



COMPARATIVE EVALUATION OF AN ALLOPLASTIC GRAFT (RESORBABLE HYDROXYAPATITE GRAFT) AND AUTOLOGOUS PLATELET RICH FIBRIN IN THE TREATMENT OF INTRABONY DEFECTS – A CLINICO – RADIOGRAPHIC STUDY

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ABSTRACT

Background: The primary goal of periodontal treatment is the maintenance of the natural dentition in health and comfortable function. Hydroxyapatite (HA) bone grafting material has been used to fill periodontal intrabony defects, which has resulted in clinically acceptable responses. Platelet rich fibrin (PRF) developed in France described by CHOUKRON et al is a second generation platelet concentrate widely used to accelerate soft and hard tissue healing. **Method:** The study sample includes 23 patients (which included 30 intrabony sites) with the age group of 20-55 years. The surgical sites were randomly divided into 3 groups, group A treated by OFD, group B treated by alloplast (resorbable HA) and group C treated by PRF. Clinical parameters (PI, PPD, CAL, GML) were recorded at baseline, 3 months, 6 months and 9 months post operatively. Radiographic parameters were recorded at baseline and 9 months post operatively. **Results:** Mean PPD reduction was 1.8(0.74) in group A; 2.8(0.74) in group B; 4.1(0.83) in group C. while mean CAL gain was 1.7(0.64) in group A; 2.8(0.65) in group B; 2.8(0.83) in group C. furthermore, significantly greater percentage of mean bone fill was found to be 66.39(17.28)% in group C compared to 50.0(7.45)% in group B and 13.7(19.73)% in group A. **Conclusion:** PRF can be used as a sole grafting material, useful alternative and economical to alloplastic graft in the treatment of intrabony defects.

KEYWORDS : Open Flap Debridement, Hydroxyapatite, Platelet Rich Fibrin, Intrabony Defect Depth, Percentage of Bone Fill.

INTRODUCTION

Periodontics is an infectious disease that cause destruction of the tooth – attachment apparatus.¹ Periodontal therapy is performed with the primary objective of gaining access to the disease sites, achieving reduction in pocket depth, arresting further disease progression and finally restoring the periodontal tissues lost due to disease process.²

Conventional flap debridement falls short of regenerating tissues destroyed by the disease, and current regenerative procedures offers a limited potential towards attaining complete periodontal restoration.⁴ Various biomaterials⁵⁻⁸ based on endogenous regenerative technology (ERT), have been used for periodontal tissue regeneration in addition to autogenous^{9,10} and allogenic bone grafts,^{11,12} but not a single graft material is considered as a gold standard in the treatment of intrabony defects.

Porous hydroxyapatite (HA) bone graft material has been used to fill periodontal intrabony defects, which has resulted in clinically acceptable responses.¹³ Hydroxyapatite (HA) has good biocompatibility when in contact with bone as its chemical composition is similar to that of bone. They are biocompatible, non-toxic, resorbable, and non-inflammatory, cause no immunological, foreign body or irritating response, and have excellent osteoconductive ability.^{14,15,16}

Among platelet concentrates, platelet rich fibrin (PRF) belongs to a group of second generation blood autologous preparation that was originally described by Choukroun¹⁷ et al in 2001 is defined as an autologous leukocyte and PRF biomaterial. First biochemical analysis of the PRF composition indicated that this biomaterial consists of an intimate assembly of cytokines, glycanic chains and structural glycoproteins enmeshed within a slowly polymerized fibrin network. The biologic activity of the fibrin molecule is enough in itself to account for the significant cicatrizing capacity of the PRF and the slow polymerization mode confers to the PRF membrane, a particularly favourable physiologic architecture to support the healing process.¹⁸

The use of PRF and HA bone graft for periodontal regenerative therapy offer an interesting and a potentially clinical useful modality to the clinician in treating periodontal osseous defects. Thus the purpose of the present study was to investigate the efficacy of autologous PRF and alloplastic graft (resorbable HA) bone graft in comparison to open flap debridement (OFD) alone in the treatment of intrabony defects.

MATERIALS AND METHODS

A clinical and radiographic study was designed and conducted at Department of Periodontics, HKE Society's S. Nijalingappa Institute of Dental Sciences and Research, Gulbarga.

The study included 30 surgical intrabony sites. Both males and females

between the age group of 20-55 years of age were included in the study.

The project was reviewed by the board of ethical committee at H.K.E Society's S. Nijalingappa Institute of Dental Sciences and Research, Gulbarga and the project was envisaged after the clearance was obtained. The patients were informed about the study design and verbal and written informed consents were obtained from all subjects prior to their enrollment in the study.

Total of 30 surgical intrabony sites included in the study were divided into 3 groups Group A (n=10 sites) – intrabony defects treated by open flap debridement. Group B (n=10 sites) – intrabony defects treated by alloplastic graft (resorbable hydroxyapatite). Group C (n=10 sites) – intrabony defects treated by autologous platelet rich fibrin (PRF).

Clinical Procedure: The defect sites was anaesthetized with 2% lignocaine. Full thickness Mucoperiosteal flap was reflected, and root surface debrided. Defect sites in each patient were randomly divided into 3 groups to be treated with OFD, resorbable Hydroxyapatite and Platelet Rich Fibrin respectively. Sutured and Periodontal dressing placed and post – operative instructions given. Patient was recalled after 7 days for suture removal. Post Surgical evaluation and follow up at 3 months, 6 months and 9 months was done. Clinical parameters were recorded using UNC- 15 probe and customized acrylic stents at 3, 6 and 9 months. Radiographic parameters were recorded using IOPA grids at baseline and at 9 months post operatively.

PRF Preparation: Preparation of PRF follows the protocol developed by Choukroun et al in Nice France. The steps involved are as follows:

- Blood specimen is collected or drawn from the patient.
- The blood specimen is placed in the centrifuge and the tubes were immediately centrifuged at 3,000 rpm for 10 minutes at room temperature.

The layers that are formed are:-

- The lower fraction containing the RBC's.
- The middle fraction containing the fibrin clot.
- The upper fraction containing the straw colored platelet poor plasma (PPP).

Radiographic parameter: Conventional IOPA radiograph with IOPA grid were taken using paralleling cone technique of the surgical sites at baseline and 9 months post operatively and following measurements were made.

- Distance from cemento-enamel junction to base of the defect (CEJ to BD).
- Distance from cemento-enamel junction to the alveolar crest (CEJ to AC).
- Intra Bony Defect (IBD) depth = CEJ to BD – CEJ to AC.

RESULTS

Clinical Assessment: Demographic data - In group A 2 males (28.71%) and 5 females (71.43%) were included in this group (table 1a & fig 1a). The mean age of patients was 28.71 years and the age range was 20-35 years (table 1b & fig 1b). 4 surgical sites were from upper and lower single rooted teeth and remaining 6 surgical sites were from upper and lower multi-rooted teeth. In group B 4 males (44.44%) and 5 females (55.56%) were included in this group (table 1a & fig 1a). The mean age of patients was 38 years and the age range was 20-54 years (table 1b & fig 1b). 1 surgical site was from lower anterior and remaining 9 surgical sites were from upper and lower multi-rooted teeth. In group C 4 males (57.14%) and 3 females (42.86%) were included in this group (table 1a & fig 1a). The mean age of patients was 31 years and the age range was 20-54 years (table 1b & fig 1b). 1 surgical site was from lower anterior and remaining 9 surgical sites were from upper and lower multi-rooted teeth.

Table 2 and fig 2 shows the mean and standard deviation of the plaque index for all the 3 groups at baseline, 3 months, 6 months and 9 months post operatively. A statistically significant reduction in the Plaque Index was observed in all the three groups from baseline to 9 months postoperatively. There was no statistically significant difference when compared between the groups.

Table 3 and fig 3 shows the mean and standard deviation of Pocket Probing Depth (PPD) for all 3 groups at baseline, 3 months, 6 months and 9 months post Operatively. Table 3 also shows the comparison between the groups by one way ANOVA test. There was statistically significant difference only in the 9th month between the groups (F value=4.29).

Table 4 and fig 4 shows the mean and standard deviation of clinical attachment level (CAL) for all 3 groups at baseline, 3 months, 6 months and 9 months post operatively. Table 4 also shows the comparison between the groups by one way ANOVA test. There was statistically significant difference at baseline between the groups (F value=4.32).

Table 5 and fig 5 shows the mean and standard deviation of gingival margin level at baseline, 3 months, 6 months and 9 months post operatively. Table 5 also shows the comparison between the groups by one way ANOVA test. There was no statistically significant difference at all periods between the groups. Table 5(a) shows the mean and standard deviation of changes in the clinical parameter with respect to baseline. Table 5(b) shows the comparison of changes in the clinical parameters (with respect to baseline) between groups. There was no statistically significant difference between group A and group C.

Radiographic Assessment: Table 6 and fig 6 shows the mean and standard deviation of CEJ to the base of the defect of all 3 groups at baseline, 3 months, 6 months and 9 months post operatively. There was statistically significant difference in group B [5.8(1.17) mm to 3.6(0.67) mm] with $p < 0.05$ respectively. There was statistically significant difference in group C [7.5(1.43) mm to 3.1(1.14) mm] with $p < 0.05$ respectively.

Table 7 and fig 7 shows the mean and standard deviation of CEJ to the alveolar crest of all 3 groups at baseline and 9 months post operatively. There was statistically significant difference in group B [3.4(0.66) mm to 2.4(0.66) mm] with $p < 0.05$ respectively. There was statistically significant difference in group C [3.6(0.92) mm to 1.6(0.92) mm] with $p < 0.05$ respectively. There was statistically significant difference between the groups at baseline (F value = 3.57) and 9 months (F value = 5.04) respectively.

Table 8 and fig 8 shows the mean and standard deviation of intra bony defect (IBD) depth of all 3 groups at baseline and 9 months post operatively. There was statistically significant difference in the means of group B [2.4(0.66) mm to 1.2(0.4) mm] with $p < 0.05$ respectively. There was statistically significant difference in the means of group C [3.9(1.14) mm to 1.5(0.81) mm] with $p < 0.05$ respectively. There was statistically significant difference between the periods in group B (T value = 4.65) and in group C (T value = 4.44). There was statistically significant difference between the groups at baseline (F value = 5.02) and 9 months (F value = 2.39) respectively.

Table 9 shows the mean and standard deviation of changes in the radiographic parameters with respect to baseline. The mean values of intra bony defect (IBD) depth of group A was 0.3(0.46) mm; group B

was 1.2(0.4) mm; group C was 3.0(1.79) mm with $p < 0.05$ respectively. The percentage of bone fill in group A was 13.7(19.73)%; group B was 50.0(7.45)%; group C was 66.39(17.28)% with $p < 0.05$ respectively.

Table 10 shows the changes in the radiographic parameters with respect to baseline between the groups. Mean IBD depth reduction: There was statistically significant difference between group A and group B with T value of 4.44; between group B and group C with T value of 4.38; between group A and group C with T value of 2.94. Mean percentage of Bone defect fill: There was statistically significant difference between group A and group B with T value of 5.06; between group B and group C with T value of 6.03; between group A and group C with T value of 2.61.



Fig 1: Armamentarium



Fig 2: UNC 15 Probe

Fig 3: Group A (Open Flap Debridement)



Fig 3 a: Detection of Periodontal Pocket



Fig 3 b: Flap Reflection and Debridement



Fig 3 c: Sutures Given

Figure 4: Group B (Resorbable hydroxyapatite)



Fig 4a: Detection of Periodontal Pocket



Fig 4b: Flap Reflection and Debridement

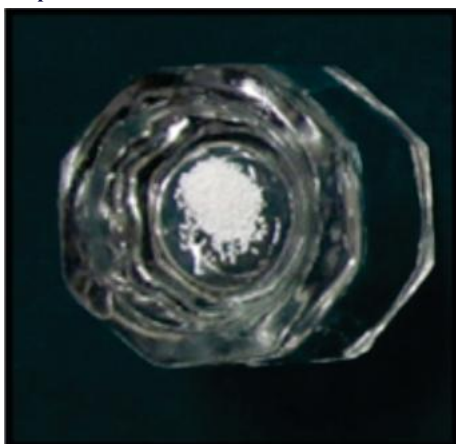


Fig 4c: Resorbable Hydroxyapatite Graft



Fig 4d: Application of Bone graft



Fig 4d: Sutures Given



Fig 4e: Periodontal Pack given

Figure 5: Group C (Platelet Rich Fibrin)



Fig 5a: Detection of Periodontal Pocket



Fig 5b: Flap Reflection and Debridement

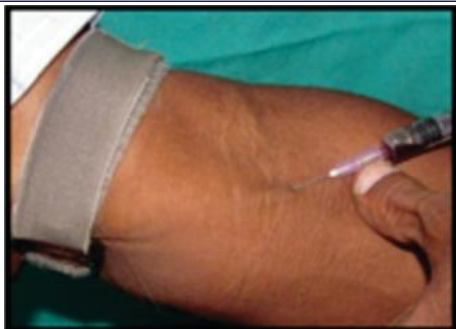


Fig 5c: Collection of Blood



Fig 5d: Centrifugation Machine

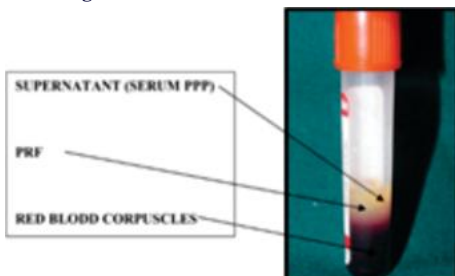


Fig 5e: The 3 Layers Formed

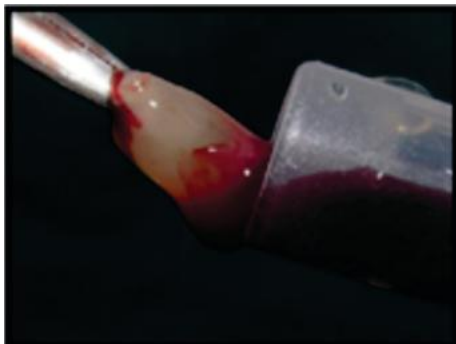


Fig 5f: PRF Obtained



Fig 5g: Application of PRF



Fig 5h: Sutures Given



Fig 5i: Periodontal Pack Give

Figure 6: Radiographic Assesment

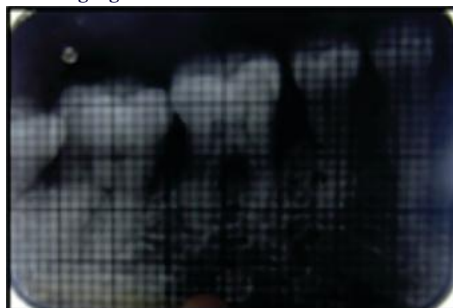


Fig 6a: Group A Baseline



Fig 6b: Group A 9 Months Post Operatively

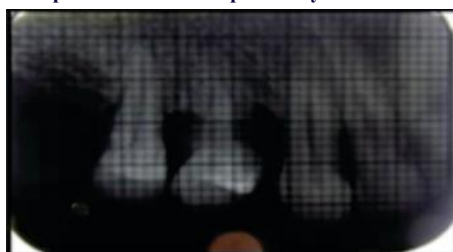


Fig 6c: Group B Baseline



Fig 6d: Group B 9 Months Post Operatively

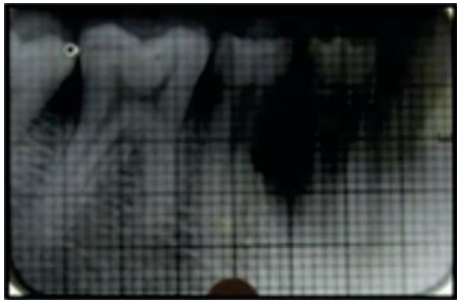


Fig 6e: Group C Baseline



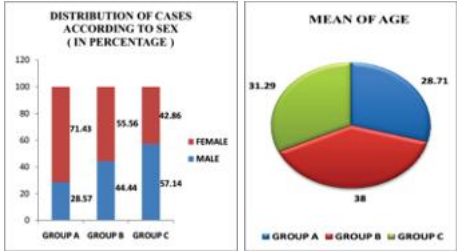
Fig 6f: Group C 9 Months Post Operatively

Table 1(a) Distribution of Cases According to Sex

GROUP	MALE		FEMALE	
	NUMBER	PERCENTAGE	NUMBER	PERCENTAGE
A	02	28.57	05	71.43
B	04	44.44	05	55.56
C	04	57.14	03	42.86

Table 1(b) Mean and Standard Deviation (sd) of Ages

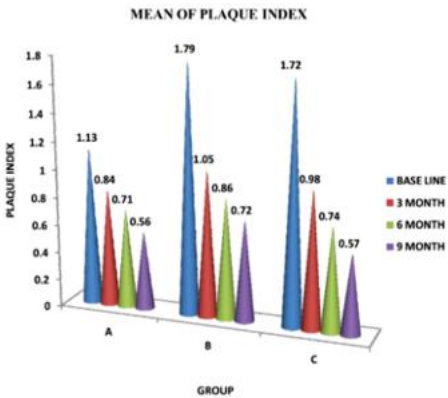
GROUP	AGE (IN YEARS)	
	MEAN	SD
A	28.71	4.16
B	38.00	11.83
C	31.29	11.99



Graph 1(a) Graph 1(b)

Table 2 Mean and Standard Deviation (sd) of Plaque Index and Comparison Between the Groups

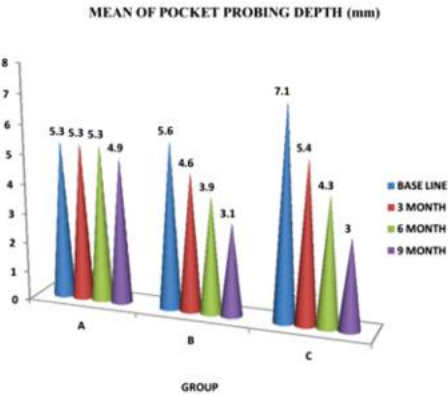
GROUP	BASE LINE		3 MONTH		6 MONTH		9 MONTH	
	MEAN	SD	MEAN	SD	MEAN	SD	MEAN	SD
A	1.13	0.33	0.84	0.28	0.71	0.21	0.56	0.26
B	1.79	1.05	1.05	0.48	0.86	0.39	0.72	0.41
C	1.72	1.23	0.98	0.49	0.74	0.29	0.57	0.25
F VALUE	0.93(NS)		0.46(NS)		0.46(NS)		0.57(NS)	



Graph 2

Table 3 Mean and Standard Deviation (sd) of Pocket Probing Depth (ppd) and Comparison Between the Groups

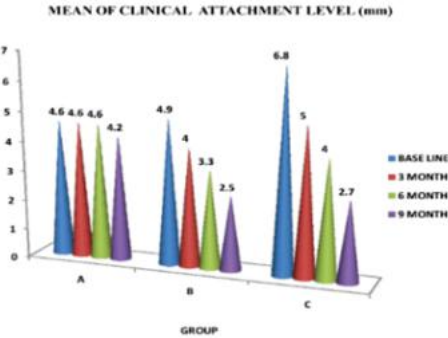
GROUP	BASE LINE		3 MONTH		6 MONTH		9 MONTH	
	MEAN	SD	MEAN	SD	MEAN	SD	MEAN	SD
A	5.3	2.1	5.3	2.1	5.3	2.1	4.9	1.81
B	5.6	1.36	4.6	1.36	3.9	1.45	3.1	1.45
C	7.1	1.81	5.4	1.56	4.3	1.42	3.0	1.34
F VALUE	2.63(NS)		0.59(NS)		1.64(NS)		4.29(S)	



Graph 3

Table 4 Mean and Standard Deviation (sd) of Clinical Attachment Level (cal) and Comparison Between the Groups

GROUP	BASE LINE		3 MONTH		6 MONTH		9 MONTH	
	MEAN	SD	MEAN	SD	MEAN	SD	MEAN	SD
A	4.6	2.06	4.6	2.06	4.6	2.06	4.2	2.04
B	4.9	1.37	4.0	1.26	3.3	1.35	2.5	1.28
C	6.8	1.66	5.0	1.41	4.0	1.34	2.7	1.35
	4.32(S)		0.87(NS)		1.46(NS)		3.06(NS)	

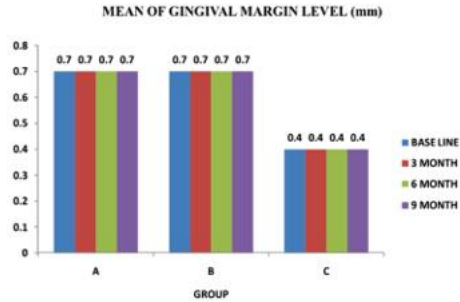


Graph 4

Table 5 Mean and Standard Deviation (sd) of Gingival Margin Level (gml) and Comparison Between the Groups.

GROUP	BASE LINE		3 MONTH		6 MONTH		9 MONTH	
	MEAN	SD	MEAN	SD	MEAN	SD	MEAN	SD
I	0.7	0.46	0.7	0.46	0.7	0.46	0.7	0.46

II	0.7	0.46	0.7	0.46	0.7	0.46	0.7	0.46
III	0.4	0.8	0.4	0.8	0.4	0.8	0.4	0.8
F VALUE	0.76(NS)		0.76(NS)		0.76(NS)		0.76(NS)	



Graph 5

Table 5 (a) Mean and Standard Deviation (sd) of Changes in the Clinical Parameters (with respect to base line)

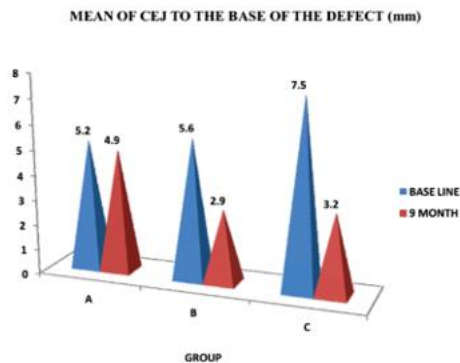
PARAMETER	GROUP	3 MONTH		6 MONTH		9 MONTH	
		MEAN	SD	MEAN	SD	MEAN	SD
MEAN PPD CHANGE (MM)	A	0	0	0.9	0.3	1.8	0.74
	B	0	0	1.6	0.67	2.8	0.74
	C	0.4	0.49	2.4	1.02	4.1	0.83
MEAN CAL (MM)	A	0	0	1	0	1.7	0.64
	B	0	0	1.7	0.46	2.8	0.65
	C	0.4	0.49	2.5	0.81	4.1	0.83
MEAN GML CHANGE (MM)	A	0	0	0	0	0	0
	B	0	0	0	0	0	0
	C	0	0	0	0	0	0

Table 5 (b) Comparison of Changes in the Clinical Parameters (with respect to base line) Between the Groups.

PARAMETER	GROUP	T VALUE		
		3 MONTH	6 MONTH	9 MONTH
MEAN PPD CHANGE (MM)	GROUP A AND GROUP B	9.00(S)	7.24(S)	5.30(S)
	GROUP B AND GROUP C	3.35(S)	3.60(S)	3.88(S)
	GROUP A AND GROUP C	0.13(NS)	0.09(NS)	0.06(NS)
MEAN CAL GAIN (MM)	GROUP A AND GROUP B	-	11.13(S)	6.68(S)
	GROUP B AND GROUP C	3.28(S)	3.76(S)	4.15(S)
	GROUP A AND GROUP C	-	0.08(NS)	0.06(NS)
MEAN GML CHANGE (MM)	GROUP A AND GROUP B	-	-	-
	GROUP B AND GROUP C	-	-	-
	GROUP A AND GROUP C	-	-	-

Table 6 Mean and Standard Deviation (sd) of CEJ to the Base of the Defect Comparison Between Periods and Between the Groups

GROUP	BASE LINE		9 MONTH		T VALUE
	MEAN	SD	MEAN	SD	
A	5.2	1.99	4.9	2.02	0.31(NS)
B	5.8	1.17	3.6	0.67	4.92(S)
C	7.5	1.43	3.1	1.14	7.22(S)
F VALUE	5.19(S)		4.66(S)		

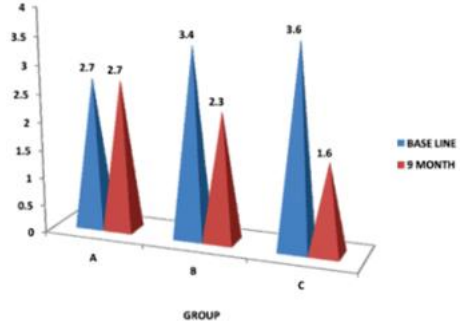


Graph 6

Table 7 Mean and Standard Deviation (sd) of CEJ to the Alveolar Crest, Comparison Between Periods and Between the Groups.

GROUP	BASE LINE		9 MONTH		T VALUE
	MEAN	SD	MEAN	SD	
A	2.7	0.64	2.7	0.64	0(NS)
B	3.4	0.66	2.4	0.66	3.20(S)
C	3.6	0.92	1.6	0.92	4.63(S)
F VALUE	3.57(S)		5.04(S)		

MEAN OF CEJ TO THE ALVEOLAR CREST (mm)

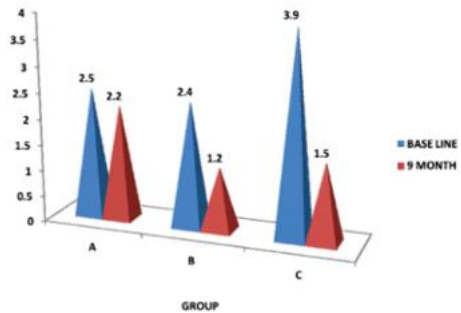


Graph 7

Table 8 Mean and Standard Deviation (sd) of Intra Bony Defect (ibd) Depth Comparison Between Periods and Between the Groups

GROUP	BASE LINE		9 MONTH		T VALUE
	MEAN	SD	MEAN	SD	
A	2.5	1.43	2.2	1.47	0.44(NS)
B	2.4	0.66	1.2	0.4	4.65(S)
C	3.9	1.14	1.5	0.81	4.44(S)
F VALUE	5.02(S)		2.39(S)		

MEAN OF INTRA BONY DEFECT DEPTH (mm)

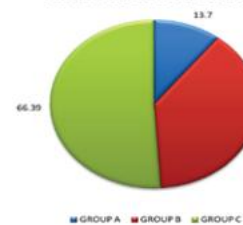


Graph 8

Table 9 Mean and Standard Deviation (sd) of Changes in the Radiographic Parameters (with respect to base line)

PARAMETER	GROUP	9 MONTH	
		MEAN	SD
MEAN IBD DEPTH REDUCTION	A	0.3	0.46
	B	1.2	0.4
	C	3.0	1.79
PERCENTAGE OF BONE DEFECT FILL	A	13.7	19.73
	B	50.0	7.45
	C	66.39	17.28

PERCENTAGE OF BONE DEFECT FILL



Graph 9

Table 10 Comparison of Changes in the Radiographic Parameters (with respect to base line) Between the Groups

PARAMETER	GROUP	T VALUE
MEAN IBD DEPTH REDUCTION	GROUP A AND GROUP B	4.44(S)
	GROUP B AND GROUP C	4.38(S)
	GROUP A AND GROUP C	2.94(S)
PERCENTAGE OF BONE DEFECT FILL	GROUP A AND GROUP B	5.06(S)
	GROUP B AND GROUP C	6.03(S)
	GROUP A AND GROUP C	2.61(S)

Statistics

- The paired t – test was applied to assess the statistical significance between time point within each group clinical and radiographic parameters.
- ONE WAY ANOVA test was applied to assess the statistical significance between time points between the groups for clinical and radiographic parameters.

DISCUSSION

Regeneration has been defined as the reproduction or reconstitution of a lost or injured part to restore the architecture and function of the periodontium (American Academy of Periodontology 1992).²

New attachment has been defined as the union of connective tissue or epithelium with a root surface that has been deprived of its original attachment apparatus (Glossary of Periodontal Terms 2001).²⁰

At present, one of the most widely used periodontal regenerative modalities is osseous graft therapy. The use of bone grafts for reconstructing osseous defects produced by periodontal disease dates back to Hegedus in 1923 and was reviewed by Nabers and O'Leary in 1965. Since that time, a number of techniques and materials have been used for regeneration.²¹

Platelet Rich Fibrin (PRF) described by Choukroun et al is a second generation platelet concentrate which contains platelets and growth factors in the form of fibrin membranes prepared from the patients own blood free of any anticoagulant or other artificial biochemical modifications. In vitro study²⁵ demonstrated that the viable platelets in PRF released six growth factors like PDGF, VEGF, TGF, IGF, EGF and bFGF in about the same concentration for the 7 – day duration. Prolong release of these growth factors, would be expected that PRF growth factors, would be expected that PRF treatment of an intrabony defect (IBD) may result in enhanced wound healing and periodontal regeneration compared with those sites treated with conventional flap debridement.

Clinical interpretation: Plaque index - Mean plaque index at 9 months was 0.56(0.26) in group A, 0.72(0.41) in group B and 0.57(0.25) in group C. A statistically significant reduction in the plaque index was observed in all the 3 groups from baseline to 9 months postoperatively.

Changes in Pocket Probing Depth (PPD) - Mean reduction in PPD was 0.8(0.28) mm in group A, 2.8(0.74) mm in group B and 4.1(0.83)mm in group C, which was similar to results obtained in a clinical study done by Kazuhiro et al²³ and AR Pradeep et al.²⁴

Changes in Clinical Attachment Level (CAL) - Mean gain in CAL was 1.7(0.64)mm in group A, 2.8 (0.65) mm in group B and 4.1(0.83) mm in group C, which was similar to the findings of Kazuhiro et al²³ and Pradeep et al.²⁴

Changes in Gingival Margin Level (GML) - Mean changes in GML was 0 for all the group.

An important clinical outcome of a periodontal reduction in PPD and CAL gain were found in all three groups when compared with baseline and 9 months.

Radiographic interpretation: Mean Intra Bony Defect (IBD) depth reduction - Mean IBD depth reduction was 0.3(0.46) mm in group A; 1.2 (0.4) mm in group B and 3.0(1.79) mm in group C. other studies by James et al²⁵, Corsair et al²⁶ and Pradeep et al²⁴ was similar to our results. Percentage of Bone Defect Fill - The percentage of bone defect fill was 13.7(19.73)% in group A, 50.0(7.45)% in group B and 66.39(17.28)% in group C. Our result was 66.39(17.28)% indicating more percentage of bone fill compared to other studies by Pradeep et al²⁴, James et al²⁵,

Corsair et al²⁶, Pradeep et al²⁷, Pradeep et al.²¹

It is prudent to introduce PRF as a graft material for the treatment of intrabony defects in a country like INDIA, where decisions still resolve around the economical viability of the treatment options. Simple way of preparation without the use of anticoagulants, inexpensive technique, less risk of disease transmission to the patient and more regenerative potential makes PRF as a demanding graft material to be used by developing countries like INDIA.

PRF is superior to other platelet concentrate like PRP due to its ease and inexpensive method of preparation and also it does not need any addition of exogenous compounds like bovine thrombin and calcium chloride. It is advantageous than autogenous graft also because an autograft requires a second surgical site and procedure.

In the current study the primary outcome was bone defect fill with the help of radiographic grid, as radiographic evaluation is a noninvasive examination for bony defects repair. Measuring the grid lines help in accurately measuring the defect as the distance between the 2 grid lines on the radiograph is 1mm, even if the images is foreshortened or elongated. Also, bone fill data derived from surgical re-entry are important to substantiate routine postoperative measurement data obtained radiographically.

In the present study grafting with (PRF) platelet rich fibrin showed more percentage of bone defect fill compared to grafting with an alloplastic (resorbable hydroxyapatite) and open flap debridement (OFD). This implies that platelet rich fibrin can be used as a sole grafting material in the treatment of intra bony defect. However, long-term, randomized, controlled clinical trials with large sample size are needed to prove its clinical and radiographic efficacy in periodontal regeneration.

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REFERENCES

- Dieter D Bosshardt, Anton Sculean. Does periodontal tissue regeneration really work? Periodontol 2000 2009;51:208–219.
- American Academy of Periodontology. Glossary of periodontal terms, 3rd edn. Chicago: American Academy of Periodontology 1992.
- Sander L, Karring T. Healing of periodontal lesions in monkeys following the guided tissue regeneration procedure. A histological study. J Clin Periodontol 1995;22:332–337.
- Pradeep AR, Shetty SK, Garg G, Pai S. Clinical effectiveness of autologous platelet-rich plasma and Peptide-enhanced bone graft in the treatment of intrabony defects. J Periodontol 2009;80:62-71.
- Döri F, Kovacs V, Arweiler NB, et al. Effect of platelet rich plasma on the healing of intrabony defects treated with an anorganic bovine bone mineral: a pilot study. J Periodontol 2009;80:1599-605.
- Siciliano VI, Andreuccetti G, Siciliano AI, Blasi A, Sculean A, Salvi GE. Clinical outcomes after treatment of non-contained intrabony defects with enamel matrix derivative or guided tissue regeneration: a 12-month randomized controlled clinical trial. J Periodontol 2011;82:62-71.
- Wu SY, Chen YT, Chen CW, et al. Comparison of clinical outcomes following guided tissue regeneration treatment with a polylactic acid barrier or a collagen membrane. Int J Periodontics Restorative Dent. 2010;30:173-179.
- Chen FM, Zhang J, Zhang M, An Y, Chen F, Wu ZF. A review on endogenous regenerative technology in periodontal regenerative medicine. Biomaterials 2010;31(31):7892-927.
- Orsini M, Orsini G, Benlloch D, Aranda JJ, Sanz M. Long-term clinical results on the use of bonereplacement grafts in the treatment of intrabony periodontal defects. Comparison of the use of autogenous bone graft plus calcium sulfate to autogenous bone graft covered with a bioabsorbable membrane. J Periodontol 2008;79:1630.
- Cochran DL, Jones A, Heijl L, Mellonig JT, Schoolfield J, King GN. Periodontal regeneration with a combination of enamel matrix proteins and autogenous bone grafting. J Periodontol. 2003;74:1269-1278.
- Nevins M, Hanratty J, Lynch SE. Clinical results using recombinant human platelet-derived growth factor and mineralized freeze-dried bone allograft in periodontal defects. Int J Periodontics Restorative Dent 2007;27:421–427.
- Hoidal MJ, Grimald BA, Mills MP, Schoolfield JD, Mellonig JT, Mealey BL. Clinical evaluation of demineralized freeze-dried bone allograft with and without enamel matrix derivative for the treatment of periodontal osseous defects in humans. J Periodontol 2008;79:2273-80.
- Kenney EB, Lekovic V, Han T, Carranza Jr FA, Dimitrijevic B. The use of a porous hydroxyapatite implant in periodontal defects I. Clinical results after six months. J Periodontol 1985;56:82-88.
- Meffert RM, Thomas JR, Hamilton KM, Brownstein CN. Hydroxylapatite as an alloplastic graft in the treatment of human periodontal osseous defects. J Periodontol 1985;56:63-73.
- Stahl SS, Fromm SJ. Histologic and clinical responses to porous hydroxylapatite implants in human periodontal defects. Three to twelve months postimplantation. J Periodontol 1987;58:689-695.
- Bowen J, Melling J, Gray J, Towle H. Comparison of decalcified freeze dried bone allograft and porous hydroxyapatite in human periodontal osseous defects. J periodontol 1989;60:647-653.

- 17) Choukroun J, Adda F, Schoeffler C, Vervelle A. An opportunity in perio-implantology: the PRF. *Implantodontie* 2001;42:55–62.
- 18) Anitua E, Andia I, Ardanza B, Nurden P, Nurden AT. Autologous platelets as a source of proteins for healing and tissue regeneration. *J Thromb Haemost.* 2004;91:4–15.
- 19) Veena Kalburgi, Shivaraj Warad, Haziel Diana Jenifer, Nipun Ashok, Vijayalaxmi Kokatnur, Application of Platelet Rich Fibrin and Osseomold Bone Graft in Different Intrabony Defects –2 Case Reports, *Int journal of clinical dental science*, march 2012; 3(1).
- 20) American Academy of Periodontology. Glossary of periodontal terms, 4th edn. Chicago: American Academy of Periodontology 2001.
- 21) A.R Pradeep, Nishanth S. Rao, Esha Agarwal, pavan Bajaj, Minal Kumari and savitha B. Naik. Comparative evaluation of autologous platelet rich fibrin and platelet rich plasma in the treatment of 3-wall intrabony defects in chronic periodontitis: A randomized controlled clinical trial. *J Periodontol* 2012;83:1499-1507.
- 22) Carroll, R. J., Amoczkzy, S. P., Graham, S. & O'Connell, S. M. (2005) Characterization of Autologous Growth Factors in Cascade Platelet Rich Fibrin Matrix (PRFM). Edison, NJ: Musculoskeletal Transplant Foundation.
- 23) Kazuhiro Okuda, Hideaki Tai, Kiyoshi Tanabe, Hironobu Suzuki, Tadashi Sato, Tomoyaki Kawai, Yoshinori Saito, Larry F Wolf. Platelet rich plasma combined with a porous hydroxyapatite graft for the treatment of intrabony periodontal defects in humans: a comparative controlled clinical study. *J Periodontol* 2005;76(6):890-898.
- 24) A.R Pradeep, Pavan Bajaj, Nishant S. Rao, Esha Agarwal, Savitha B.Naik. Platelet rich fibrin combined with a porous hydroxyapatite graft for the treatment of three-wall intrabony defects in chronic periodontitis: A randomized controlled clinical trial. *J Periodontol* 2017 Dec;88(12):1288-1296.
- 25) Bowen JA, Mellonig JT, Gray JL, Towle HT. Comparison of decalcified freeze dried bone allograft and porous particulate hydroxyapatite in human periodontal osseous defects. *J Periodontol* 1989;60:647-653.
- 26) Corsair A. A clinical evaluation of resorbable hydroxyapatite for the repair of human intraosseous defects. *J Oral Implantol* 1990;16(2):125-128.
- 27) Lee HJ, Choi BH, Junq JH, Zhu ST, Lee SH, Huh JY, You TM, Li J. Maxillary sinus floor augmentation using autogenous bone grafts and platelet – enriched fibrin glue with simultaneous implant placement. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2007;103(3):329-33.