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**Vinay Kumar Tripathi**  
Assistant Professor, Department  
of Orthopaedics Mayo Institute  
of Medical Sciences Barabanki,  
Uttar Pradesh, India

**Dr. Parijat gupta**  
Professor, Department of  
Orthopaedics Mayo Institute of  
Medical Sciences Barabanki,  
Uttar Pradesh, India

## Management of diabetic foot ulcers with platelet rich plasma: A clinical study

**Vinay Kumar Tripathi and Dr. Parijat gupta**

### Abstract

**Background:** Diabetic foot is a frequent site for complication in DM. Ulceration occurs as a result of trauma in the presence of neuropathy and/or peripheral vascular disease with infection as a secondary phenomenon following disruption of the protective epidermis. The present study was conducted to determine the role of autologous platelet rich plasma in healing of diabetic foot ulcers.

**Materials & Methods:** The present study was conducted on 20 cases of diabetic foot ulcers.

In all patients, ulcer size as per maximum length and breadth and area were recorded and two PRP dressings were given on day 0 and second after first week. The size was measured in maximum length and maximum breadth by using a scale and area was calculated.

**Results:** Out of 20 patients, males were 10 and females were 10. The difference was non-significant (P=1). Age group 30-50 years had 2 males and 1 females, 50-70 years had 3 males and 5 females and >70 years had 5 males and 4 females. The difference was significant (P<0.05).

There was <60% reduction in 1 male, 61-70% reduction in 1 male and 1 female, 71-80% reduction in 2 males and 4 females and >80% reduction in 6 males and 5 females. The difference was significant (P<0.05).

**Conclusion:** PRP seem to be an effective and safe preparation used for therapy of DFU. PRP is an autologous product, therefore constitutes no risk of viral hepatitis or HIV infection.

**Keywords:** Diabetic foot, platelet rich plasma, wound

### Introduction

Diabetes mellitus is a metabolic disorder characterized with increased blood glucose level. It may be due to decrease insulin level in the body or due to inability of insulin receptors to act. Patient with diabetes is vulnerable to various complications. There are several complications of DM. Among all, diabetic foot ulcer, diabetic neuropathy, diabetic nephropathy, diabetic retinopathy and diabetic foot ulcer are common [1].

Diabetic foot is a frequent site for complication in DM. Ulceration occurs as a result of trauma in the presence of neuropathy and/or peripheral vascular disease with infection as a secondary phenomenon following disruption of the protective epidermis. The disease often leads to the development of serious health threatening complications. Of all diabetic complications, diabetic foot syndrome (DFS) is one of the most devastating [2].

In 15-25% of DM patients, diabetic foot ulceration (DFU) develops. Approximately 10-25% of those cases require amputation. Studies have shown that amputation is 20-35 times higher among patients with diabetes than without it. It has been observed that more than 65% of diabetic foot ulcers are the result of underlying neuropathy. One of the more commonly described mechanisms of action is the polyol pathway. Peripheral arterial disease (PAD) is a contributing factor to the development of foot ulcers in up to 50% of cases [3].

Among various treatment modalities for diabetic foot ulcers, use of autologous platelet-rich plasma (PRP) in the form of local application obtained by centrifugation of whole blood and addition of an activator, clotting agent is designed for the creation of local conditions favourable to healing processes [4]. The present study was conducted to determine the role of autologous platelet rich plasma in healing of diabetic foot ulcers.

### Materials & Methods

The present study was conducted in the department of orthopedics. It comprised of 20 cases of both genders. All were informed regarding the study and written consent was obtained.

### Correspondence

**Dr. Parijat gupta**  
Professor, Department of  
Orthopaedics Mayo Institute of  
Medical Sciences Barabanki,  
Uttar Pradesh, India

Ethical clearance was obtained from institutional ethical committee prior to the study. Patient with liver cell failure, patient with severe cardiomyopathy and patient with major lower limb amputation were excluded from the study.

General information such as name, age, gender etc. was recorded. In all patients, ulcer size as per maximum length and breadth and area were recorded and two PRP dressings were given on day 0 and second after first week. The size was measured in maximum length and maximum breadth by using a scale and area was calculated. Assessment of the DFUs was done as per University of Texas classification of DFU. Results thus obtained were subjected to statistical analysis for correct inferences. P value less than 0.05 was considered significant.

## Results

**Table 1:** Distribution of patients

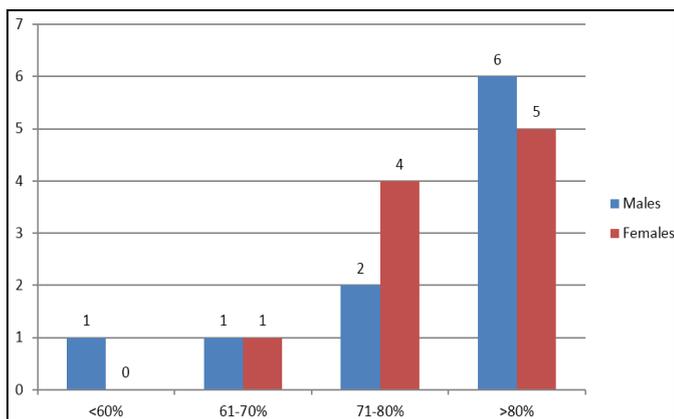
Total- 20		
Males	Females	P value
10	10	1

Table I shows that out of 20 patients, males were 10 and females were 10. The difference was non-significant (P-1).

**Table 2:** Age wise distribution of patients

Age group (years)	Males	Females	P value
30-50	2	1	0.05
50-70	3	5	
>70	5	4	

Table II shows that age group 30-50 years had 2 males and 1 females, 50-70 years had 3 males and 5 females and >70 years had 5 males and 4 females. The difference was significant (P-0.05).



**Graph 1:** Reduction in length and breadth of ulcer in patients

Graph I shows that there was <60% reduction in 1 male, 61-70% reduction in 1 male and 1 female, 71-80% reduction in 2 males and 4 females and >80% reduction in 6 males and 5 females. The difference was significant (P<0.05).

## Discussion

Diabetes mellitus is a disorder characterized by presence of multiple complications such as diabetic neuropathy, nephropathy, retinopathy and peripheral vascular disease etc. Uncontrolled diabetes contributes to the development of neuropathy and peripheral arterial disease by complex metabolic pathways. Loss of sensation caused by peripheral neuropathy, is

chaemia due to peripheral arterial disease, or a combination of these may lead to foot ulcers. Leg ulcers are classified as acute or chronic according to the duration. Chronic ulcers come with significant cost and morbidity for the patients and society. These non-healing ulcers of lower extremity develop as a result of peripheral neuropathy, ischemia, or trauma and are often difficult to treat [5].

In present study, out of 20 patients, 10 patients were seen each in males and females. We found that age group 30-50 years had 2 males and 1 females, 50-70 years had 3 males and 5 females and >70 years had 5 males and 4 females. This is in agreement with Junichi *et al.* [6].

Vickie *et al.* [7] in their study, 24 patients with non-healing ulcers of different etiologies were treated with single dose of subcutaneous PRP injections along with topical application of PRP gel under compassionate use. The mean age was 62.5±13.53 years and 24 weeks follow up period was used in the study. All the patients showed signs of wound healing with reduction in wound size, and the mean time duration to ulcer healing was 8.2 weeks. There was 5 fold increases in the platelet concentrate in the final PRP product obtained using the rapid point-of-care device, and the average platelet dose administered to the patients was 70.10×10<sup>8</sup> [8].

PRP contains additionally leukocytes that increase its antibacterial properties and synthesize interleukins as part of a nonspecific immune response. Platelets contain numerous natural growth factors released from their  $\alpha$  granulations and stimulating healing processes. PDGF is a growth factor found also in macrophages and endothelial cells. TGF is also found in macrophages, and EGF is present in macrophages, monocytes, and keratinocytes. VEGF is found mostly in endothelium, and IGF-1 is predominantly produced in the liver [8].

In present study, there was <60% reduction in 1 male, 61-70% reduction in 1 male and 1 female, 71-80% reduction in 2 males and 4 females and >80% reduction in 6 males and 5 females. This is in agreement with Knighton *et al.* [9].

It has been observed that the antibacterial effect of PRP is a result of activity of PGDF, by activation of macrophages, and VEGF, by stimulation of macrophages and monocytes. Re-epithelialisation and fibroblast proliferation are mostly the effect of PDGF, TGF, EGF, and IGF-1, stimulating the deposition of extracellular matrix (ECM). PDGF and IGF-1 are responsible for stimulation of other growth factors and cytokines, and angiogenic effect is shown mostly by VEGF, EGF, and PDGF [10].

Sunil *et al.* [11] conducted a study on 50 patients of chronic non healing ulcers who were grouped into two groups-PRP and conventional dressing group of 25 patients each. PRP group showed better results in term of healing rate and required significantly less number of dressings for wound closure. Authors suggested that PRP is a safe and effective treatment modality for chronic non-healing ulcers. PRPT is a rich source of locally active growth factors and cytokines that improve conditions of wound healing. Relatively simple and cheap production of PRP argues for continued interest in that adjunct method. It seems that specific cellular therapy constitutes an additional and valuable option in therapy of DFU resistant to the conventional therapy.

## Conclusion

PRP seem to be an effective and safe preparation used for therapy of DFU. PRP is an autologous product, therefore constitutes no risk of viral hepatitis or HIV infection.

## References

1. Saad Setta H, Elshahat A, Elsherbiny K, Massoud K, Safe I. Platelet-rich plasma versus platelet-poor plasma in the management of chronic diabetic foot ulcers: a comparative study. *Int Wound J.* 2011; 8(3):307-12.
2. Steed DL. Clinical evaluation of recombinant human platelet-derived growth factor for the treatment of lower extremity ulcers. *Plast Reconstr Surg.* 2006; 117:143-149.
3. Robert G, Frykberg Vickie R, Driver Donna Carmen. Chronic wounds treated with a physiologically relevant concentration of Platelet rich plasma gel: A prospective case series. *Ostomy Wound Management.* 2010; 56(6):36-4.
4. Brem H, Tomic-Canic M. Cellular and molecular basis of wound healing in diabetes. *J Clin Invest.* 2007; 117:1219-22.
5. Anderson I. Aetiology, assessment and management of leg ulcers. *Wound Essent.* 2006; 1:20-36.
6. Junichi Sakata, Shigeru Sasaki, Kazuyoshi Handa, Takashi Uchino, Tsukasa Sasaki, Ryuji Higashita *et al.* A retrospective, longitudinal study to evaluate healing of lower extremity wounds in patients with diabetes mellitus and ischemia using standard protocols of care and platelet-rich plasma gel in a Japanese wound care program. *Ostomy Wound Manage.* 2012; 58(4):36-49.
7. Vickie Driver R, Jason Hanft, Carelyn Fylling P. A prospective, randomized, controlled trial of autologous platelet-rich plasma gel for the treatment of diabetic foot ulcers. *Ostomy wound management.* 2006; 52(6):68-87.
8. Suryanarayan S, Budamakuntla L, Khadri SIS *et al.* Efficacy of Autologous platelet-rich plasma in the treatment of chronic non-healing leg ulcers. *Plast Aesthet Res.* 2015; 1:65-9.
9. Knighton DR, Ciresi K, Fiegel VD, Schumerth S, Butler E, Cerra F. Stimulation of repair in chronic, non-healing cutaneous ulcers using platelet derived wound healing formula. *Surg Gynecol Obstet.* 1990; 1:56-60.
10. Steed DL. Clinical evaluation of recombinant human platelet-derived growth factor for the treatment of lower extremity diabetic ulcers. Diabetic Ulcer Study Group. *J Vasc Surg.* 1995; 21(1):71-81.
11. Sunil Hanft J, Fylling P. A prospective, randomized, controlled trial of autologous platelet rich plasma for the treatment of diabetic foot ulcers. *Ostomy Wound Manage.* 2006; 52(6):68-87.