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Evaluation of Wound Healing in Diabetic Foot Ulcer Using Platelet-rich Plasma Gel: A Single-arm Clinical Trial

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Short title: PRP Gel Therapy in Diabetic Foot Ulcers

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Abstract

The aim of present study was to evaluate the effectiveness of using autologous platelet-rich plasma (PRP) gel for treatment of DFU during the first 4 weeks of the treatment. In this longitudinal and Single-arm trial, 100 patients were randomly selected after meeting certain inclusion and exclusion criteria which 70 (70% of them) patients were enrolled in the trial. After the primary caring actions such as wound debridement, the area of each wound was calculated and recorded. The PRP therapy (2ml/cm2 of ulcers) was performed weekly until the healing time for each patient. We used one sample T-test for healing wounds and Bootstrap resampling approach for reporting confidence interval with 1000 Bootstrap samples. The p-value<0.05 were considered statistically significant. The mean (SD) of DFU duration was 19.71 (4.94) weeks for units sampling. The ratio of subjects who withdraw from the study was calculated to be 2 (2.8% of them) of patients. Average area of 71 ulcers in mentioned number of cases was calculated 6.11 cm² (SD: 4.37). Also, the mean, median (SD) of healing time was 8.7, 8 (SD: 3.93) weeks except for 2 mentioned cases. According to one sample T-test, wound area (cm²) significantly decreased to 51.9 (CI: 46.7-57.1) % through the first four weeks of therapy, averagely. Furthermore, significant correlation (0.22) was not found between area of ulcers and healing duration (pvalue>0.5). As results showed, PRP could be considered as a candidate treatment for non-healing DFUs by preventing future complications such as amputation or death in this pathological phenomenon.

Key words: Diabetes, Diabetic Foot Ulcer, Platelet-Rich Plasma, Wound Healing

1. Introduction

Diabetic foot ulcer (DFU) is a common accompanying complication and the most important cause of hospitalization among diabetic patients. This phenomena with the incidence of 15% in diabetic population, is an important issue for health and care services [1-3]. During the life time of a diabetic patient, the risk of any lower extremity involvement with DFU is estimated about 25% which is affected by several risk factors including arterial disorders, peripheral neuropathy and infection. Among diabetic patients, 20% are diagnosed with inadequate blood flow and 50% with peripheral neuropathy. These incidences are totally different in DFU population so that 80% of them are suffering from both conditions [4]. Moreover, the vascular problems in these cases not only postpone the wound healing process but also hinder the reaction of immune system to the accompanying infections. The vascular complications in diabetic patients mostly develop as 3 major disorders of thrombosis or arteritis of arterioles, peripheral neuropathy (mostly due the ischemic situations) and atherosclerosis of arteries. Beside the mentioned risk factors, physical and mechanical traumas causing neuropathy in lower extremity may also lead to DFUs [5-7]. Chronic DFU is defined as an ulcer not decreased by 50% of the primary size during a month [4]. Proper treatments suggested for DFU mostly include local actions such as ulcer debridement, antibiotic therapy and bedside surgery [8, 9]. Although ulcer debridement is suggested as the primary step, it could only be helpful when patient does not suffer from arterial insufficiency. So far, different surgical methods such as percutaneous transluminal angioplasty [10], luminal stenting and arterial reconstruction surgery have been practiced in order to improve blood supply in patients with ischemic DFUs [11, 12]. Moreover, new

strategies such as hyperbaric oxygen therapy [13], bioengineered tissues [14], electrical stimulation, phototherapy [15] and platelet derived growth factors [16] are also recommended and applied. Platelet derived growth factors are biologic active compounds act in different mechanisms and pathways including activation or induction of chemotaxis, cellular proliferation and angiogenesis to induce and accelerate wound healing and are being available in clinic since 1985 [17].

The aim of this longitudinal and Single-Arm clinical trial was to evaluate the effectiveness of using autologous PRP gel for treatment of DFU during the first 4 weeks of treatment.

2. Materials and Methods

2.1. Ethics

This clinical trial was approved by Medical Ethic Committee of Research Institute for Hematology, Oncology and Stem Cell Transplantation of Shariati Hospital, Tehran University of Medical Sciences (approval code: 1394.103.3). Also all the team members considered 1975 Declaration of Helsinki and its next revisions during the trial. The aim of this study was clearly explained for each patient. Eventually, all patients were requested to sign a written consent form to declare joining this study freely after being explained (according to their knowledge) the purpose of the study. This clinical trial was registered in Iranian Registry of Clinical Trials (under supervision of Ministry of Health and Medical Education) with IRCT2015123018842N10 number. This study was a part of bigger one with a different IRCT code. But unfortunately due to the financial problems it was canceled and

removed from IRCT. Thus we had to submit for another IRCT code. Although during the study, this trial was under supervision of Ethic Committee.

2.2. Study Design and Patients

This longitudinal and Single-Arm clinical trial was conducted between May 2014 and December 2015 in Shariati Hospital (Tehran University of Medical Sciences, Tehran, Iran). In the mentioned period of time, 100 patients with chronic DFUs were selected in the study by simple random sampling. Then the inclusion and exclusion criteria were applied to all patients. Chronic DFU, as defined before, was considered the wound in an extremity (mostly foot) of a known diabetic patient which has not been decreased by 50% of the primary size in one month (Inclusion criteria). No other limitations such as age, sex, fasting blood sugar (FBS) or hemoglobin A1C (Hb A1C) levels were applied. Exclusion criteria included osteomyelitis, malignant arterial insufficiency, exposed bone in an ulcer, antibiotic resistance DFU, wounds with Charcot deformity, history of anti-proliferative medication or radiation in past 3 months, serum Hb<10 mg/dL, platelet count<10³/µL and history of growth factor therapy within last 2 weeks. Finally, according to exclusion criteria, 70 (70% of them) patients as units sampling were enrolled in the trial (Fig 1).

2.3. Primary care

First of all, demographic data and full medical history including the present illness and drug history were collected from all patients. Then primary laboratory tests such as FBS and HbA1C were re-checked. For those patients with imbalanced glucose level (abnormal FBS and/or Hb A1C), new insulin prescriptions were ordered by consulting with an expert

endocrinologist while consequent laboratory results were evaluated again. For the infection complications, a specialist was consulted with to prescribe proper antibiotics for prophylaxis in all cases (intra–venous or oral). Also debridement was performed wherever required by an expert surgeon under sedation (in some cases) in an operating room in order to remove all necrotic tissues, foreigner bodies and sinuses. Afterward, the area (length and wide) of the ulcer was measured and recorded separately for each patient by a trained physician in order to find the ulcer area. For each ulcer, the most likely geometric shape was estimated and then the area was calculated according to the appropriate shape formula (such as triangle, square, rectangular or elliptic).

2.4. Platelet rich plasma preparation

PRP preparation was performed by Rooyagen[®] PRP Gel kit (Arya Mabna Tashkhis co, Iran). According to Kit instructions, first, 27mL peripheral blood was drawn from donor using a 30mL syringe containing 3mL anticoagulant: sodium citrate. Then, the blood was shaken gently 4 times. Afterwards, it was transferred into three 10ml tubes using the transfusion Kit adaptor connected to the syringe, and centrifuged in 2000×g for 10 minutes in 24°C. After first centrifuge, the 2 fold rich platelet in the supernatant plasma was achieved. This PRP was then transferred to second tube containing 2ml 25mM CaCl2 leading to gel formation after 20 minute. Finally, the platelet gel was applied topically on the wound which would be explained in details in the next step.

2.5. Ulcer dressing and evaluation

After debridement and washing the DFU, 2 ml/cm² of PRP was applied on the ulcers as a covering layer. Then a non-absorbing wet dressing was layered on the lesion as the first contact surface. Every week until the healing was achieved, this procedure was repeated and wound area was re-measured and recorded by the physician over times. All photos were taken using a 10 mega pixel camera (Sony). At the end of the treatment, all patients were asked to follow discharge orders to prevent possible episodes of diabetic foot ulcer remission and also keep a scheduled visit with an endocrinologist. After the first four weeks, the changes between measured ulcer areas before and after treatment, were calculated by dividing the pre-treatment and post-healing (or measured ulcer area at the end of the first four weeks) areas to be considered as the effect size of PRP therapy in this study.

2.6. Statistical Methods

From the first visit to the end of the trial, all collected information were added to patient's file in Excel 2007 (Microsoft Office, Microsoft, Washington D.C, USA). According to Normality assumption of wound changes, we used one sample T-test for evaluating the wound changes in pre-treatment to 4th week of treatment and for reporting confidence interval of the effect size, Bootstrap resampling approach with 1000 Bootstrap samples was applied. The p-value<0.05 was considered statistically significant. Data were analyzed using Statistical Package for Social Sciences version 18.0 (SPSS, Inc., Chicago, IL, USA).

3. Results

During the mentioned time period, 100 patients were evaluated from which 30 were excluded according to the already defined exclusion criteria. Among the remaining 70 cases

with 71 ulcers, 58 (83%) individuals were male and 12 (17%) were female. The average age of participants (in the range of 30-79 years old) was calculated 53.8 years (SD: 10.59). As already explained, a full medical history of patients was collected and according to these histories, duration of diabetes was recorded between 2-32 years with the mean of 16.2 years (SD: 7.9). Also the average ulcer duration time was found to be 136.9 days (SD: 32.7 in the range of 74-207 days). Moreover, Hb A1C mean was 6.2 (SD: 0.65) % with Min of 4.6% and Max of 8.0%. For those cases with abnormal Hb A1C levels, new insulin orders was prescribed by an expert endocrinologist and continued to reach a proper serum glucose levels.

After measuring and calculating the ulcer area in the pre-treatment step, the median value was obtained 6.11 (SD: 4.37) cm², holding a range between 0.2 and 15.12 cm². Also the average ulcer at the end of 4th week was calculated 2.82 (SD: 2.25 in the range of 0.07-9.42) cm². At week 11 and 12, two of the patients left the study due to personal problems not related to the process of the study. At the end, 69 remaining ulcers were healed in an average of 8.7 (SD: 3.93 in the range of 4-17) week (Fig 2). Although wounds were re-measured every week as the ulcer dressing was changed, but inferable evaluation was documented at the 4th week.

According to the results of Kolmogorov-Smirnov test for normality of effect size, one sample Ttest was used. Result of this test showed that the wound area (cm2) was significantly decreased up to 51.9 (CI: 46.7-57.1) % from the beginning of treatment to the end of the first four weeks by PRP therapy, averagely. The correlation between ulcers area in pre-treatment and at the end of 4^{th} week (after treatment by PRP gel) was 0.78 (P value= .008), then the correlation between the

primary wound area and the healed time (by week) was calculated 0.22, which was not significant (P value>0.5). Also no significant relation was detected between effect size and other variables such as age, HbA1C, gender and duration of diabetes (p-values>0.05).

4. Discussion

Diabetes has been long known as a serious issue in health care system due to its growing prevalence. According to data, patients with diabetes are estimated to be 422 million (2014) [18] which could reach to 552 million people in 2030 [19]. As any other chronic disease, diabetes is associated with various complications in different organs [20, 21]. Among these complications, DFUs which almost always are observed in lower extremity occurs in 15% of diabetic population [22]. This phenomenon may lead to infection, gangrene, amputation [23] or even the patient's death [24]. This fact could reach a new level of importance when data reveals that 50-70% of lower extremity amputations are caused by DFUs [3]. In the United States, 66000 lower extremity amputations followed after untreated DFUs are recorded in 2006 [25]. In this issue any non-invasive therapeutic process is preferred to invasive ones especially when the results are predicted to be the same. So, in the cases of non-healing DFUs, amputation is considered as the last choice in the treatment process [26]. In this clinical trial, the effect of autologous PRP gel application as a non-invasive method was evaluated on the treatment of chronic DFUs. The results showed a significant decrease in the ulcer area in the pre-treatment step compared to 4th week (after beginning of treatment). Also it was shown that there is no significant correlation between primary ulcer area and the healed time which means that there is no limitation for any case with any wound area to undergo PRP therapy while considering the exclusion criteria. Thus a wide range of patients

with varying levels (are of wound) of DFUs are able to use this treatment. Also, most of the HbA1Cs did not increase dramatically before the treatments. As data showed, age and gender as confounding variables have no significant association with the ulcers area. Fortunately, at the end of the current clinical trial, amputation was not necessary in any case although no further information was collected from two leaved cases. So far, several studies including case studies and clinical trials have reported using PRP for treatment of DFUs with different success ratios. Scimeca et al successfully treated a 49-year old man with a 15cm² DFU (3 months wound) using PRP in 7 weeks [27]. Also, according to what Suresh et al [28] declared, a 57-year old man suffering from a 15 cm² non-healing DFU for 4 years was treated by PRP and healed after 7 weeks or six settings of PRP. In a clinical trial held in the United States in 2006 by Driver et al, efficacy of PRP treatment on non-healing chronic DFUs was investigated on two groups of patients: controls and standard care (PRP gel) with the wound area of 3.2 cm² (SD: 3.5) and 4cm² (SD: 5.3). In PRP treated group (72 cases) during 12 weeks, 68.4% of ulcers were healed while in controls 42% was cured [29]. Although after adjacent outliers for PRP treated group, this percentage raised to 81.3% versus 42% in controls. In another study carried out in Japan in 2012 by Sakata et al. on 39 cases with 40 DFUs in lower extremity with a wound mean of 16.8 cm² (SD 26.67), it was shown that 83% of DFUs were healed completely in 145.2 days with a significant p-value [16].

Platelet rich plasma is known as an autologous fraction from the blood containing high amounts of platelet and growth factors [30]. PRP renowned as a chemotaxis and mitogenic agent is commonly and successfully used in clinic for treatment of wound in different fields of periodontal and oral, maxillofacial, orthopedics, cosmetic and plastic surgeries [17]. In

the general aspect, the mechanism by which PRP associates with wound healing begins by α -granules degradation [31]. This phenomenon in turn activates other related growth factors including platelet derived growth factor (PDGF), vascular endothelial growth factor (VEGF), platelet derived endothelial growth factor (PDEGF), transforming growth factor β (TGF- β), epidermal growth factor (EGF), insulin like growth factor (IGF), platelet growth factor 4 (PF-4) and interleukin 1 (IL-1). All these mentioned growth factors been proved to play roles in wound healing process [17, 30]. PRP is also capable of recruiting macrophages due to its chemotaxis properties and act as an anti-infection agent as well [32].

Angiogenesis, the formation of new blood vessels from pre-existing ones mostly occurs in pathological conditions such as tumor growth, corneal neovascularization and psoriasis [33, 34]. On the other hand in physiological conditions this process is mainly responsible for two main events: wound healing and female reproductive cycle [35]. Although PRP is consists of different growth factors affecting different cells, it seems that one of the possible pathways of its action is through the induction of angiogenesis. In angiogenesis process VEGF is a strong inducer of angiogenesis that affects proliferation and migration of endothelial cells (ECs) which are two main steps of angiogenesis. Also VEGF is responsible for induction of other angiogeneic factors [36-38]. Also PDGF, TGF-ß and IGF in PRP are noted as strong angiogenic factors [39]. In this condition the new vascular branches could provide enough blood supply (both nutritional and oxygenation support) and also are able to remove dusts of wound healing process [40].

This study affirmed a highly efficient application of PRP gel in treatment and healing of chronic non-healing DFUs in a way that among all 70 evaluated cases, a relative improvement and healing results were observed.

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References

[1] Naidoo P, Liu V, Mautone M, Bergin S. Lower limb complications of diabetes mellitus: a comprehensive review with clinicopathological insights from a dedicated high-risk diabetic foot multidisciplinary team. Br J Radiol 2015;88:20150135.

[2] Association AD. Economic costs of diabetes in the US in 2007. Diabetes Care 2008;31:596-615.

[3] Boulton AJ, Vileikyte L, Ragnarson-Tennvall G, Apelqvist J. The global burden of diabetic foot disease. Lancet 2005;366:1719-1724.

[4] Ahmad J. The diabetic foot. Diabetes Metab Syndr 2015;10(1):48-60.

[5] Weledji EP, Fokam P. Treatment of the diabetic foot-to amputate or not? BMC Surg 2014;14:83.

[6] Shalaby SY, Blume P, Sumpio BE. New modalities in the chronic ischemic diabetic foot management. Clin Podiatr Med Surg 2014;31:27-42.

[7] Nickerson DS. Rationale, Science, and Economics of Surgical Nerve Decompression for Diabetic Neuropathy Foot Complications. Clin Podiatr Med Surg 2016;33:267-282.

[8] Brem H, Lyder C. Protocol for the successful treatment of pressure ulcers. Am J Surg 2004;188:9-17.

[9] Braun LR, Fisk WA, Lev-Tov H, Kirsner RS, Isseroff RR. Diabetic foot ulcer: an evidencebased treatment update. Am J Clin Dermatol 2014;15:267-281.

[10] Kerselaers L, Verbist J, Keirse K, Deloose K, Bosiers M, Peeters P. Management of Diabetic Foot following Percutaneous Transluminal Angioplasty. Urgent Interventional Therapies 2014:348.

[11] Randon C, Vermassen F, Jacobs B, De Ryck F, Van Landuyt K, Taes Y. Outcome of arterial reconstruction and free-flap coverage in diabetic foot ulcers: long-term results. World J Surg 2010;34:177-184.

[12] Sumpio BE, Lee T, Blume PA. Vascular evaluation and arterial reconstruction of the diabetic foot. Clin Podiatr Med Surg 2003;20:689-708.

[13] Fagher K, Katzman P, Löndahl M. Hyperbaric oxygen therapy reduces the risk of QTc interval prolongation in patients with diabetes and hard-to-heal foot ulcers. J Diabetes Complications 2015;29:1198-1202.

[14] Teng Y, Li Y, Wang J, Yang K, Zhang Y, Wang Y, et al. Bioengineered skin in diabetic foot ulcers. Diabetes Obes Metab 2010;12:307-315.

[15] Mulder G, Tenenhaus M, D'Souza GF. Reduction of diabetic foot ulcer healing times through use of advanced treatment modalities. Int J Low Extrem Wounds 2014;13:335-346.
[16] Sakata J, Sasaki S, Handa K, Uchino T, Sasaki T, Higashita R, et al. A retrospective, longitudinal study to evaluate healing 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;58:36-49.

[17] Lacci KM, Dardik A. Platelet-rich plasma: support for its use in wound healing. Yale J Biol Med 2010;83:1-9.

[18] Collaboration NRF. Worldwide trends in diabetes since 1980: a pooled analysis of 751 population-based studies with 4. 4 million participants. Lancet 2016;387:1513-1530.

[19] Zhou X, Guan H, Zheng L, Li Z, Guo X, Yang H, et al. Prevalence and awareness of diabetes mellitus among a rural population in China: results from Liaoning Province. Diabet Med 2015;32:332-342.

[20] Bril V. Neuromuscular complications of diabetes mellitus. Continuum 2014;20:531-544.

[21] Oei L, Rivadeneira F, Zillikens MC, Oei EH. Diabetes, diabetic complications, and fracture risk. Curr Osteoporos Rep 2015;13:106-115.

[22] Fatty N. Prevalence and Risk Factors of Diabetic Foot Problems in Taiwan. Diabetes Care 2003;26:3351.

[23] Gehling DJ, Lecka-Czernik B, Ebraheim NA. Orthopedic complications in diabetes. Bone 2016;82:79-92.

[24] Walsh J, Hoffstad O, Sullivan M, Margolis D. Association of diabetic foot ulcer and death in a population-based cohort from the United Kingdom. Diabet Med 2016. DOI: 10.1111/dme.13054

[25] Snow K. Principles of Care in the Diabetic Surgical Patient. The Diabetic Foot: Springer;2012. p. 3-16.

[26] Yazdanpanah L, Nasiri M, Adarvishi S. Literature review on the management of diabetic foot ulcer. World J Diabetes 2015;6:37.

[27] Scimeca CL, Bharara M, Fisher TK, Kimbriel H, Armstrong DG. Novel use of platelet-rich plasma to augment curative diabetic foot surgery. J Diabetes Sci Technol 2010;4:1121-1126.
[28] Suresh DH, Suryanarayan S, Sarvajnamurthy S, Puvvadi S. Treatment of a non-healing diabetic foot ulcer with platelet-rich plasma. J Cutan Aesthet Surg 2014;7:229.

[29] Driver VR, Hanft J, Fylling CP, Beriou JM. A prospective, randomized, controlled trial of autologous platelet-rich plasma gel for the treatment of diabetic foot ulcers. Ostomy Wound Manag 2006;52:68.

[30] Kabiri A, Esfandiari E, Esmaeili A, Hashemibeni B, Pourazar A, Mardani M. Platelet-rich plasma application in chondrogenesis. Adv Biomed Res 2014;3.

[31] Knighton DR, Doucette M, Fiegel VD, Ciresi K, Butler E, Austin L. The use of platelet

derived wound healing formula in human clinical trials. Prog Clin Biol Res 1987;266:319-329.

[32] Sánchez-González DJ, Méndez-Bolaina E, Trejo-Bahena NI. Platelet-rich plasma peptides:

key for regeneration. Int J Pept 2012;2012.

[33] Norooznezhad AH, Norooznezhad F, Ahmadi K. Next target of tranilast: Inhibition of corneal neovascularization. Med Hypotheses 2014;82:700-702.

[34] Norooznezhad AH, Norooznezhad F. How could cannabinoids be effective in multiple evanescent white dot syndrome? A hypothesis. J Rep Pharm Sci 2016;5:49-52.

[35] Bodnar RJ. Chemokine regulation of angiogenesis during wound healing. Adv Wound Care 2015;4:641-650.

[36] Gianni-Barrera R, Bartolomeo M, Vollmar B, Djonov V, Banfi A. Split for the cure: VEGF,PDGF-BB and intussusception in therapeutic angiogenesis. Biochem Soc Trans 2014;42:1637-1642.

[37] Hoeben A, Landuyt B, Highley MS, Wildiers H, Van Oosterom AT, De Bruijn EA.

Vascular endothelial growth factor and angiogenesis. Pharmacol Rev 2004;56:549-580.

[38] Johnson KE, Wilgus TA. Vascular endothelial growth factor and angiogenesis in the

regulation of cutaneous wound repair. Adv Wound Care 2014;3:647-661.

[39] Bir SC, Esaki J, Marui A, Yamahara K, Tsubota H, Ikeda T, et al. Angiogenic properties of sustained release platelet-rich plasma: characterization in-vitro and in the ischemic hind limb of the mouse. J Vasc Surg 2009;50:870-879. e872.

[40] Demidova-Rice TN, Durham JT, Herman IM. Wound healing angiogenesis: innovations and challenges in acute and chronic wound healing. Adv Wound Care 2012;1:17-22.

Figures legends

Fig 1. A follow diagram of study design and methodology

Fig 2. The treatment process of PRP therapy in one case (selected randomly) in different weeks from beginning of the treatment until the healing time

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