patients were given 3 doses of injection, first at date of first visit, second at first month post first injection, and third at 2 months post first injection and were followed up till 6 months post first injection. On each visit, patients were assessed with VAS and AOFAS.

PRP injection was prepared in our blood bank using centrifugation technique after withdrawing 40 mL blood from antecubital site of the patient. Blood was anticoagulated with citrate phosphate dextrose adenine (CPDA) with a ratio 1:6 to the blood. After ten minutes of centrifugation at 2,000 rpm, the blood got layered in three basic components: red blood cell, platelets, and platelet poor plasma (PPP). Because of the different sediment coefficients, the red blood cells were at the lowest level, platelets were in the middle, and the plasma was at the top. Supernatant, including platelets and plasma, were drawn from the tube, leaving red blood cells at the bottom. This supernatant containing platelets and plasma was put in other tubes and agitated for several seconds. Then it was again centrifuged, but at 2,600 rpm for 10 minutes. After centrifugation, two layers were formed in which the supernatant was PRP while the lower layer was concentrated platelets. About three quarters of the supernatant was discarded, the residual PRP (approximately 4 mL) was drawn into a syringe.

Affected foot was exposed and after proper cleaning and draping, the foot was approached through medial site using 20 G disposable steel needle. PRP was injected in the foot with the help of a 10 cc syringe. Needle was removed and aseptic dressing was done. Patient was instructed to flex and extend the ankle several times to allow equal distribution of PRP throughout. Patient was discharged from OPD with advice of cold compresses of foot for 24 hours, full weight bearing, oral antibiotics (tablet levofloxacin 750 mg OD) for three days and not to use analgesics for treatment. Patient was advised to come after one month for next injection, then after 2 months for third injection, and then as per follow-up schedule. Patient was instructed to do regular flexion and extension exercises and not to take any oral analgesic during the period of follow-up.

In case of corticosteroid injection group, affected foot was exposed and after proper cleaning and draping, the foot was approached through medial site using 20 G disposable steel needle. 2 mL methylprednisolone along with 1 mL local anaesthetic was injected in the medial aspect of the foot.

Patients were asked to come for follow-up at 0, 1, 2, and 6 month interval. On each follow-up, patients were evaluated again using VAS and ankle-hind foot scale. Adverse effects if any were also recorded.

## RESULTS

In our study, 41–50 years is the age group which got most affected in both corticosteroid (A) [48%] and PRP injection (B) [40%] groups. The mean age of the patients in group A was  $42.76 \pm 9.38$  years and in group B was  $40.40 \pm 9.95$ .

In our study, females got more affected than males in both corticosteroid (A) [56% females] and PRP injection (B) [68% females] groups. Right side got more affected than left side in both corticosteroid (A) and PRP injection (B) groups.

Mean age of the patients in groups A and B was 42.76 and 40.40 years, respectively. Mean VAS score in groups A and B before injection was 6.08 and 5.88, respectively, which improved to 2.80 and 1.96, respectively, at one month and deteriorated for group A and improved for group B, 2.92 and 1.96, respectively, at 2 months of follow-up. At the end of 6 months, mean VAS was 2.92 and 1.96, respectively, for groups A and B. Mean AOFAS ankle-hindfoot score in groups A and B before injection was 67.00 and 68.44, respectively, which improved to 85.76 and 89.56, respectively, at one month of follow-up and deteriorated for group A, 84.14 and improved for group B, 89.84. At the end of 6 months, AOFAS ankle-hindfoot score deteriorated further for group A, 83.92 and improved further for group B, 89.92.

Statistically significant reduction in heel pain was seen in group B patients, as average VAS score improved from  $5.88 \pm 0.86$  at the initiation of treatment to  $1.96 \pm 0.45$  at 1 month,  $1.96 \pm 0.45$  at 2 months. The improvement was sustained throughout the study; at 6 months, the average VAS score was  $1.96 \pm 0.45$ . Average AOFAS ankle-hindfoot score improved from  $68.44 \pm 17.8$  at the initiation of treatment to  $89.56 \pm 0.91$  at one month,  $89.84 \pm 0.55$ . At the end of 6 months, AOFAS ankle-hindfoot score further improved to  $89.92 \pm 0.40$ .

In group A patients, after initial improvement in average VAS score at 1-month outcome measure  $2.80 \pm 0.76$  from initial VAS score of  $6.08 \pm 0.86$ , no statistically significant reduction in VAS score was seen throughout the study.

Infact, VAS score increased  $2.92 \pm 0.75$  at the end of 2 months, VAS remained constant at  $2.92 \pm 0.75$  and did not improve further.

**Table 2:** Comparison of VAS at pre injection, one month post injection, 2 months post first injection, and at 6 months post first injection

		Mean	SD	Median	p value	Significance
VAS pre injection	Group A	6.0800	0.86217	6.00	<0.001	HS
	Group B	5.8800	0.78102	6.00		
VAS at one month	Group A	2.8000	0.76376	3.00		
	Group B	1.9600	0.45461	2.00		
VAS at 2 months	Group A	2.9200	0.75939	3.00		
	Group B	1.9600	0.45461	2.00		
VAS at 6 months	Group A	2.9200	0.75939	3.00		
	Group B	1.9600	0.45461	2.00		

**Table 3:** Comparison of AOFAS at pre injection, one month post injection, 2 months post first injection, and at 6 months post first injection

		Mean	S.D	Median	p value	Significance
AOFAS ankle–hindfoot score before injection	Group A	67.0000	10.00000	75.00	<0.001	HS
	Group B	68.4400	17.78595	75.00		
AOFAS ankle–hindfoot score at one month	Group A	85.7600	5.44120	88.00		
	Group B	89.5600	0.91652	90.00		
AOFAS ankle–hindfoot score at 2 months	Group A	84.1600	5.94895	88.00		
	Group B	89.8400	0.55377	90.00		
AOFAS ankle–hindfoot score at 6 months	Group A	83.9200	5.84466	85.00		
	Group B	89.9200	0.40000	90.00		

Average AOFAS ankle–hindfoot score improved from  $67.00 \pm 10.00$  at the initiation of treatment to  $85.76 \pm 5.44$  at one month. At the end of 2 months, it started deteriorating to  $84.16 \pm 5.94$  and even deteriorated further to  $83.92 \pm 5.84$  at the end of 6 months.

The *p* value for mean VAS scores before the initiation of treatment for both groups was  $>0.393$  and at 1 month was  $<0.001$ , at 2 months  $<0.001$ , and at 6 months was  $<0.001$ , which showed a highly significant statistical difference in mean VAS scores throughout the study period between the two study groups (Table 2).

The *p* value for mean AOFAS ankle–hindfoot scores before the initiation of treatment for both groups was  $>0.034$  and at 1 month  $<0.001$ , at 2 months  $<0.001$ , and at 6 months was ( $<0.001$ ), which showed a highly significant statistical difference in mean AOFAS ankle–hindfoot scores throughout the study period between the two study groups (Table 3).

## DISCUSSION

This study was designed to compare the efficacy of corticosteroid therapy to PRP therapy for chronic PF. PRP contains a more concentrated amount of platelets than does whole blood. Within platelets are powerful growth factors, including platelet-derived growth factor, transforming growth factor beta, and epidermal growth factor. The injection of PRP into the affected tissue initiates the healing stages necessary to reverse the degenerative process at the base of the plantar fascia. The individual cytokines present in the platelet alpha-granules have been shown to enhance fibroblast migration and proliferation, upregulate vascularisation, and increase collagen deposition in a variety of *in vitro* and *in vivo* settings. Additionally, many of these cytokines have been seen to work in a dose-dependent manner. The concentrated growth factors work in a synergetic manner to initiate a tendon healing response. Transforming growth factor b1 is shown to significantly

increase type I collagen production by tendon sheath fibroblasts. This same mechanism is likely to be active in chronic PF.

In our patient group, the age range varies between 20 years and 60 years of age. The range of participants age's reflects that there is little doubt that planter fasciitis pain affects the adults specially in middle to later age of life. Females were affected more than males in our study. On the basis of laterality, right side was more affected.

Many authors consider PF a degenerative tissue condition rather than inflammation at the site of origin of the plantar fascia at the medial tuberosity of the calcaneus. Degeneration of collagen occurs at the site of the lesion because of microtears of the fascia that do not heal. The histologic features of chronic PF show no inflammatory cell invasion at the site of the lesion, and the normal fascia and surrounding tissue are replaced by angiofibroblastic hyperplastic tissue. PRP injection delivers platelets with growth factors in high concentrations directly to the site of the lesion, which otherwise is inaccessible to growth factors because of hypovascularity and hypocellularity. The cytokines in platelet alpha granules affect the healing stages necessary to reverse chronic PF by enhancing fibroblast migration and proliferation, increase vascularization, and improve collagen deposition.

Complications related to corticosteroid injection in PF like plantar fascia rupture and infection have been reported in literature. But in our study, we did not encounter any such complications.

Previous studies described PRP injection as an effective treatment option for chronic PF (Table 4).

Our PRP vs steroid comparison matched the results of recent studies as that of Omar et al., who found a significant difference as regards mean VAS between the two groups, favoring the PRP group at 1.5 months follow-up ( $p < 0.05$ ) and also that of Monto who demonstrated that both PRP and steroid groups continued to improve up to 3 months and found that the improvement in the steroid group started to decline after 3 months and was sustained for longer periods in the PRP group. Our results also matched the

**Table 4:** Previous studies of PRP treatment in chronic plantar fasciitis

Study	Design	Doses of PRP	Assessment method	Follow-up	Conclusion
Shetty et al. <sup>9</sup>	Comparison between PRP injection and corticosteroid injection (60 patients)	1	VAS score FADI AOFAS score	3 months	No significant difference between PRP and corticosteroid injection
Monto et al. <sup>10</sup>	Comparison between PRP injection and corticosteroid injection (50 patients)	1	AOFAS score	Preinjection; 3, 6, 12, 24 months	PRP more effective and durable than corticosteroid injection
Kumar et al. <sup>11</sup>	PRP injection in 44 patients (50 heels)	1	VAS score AOFAS score roles and maud-sley score	6 months	PRP injection effective in treatment
Aksahin et al. <sup>12</sup>	Comparison between PRP injection and corticosteroid injection (60 patients)	1	VAS score modified roles and maud-sley score	3 week; 6 months	PRP injection as effective as corticosteroid injection
Ragab and Othman <sup>13</sup>	PRP injection in 25 patients	1	VAS score ultrasound	Preinjection; 2, 6 weeks; 6, 12 months	PRP injection effective in treatment
Barrett and Erredge <sup>14</sup>	Autologous blood injection in 9 patients	1	Ultrasound	1 week; 1, 2, 3, 12 months	Autologous blood treatment effective in treatment

Abbreviations: AOFAS, American orthopaedic foot and ankle society; FAAM, foot and ankle ability measure; FADI, foot and ankle disability index; PRP, platelet-rich plasma; SANE, single assessment numeric evaluation; SF-12v2, short form-12 health survey version 2; VAS, visual analog scale

study done by Kumar et al., on efficacy of corticosteroid and PF based on VAS score and at the end of 6 months follow-up our results matched with their results based on VAS score ( $p < 0.001$ ), where PRP results improved at one month significantly and remained constant at the end of 6 months, whereas corticosteroid group VAS score results also improved significantly at one month and deteriorated at the end of 6 months and it also matched their AOFAS scores where scores improved for PRP group constantly at each follow up and declined for steroid group at long interval ( $p < 0.001$ ) (Table 5).

The early improvement with PRP is most probably mediated by the excessive amount of growth factors and cytokines that creates an inflammatory response that subsequently restarts the cycle of tendon repair interrupting the stagnant healing environment. While with steroid injections, it only serves as an antiinflammatory agent

that ceases the inflammation early within days and has a negligible effect on regeneration, remodeling, and maturation phase which occurs at a much slower rate compared with the PRP environment rich in growth factors.

PRP was administered at the point of maximum tenderness of the heel. Some studies advocate an ultrasound-guided technique for administering injection in PF. However, Kane et al. reported no significant difference in their comparative study between ultrasound-guided and palpation-guided injection techniques in the management of idiopathic PF. In previous studies, PRP injection was administered with a peppering technique where the fascia was injected at multiple sites through a single skin portal. Other authors used a medial approach to administer PRP. It is not known whether either technique is superior. In the current study, the palpation-guided injection technique was used.

**Table 5:** Comparing the final results of various outcome studies with our study

References	Score	Baseline score outcomes	Last follow-up score outcomes	Complications
Monto et al. <sup>10</sup>	AOFAS	PRP group: 37 (range 30–56), CCS group: 52 (range 24–60)	PRP group: 92 (range 77–100), CCS group: 56 (range 30–75)	No
Kumar et al. <sup>11</sup>	AOFAS	60.6 ± 13.1	81.9 ± 16.6	No
	R and M	7.7 ± 1.4	4.2 ± 3.2	
	VAS	4 (inter-quartile 0.0)	2 (inter-quartile 1.0)	
Shetty et al. <sup>9</sup>	VAS	PRP group 8.1 ± 1.32 Steroid group 7.8 ± 1.12	PRP group 1.8 ± 1.12 Steroid group 4.27 ± 1.41	No
	AOFAS	PRP group 33.9 ± 8.15 Steroid group 32.5 ± 7.15	PRP group 83.1 ± 10.11 Steroid group 70.5 ± 9.18	
Omar et al. <sup>15</sup>	VAS	PRP group 8.2 ± 1.3, CCS group 8.8 ± 0.9	PRP group 2.6 ± 2.1, CCS group 6.5 ± 2.6	No
	FHSQ	PRP group 58.5 ± 9.6, CCS group 57.5 ± 9.4	PRP group 25.1 ± 12.4, CCS group 49.0 ± 19.1	
Our study	VAS	PRP group 5.88 ± 0.86 Steroid group 6.08 ± 0.78	PRP group 1.96 ± 0.45 Steroid group 2.92 ± 0.75	No
	AOFAS	PRP group 68.44 ± 17.7 Steroid group 67.00 ± 10.0	PRP group 89.92 ± 0.40 Steroid group 83.92 ± 5.84	

By this study, we are able to keep PRP therapy more superior than the steroid therapy for long-term benefit in planter fasciitis patients.

## CONCLUSION

Chronic heel pain is a difficult condition to treat and takes a long time to resolve. Various treatment modalities have been described in literature including surgery. But conservative methods are the treatment of choice; of these, local corticosteroid injections and PRP injections are the treatment of choice these days.

In our study, we concluded that:

- PRP is as effective as corticosteroid injection at achieving symptom relief at one and two months after first injection, for the treatment of PF, but unlike corticosteroid, its effect does not wear off with time. At 6 months follow up after first injection, PRP is significantly more effective than corticosteroid, making it better and more durable than corticosteroid injection. The PRP injection is better for long-term pain relief in planter fasciitis.
- Although there are no complications related to corticosteroids like fat pad atrophy, osteomyelitis of the calcaneus and iatrogenic rupture of the plantar fascia were observed in our study. Considering the complications mentioned in the literature, there is a need for larger sample with long-term follow-up to verify the safety of corticosteroid injection in the treatment of PF.

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