

## Platelet Rich Plasma (PRP) Therapy in Pediatric Surgical Wound Care My Experience in a Tertiary Care Hospital in Bangladesh

Dr. S.M. Khalid Mahmud<sup>1\*</sup>, Dr. Jahanara Laizu<sup>2</sup>, Prof. Dr. Aminur Rashid<sup>3</sup>, Dr. Aminul Islam<sup>4</sup>, Dr. S. M. Mahmud<sup>5</sup>

<sup>1</sup>Associate Professor, Department of Pediatrics Surgery, Bangladesh Shishu Hospital & Institute, Dhaka, Bangladesh

<sup>2</sup>Associate Professor, Department of Pharmacology, Uttara Adhunik Medical College, Dhaka, Bangladesh

<sup>3</sup>Professor and Head, Department of Pediatric Surgery, Bangladesh Shishu Hospital & Institute, Dhaka, Bangladesh

<sup>4</sup>MS Student, Department of Pediatric Surgery, Bangladesh Shishu Hospital & Institute, Dhaka Bangladesh

<sup>5</sup>Registrar, Department of Pediatric Surgery, Bangladesh Shishu Hospital & Institute, Dhaka, Bangladesh

DOI: 10.36347/sjams.2023.v11i01.019

| Received: 08.12.2022 | Accepted: 14.01.2023 | Published: 23.01.2023

\*Corresponding author: Dr. S.M. Khalid Mahmud

Associate Professor, Department of Pediatrics Surgery, Bangladesh Shishu Hospital & Institute, Dhaka, Bangladesh

### Abstract

### Original Research Article

**Background:** In recent years, platelet-rich plasma (PRP) has been used in complicated pediatric surgical wound care; burn wound repair, plastic surgery, bone and tendon ligament injury repair and other treatment. Clinical studies indicate that, PRP has a good curative effect on wound care. But in Bangladesh, we have very limited research- based information regarding the outcome of platelet rich plasma (PRP) therapy in complicated pediatric surgical wound care. **Aim of the Study:** The aim of this study was to evaluate the outcome of platelet rich plasma (PRP) therapy in complicated pediatric surgical wound care. **Methods:** This prospective observational study was conducted in Bangladesh Shishu Hospital & Institute Dhaka, Bangladesh during the period from January 2021 to June 2022. In total 34 pediatric patients with complicated surgical wound were recruited as the study population. In all cases, the treatment procedure was carried out using autologous donations. Collected data were processed, analyzed and disseminated by using MS Excel and SPSS version 23.0 program as per necessity. **Results:** In this observational study, in analyzing the final outcomes we observed that, among total 34 complicated pediatric patients with surgical wound, in more than one third of the cases (68%) wounds were healed whereas in the rest 32% cases wound were not healed. The mean  $\pm$ SD hospital staying period was found as  $6.45 \pm 2.16$  days whereas the mean  $\pm$ SD healing time was found as  $11.47 \pm 3.29$  weeks. **Conclusion:** As per the findings of this study we can conclude that, platelet-rich plasma (PRP) has a good curative effect on complicated pediatric surgical wound care. This study can provide reliable evidence for the clinical use of PRP in the clinical cure/repair of several wound.

**Keywords:** Platelet Rich Plasma, PRP, Complicated, Pediatric, Surgical wound care.

Copyright © 2023 The Author(s): This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International License (CC BY-NC 4.0) which permits unrestricted use, distribution, and reproduction in any medium for non-commercial use provided the original author and source are credited.

## INTRODUCTION

Platelet-rich plasma is well-defined as a slice of the plasma fraction of autologous blood taking a platelet concentration beyond baseline [1, 2]. Platelet-rich plasma (PRP) also has been referred to as platelet-enriched plasma, platelet-rich concentrate, autologous platelet gel, and platelet releasate [1]. Platelet releasates have been used to treat wounds since 1985 [3]. Platelet-rich plasma (PRP) serves as a growth factor agonist [4] and has both mitogenic and chemo-tactic properties [5, 6]. In accumulation to use in the treatment of chronic skin and soft tissue ulcerations [7, 8], publications regarding the use of platelet-rich plasma (PRP) include periodontal and oral surgery [9, 10] as well as maxillofacial surgery [11]. In many studies, it was

reported that, platelet-rich plasma functions as a tissue sealant and drug provision system [12], with the platelets initiating wound repair by releasing locally acting growth factors [13] via  $\alpha$ - granules degranulation [14]. Platelet-rich plasma (PRP) is easy to yield with minimal effort [15] and can be prepared as needed at the point of care [16]. In a two-step procedure, whole blood from the patient is first centrifuged to separate the plasma from packed red blood cells and then further centrifuged to separate PRP from platelet-poor plasma [17]. This focus is then activated with the addition of thrombin or calcium, causing in a gelatinous platelet gel [16, 18]. Clinically valuable Platelet-rich plasma contains at least one million platelets per microliter [16]. Not all presently marketed PRP devices are equivalent for not all concentrate viable platelets in

**Citation:** S.M. Khalid Mahmud, Jahanara Laizu, Aminur Rashid, Aminul Islam, S. M. Mahmud. Platelet Rich Plasma (PRP) Therapy in Pediatric Surgical Wound Care My Experience in a Tertiary Care Hospital in Bangladesh. Sch J App Med Sci, 2023 Jan 11(1): 120-126.

satisfactory numbers to enhance healing, with these differences accounting for many of the criticisms regarding the efficacy of platelet-rich plasma [19]. Although previous platelet-rich plasma studies have used a wide range of devices for the preparation of platelet-rich plasma [20], not all have been approved for use in humans.

## METHODOLOGY

This was a prospective observational study which was conducted in Bangladesh Shishu Hospital & Institute Dhaka, Bangladesh during the period from January 2021 to June 2022. In total 34 pediatric patients with complicated surgical wound were recruited as the study subjects. Proper written consents were taken from all the participants before data collection. In all cases, the treatment procedure was carried out using autologous donations. The whole intervention was conducted in accordance with the principles of human research specified in the Helsinki Declaration [21] and executed in compliance with currently applicable regulations and the provisions of the General Data Protection Regulation (GDPR) [22]. As per the inclusion criteria of this study, only paediatric patients with patients with complicated surgical wound were included. On the other hand, according to the exclusion criteria of this study, patients with a poor nutritional status, advanced age, over 15 years old; or inability to perform their own wound care or have it performed by a caregiver every day for an extended period until the wound healed were excluded. Donor whole blood was fractionated by centrifugation using the Spectra Optia Apheresis System (Terumo BCT, Lakewood, CO, USA). Blood was collected from large vein and transferred through a sterile device to a centrifuge spinning at 2400–2800 rpm depending on the hematocrit value, platelet count and target performance for the procedure. The blood cell separator identifies the different components, based on their weight and density. The procedure takes around 90 min and donors must meet the eligibility requirements for apheresis donation: a bodyweight > 50 kg, good general health with no abnormal lab results, age > 18 and <65 years, good venous access and no history of blood borne

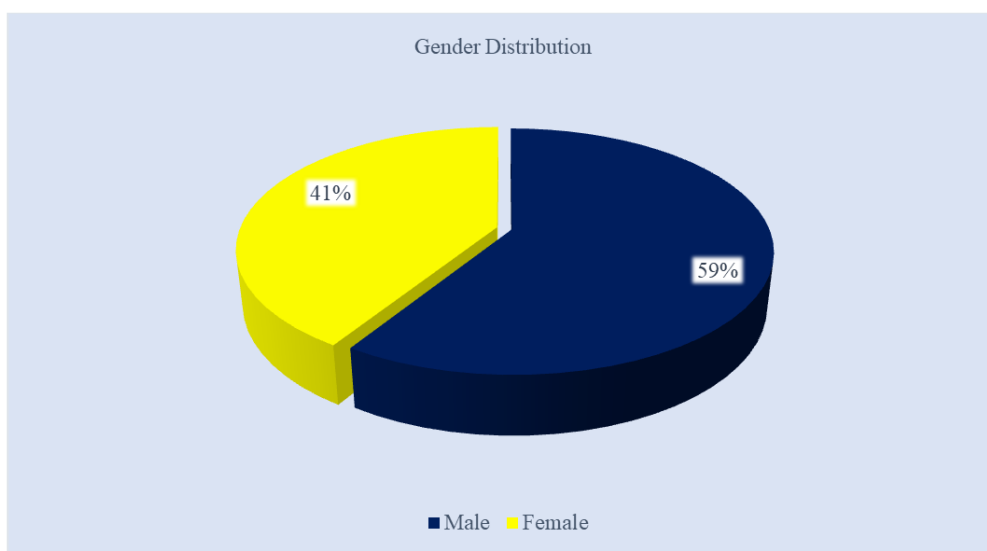
illnesses or abnormal bleeding. Once the a1-PRP was thawed, it was placed directly on the wound, left uncovered for half an hour. Subsequently, the wound and product were covered with a sterile gauze until the following day. All the demographic and clinical data of the participants were recorded. A predesigned questioner was used in data collection. All data were processed, analyzed and disseminated by using MS Excel and SPSS version 23.0 program as per necessity.

## RESULTS

In this study, among total 34 participants, 59% were male whereas the rest 41% were female. So male participants were dominating in number and the male-female ratio was 1.4:1. The mean  $\pm$ SD age of the participants was  $3.18 \pm 1.34$  years. In blood group analysis, we observed that, majority (53%) of our patients were with blood group 'O'. Besides these cases, with group 'A' and 'B' were found as 32% and 15% respectively. Rh positive cases were 79% whereas negative were 21%. The mean  $\pm$ SD haemoglobin level (g/dL) of our participants was  $13.66 \pm 1.26$  where normal status was found among 76% cases. The mean  $\pm$ SD white blood cell count ( $1 \times 10^3 \mu\text{L}$ ) of our participants was  $8.93 \pm 3.88$  where normal count was found among 62% cases. The mean  $\pm$ SD platelet count ( $1 \times 10^3 \mu\text{L}$ ) of our participants was  $256.74 \pm 189.37$  where normal count was found among 56% cases. The mean  $\pm$ SD Lymphocyte count ( $1 \times 10^3 \mu\text{L}$ ) of our participants was  $1.73 \pm 0.62$  where normal count was found among 71% cases. In this study, the mean  $\pm$ SD cholesterol level (mg/dL) of our participants was  $157.46 \pm 41.52$  where normal status was found among 56% cases. The mean  $\pm$ SD Albumin level (g/dL) of our participants was  $4.18 \pm 0.83$  where normal status was found among 91% cases. In this observational study, in analyzing the final outcomes we observed that, among total 34 complicated pediatric patients with surgical wound, in more than one third of the cases (68%) wounds were healed whereas in the rest 32% cases wound were not healed. The mean  $\pm$ SD hospital staying period was found as  $6.45 \pm 2.16$  days whereas the mean  $\pm$ SD healing time was found as  $11.47 \pm 3.29$  weeks.

**Table 1: Demographic status of participants (N=34)**

Variable	n (%) / Mean $\pm$ SD	
<b>Gender distribution</b>		
Male	20	59%
Female	14	41%
Age (Year)	$3.18 \pm 1.34$	



**Figure I: Pie chart showed gender wise participants (N=34)**

**Table 2: Baseline blood test results among participants (N=34)**

Variables	n (%) / Mean $\pm$ SD	
<b>Blood group</b>		
O	18	53%
A	11	32%
B	5	15%
<b>Rh</b>		
Positive	27	79%
Negative	7	21%
<b>Haemoglobin level (g/dL)</b>		
Mean $\pm$ SD	13.66 $\pm$ 1.26	
Normal	26	76%
Elevated	3	9%
Reduced	5	15%
<b>White blood cell count (<math>1 \times 10^3 \mu\text{L}</math>)</b>		
Mean $\pm$ SD	8.93 $\pm$ 3.88	
Normal	21	62%
Elevated	13	38%
<b>Platelet count (<math>1 \times 10^3 \mu\text{L}</math>)</b>		
Mean $\pm$ SD	256.74 $\pm$ 189.37	
Normal	19	56%
Elevated	5	15%
Reduced	10	29%
<b>Lymphocyte count (<math>1 \times 10^3 \mu\text{L}</math>)</b>		
Mean $\pm$ SD	1.73 $\pm$ 0.62	
Normal	24	71%
Lymphopaenia	10	29%
<b>Cholesterol level (mg/dL)</b>		
Mean $\pm$ SD	157.46 $\pm$ 41.52	
Normal	19	56%
Elevated	9	26%
Reduced	6	18%
<b>Albumin level (g/dL)</b>		
Mean $\pm$ SD	4.18 $\pm$ 0.83	
Normal	31	91%
Reduced	3	9%

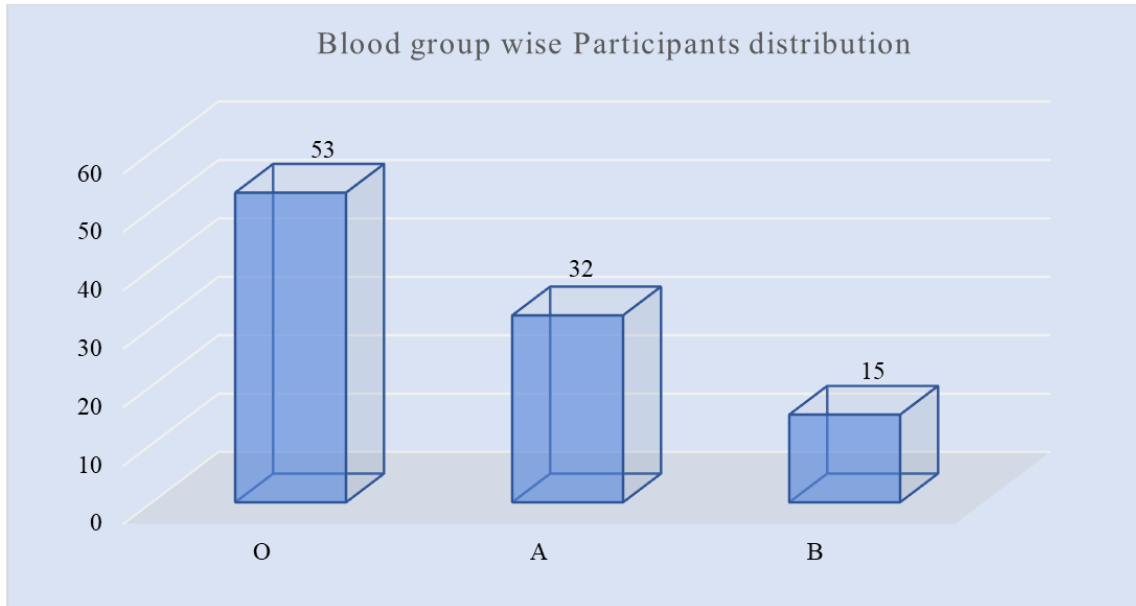


Figure II: Bar chart showed blood group wise participants blood test results (N=34)

Table 3: Outcomes among participants (N=34)

Characteristics	n (%) / Mean $\pm$ SD	
Healed	23	68%
Not healed	11	32%
Hospital staying (Day)	6.45 $\pm$ 2.16	
Healing time (Week)	11.47 $\pm$ 3.29	

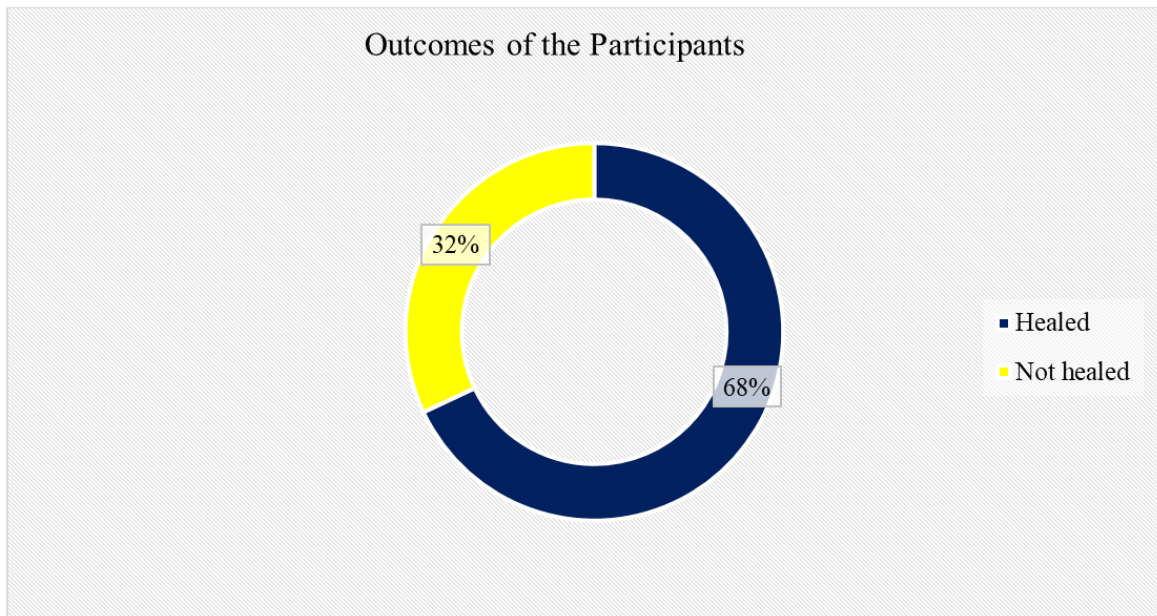


Figure III: Ring chart showed outcomes among participants (N=34)



1



2

**Picture 1 & 2: PRP Surgical Procedures**



3



4

**Picture 3 & 4: PRP Surgical Procedures**

## DISCUSSION

The aim of this study was to evaluate the outcome of platelet rich plasma (PRP) therapy in complicated pediatric surgical wound care. In this study, among total 34 participants, 59% were male whereas the rest 41% were female. So male participants were dominating in number and the male-female ratio was 1.4:1. The mean  $\pm$ SD age of the participants was

3.18 $\pm$ 1.34 years. In blood group analysis, we observed that, majority (53%) of our patients were with blood group 'O'. Besides these cases, with group 'A' and 'B' were found as 32% and 15% respectively. Rh positive cases were 79% whereas negative were 21%. By promoting safe and natural healing, platelet rich plasma is a promising alternative to the current standard of care for hard-to-heal wounds such as those patients, many having a history of DM and/or treatment with

hydroxyurea and all a poor response to other treatments [23]. It is obtained from density gradient centrifugation of peripheral venous blood and contains a wide variety of growth factors, fibroblasts, mesenchymal stem cells, and white blood cells [24]. Some studies have confirmed that, PRP can promote wound healing, inhibit the growth of various bacterial species, improve angiogenesis and reduce postoperative pain [25, 26]. In this observational study, in analyzing the final outcomes we observed that, among total 34 complicated pediatric patients with surgical wound, in more than one third of the cases (68%) wounds were healed whereas in the rest 32% cases wound were not healed. The mean  $\pm$ SD hospital staying period was found as  $6.45\pm 2.16$  days whereas the mean  $\pm$ SD healing time was found as  $11.47\pm 3.29$  weeks. There is growing evidence that, platelet rich plasma (PRP) improves the healing of chronic wounds and the use of this product is increasing in the field of wound repair [27]. In a study, it was reported that, the rate of proliferation of granulation tissue was significantly faster in al-PRP-treated patients than in controls during the first 2 weeks of treatment, suggesting that, al-PRP can stimulate granulation tissue proliferation in the early stages of healing [28]. Recently, significantly faster healing has been found with either al- or au-PRP, wounds healing in a mean of <60 days versus >85 days with conventional wound care [29]. Some systematic reviews as well as meta-analyses have found that au-PRP therapy improves the rate of complete and partial healing of hard-to-heal wounds compared to standard wound care [30, 31].

#### Limitation of the Study

Though it was a single centered study with small sample size, was conducted in a very short period of time the findings of this study may not reflect the exact scenario of the whole country.

## CONCLUSION & RECOMMENDATION

As per the findings of this study we can conclude that, platelet-rich plasma (PRP) has a good curative effect on complicated pediatric surgical wound care. This study can provide reliable evidence for the clinical use of PRP in the clinical cure/repair of several wound. We recommend for wider use of platelet-rich plasma (PRP) in several wound care.

## REFERENCES

1. Mehta, S., & Watson, J. T. (2008). Platelet rich concentrate: basic science and current clinical applications. *J Orthop Trauma*, 22(6), 432-8.
2. Marx, R. E. (2001). Platelet-rich plasma (PRP): what is PRP and what is not PRP? *Implant Dent.*, 10(4), 225-8.
3. Driver, V. R., Hanft, J., Fylling, C. P., & Beriou, J. M. (2006). Autologous Diabetic Foot Ulcer Study Group. A prospective, randomized, controlled trial of autologous platelet-rich plasma gel for the treatment of diabetic foot ulcers. *Ostomy Wound Manage*, 52(6), 68-70, 72, 74 passim.
4. Petrova, N., & Edmonds, M. (2006). Emerging drugs for diabetic foot ulcers. *Expert Opin Emerg Drugs*, 11(4), 709-24.
5. Millington, J. T., & Norris, T. W. (2000). Effective treatment strategies for diabetic foot wounds. *J Fam Pract.*, 49(11 Suppl), S40-8.
6. Steed, D. L., Goslen, J. B., Holloway, G. A., Malone, J. M., Bunt, T. J., & Webster, M. W. (1992). Randomized prospective double-blind trial in healing chronic diabetic foot ulcers. CT-102 activated platelet supernatant, topical versus placebo. *Diabetes Care*, 15(11), 1598-1604.
7. Pietrzak, W. S., & Eppley, B. L. (2005). Platelet rich plasma: biology and new technology. *J Craniofac Surg.*, 16(6), 1043-54.
8. Salemi, S., Rinaldi, C., Manna, F., Guarneri, G. F., & Parodi, P. C. (2008). Reconstruction of lower leg skin ulcer with autologous adipose tissue and platelet-rich plasma. *J Plast Reconstr Aesthet Surg.*, 61(12), 1565-7.
9. Lindeboom, J. A., Mathura, K. R., Aartman, I. H., Kroon, F. H., Milstein, D. M., & Ince, C. (2007). Influence of the application of platelet-enriched plasma in oral mucosal wound healing. *Clin Oral Implants Res.*, 18(1), 133-9.
10. El-Sharkawy, H., Kantarci, A., Deady, J., Hasturk, H., Liu, H., Alshahat, M., & Van Dyke, T. E. (2007). Platelet-rich plasma: growth factors and pro-and anti-inflammatory properties. *Journal of periodontology*, 78(4), 661-669.
11. Nikolidakis, D., & Jansen, J. A. (2008). The biology of platelet-rich plasma and its application in oral surgery: literature review. *Tissue Engineering Part B: Reviews*, 14(3), 249-258.
12. Eppley, B. L., Woodell, J. E., & Higgins, J. (2004). Platelet quantification and growth factor analysis from platelet-rich plasma: implications for wound healing. *Plastic and reconstructive surgery*, 114(6), 1502-1508.
13. Knighton, D. R., Ciresi, K. F., Fiegel, V. D., Austin, L. L., & Butler, E. L. (1986). Classification and treatment of chronic nonhealing wounds. Successful treatment with autologous platelet-derived wound healing factors (PDWHF). *Annals of surgery*, 204(3), 322.
14. Knighton, D. R., Doucette, M., Fiegel, V. D., Ciresi, K., Butler, E., & Austin, L. (1988). The use of platelet derived wound healing formula in human clinical trials. *Prog Clin Biol Res.*, 266, 319-29.
15. Bhanot, S., & Alex, J. C. (2002). Current applications of platelet gels in facial plastic surgery. *Facial Plast Surg.*, 18(1), 27-33.
16. Mishra, A., Woodall, J. Jr., & Vieira, A. (2009). Treatment of tendon and muscle using platelet-rich plasma. *Clin Sports Med.*, 28(1), 113-25.

17. Gandhi, A., Bibbo, C., Pinzur, M., & Lin, S. S. (2005). The role of platelet-rich plasma in foot and ankle surgery. *Foot Ankle Clin.*, 10(4), 621-37, viii.
18. Rozman, P., & Bolta, Z. (2007). Use of platelet growth factors in treating wounds and soft-tissue injuries. *Acta Dermatovenerol Alp Panonica Adriat.*, 16(4), 156-65.
19. Marx, R. E. (2004). Platelet-rich plasma: evidence to support its use. *J Oral Maxillofac Surg.*, 62(4), 489-96.
20. Mazzucco, L., Balbo, V., Cattana, E., & Borzini, P. (2008). Platelet-rich plasma and platelet gel preparation using Plateltex. *Vox Sang.*, 94(3), 202-8.
21. World Medical Association. (2001) . World Medical Association Declaration of Helsinki. Ethical principles for medical research involving human subjects. Bulletin of the World Health Organization, 79 (4) , 373 - 374. World Health Organization.  
<https://apps.who.int/iris/handle/10665/268312>.
22. Voigt, P., & Axel von dem, B. (2017). "Enforcement and fines under the GDPR." The EU General Data Protection Regulation (GDPR). Springer, Cham, 201-217.
23. Vidán-Estévez, J., Sánchez- Herráez, S., Escalante-Barrigón, F., & Seco-Calvo, J. (2021). Healing of Chronic Wounds with Platelet-Derived Growth Factors from Single Donor Platelet-Rich Plasma following One Freeze-Thaw Cycle. A Cross-Sectional Study. *J. Clin. Med.*, 10, 5762. <https://doi.org/10.3390/jcm10245762>.
24. Marx, R. E. (2001). Platelet-rich plasma (PRP): What is PRP and what is not PRP? *Implant. Dent.*, 10, 225–228. [CrossRef] [PubMed].
25. de Leon, J. M., Driver, V. R., Fylling, C. P., Carter, M. J., Anderson, C., Wilson, J., ... & Rappl, L. M. (2011). The clinical relevance of treating chronic wounds with an enhanced near-physiological concentration of platelet-rich plasma gel. *Advances in skin & wound care*, 24(8), 357-368. [CrossRef].
26. Serra, R., Buffone, G., Dominijanni, A., Molinari, V., Montemurro, R., & de Franciscis, S. (2013). Application of platelet-rich gel to enhance healing of transmetatarsal amputations in diabetic dysvascular patients. *Int. Wound J.*, 10, 612–615. [CrossRef] [PubMed].
27. Xia, Y., Zhao, J., Xie, J., Lv, Y., & Cao, D. S. (2019). The Efficacy of Platelet-Rich Plasma Dressing for Chronic Nonhealing Ulcers: A Meta-Analysis of 15 Randomized Controlled Trials. *Plast. Reconstr. Surg.*, 144, 1463–1474. [CrossRef].
28. McAleer, J. P., Sharma, S., Kaplan, E. M., & Persich, G. (2006). Use of autologous platelet concentrate in a nonhealing lower extremity wound. *Adv Skin Wound Care*, 19(7), 354-63.
29. Rozman, P., & Bolta, Z. (2007). Use of platelet growth factors in treating wounds and soft-tissue injuries. *Acta Dermatovenerol Alp Panonica Adriat.*, 16(4), 156-65.
30. O'Connell, S. M., Impeduglia, T., Hessler, K., Wang, X. J., Carroll, R. J., & Dardik, H. (2008). Autologous platelet-rich fibrin matrix as cell therapy in the healing of chronic lower-extremity ulcers. *Wound Repair and Regeneration*, 16(6), 749-756.
31. Margolis, D. J., Bartus, C., Hoffstad, O., Malay, S., & Berlin, J. A. (2005). Effectiveness of recombinant human platelet-derived growth factor for the treatment of diabetic neuropathic foot ulcers. *Wound repair and regeneration*, 13(6), 531-536.