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## Treatment of localized gingival recession using Vestibular Incision Sub periosteal Tunnel Access Technique in conjunction with albumin Platelet Rich fibrin

### Abstract:

#### Background:

The current randomized control study(RCT)was carried out to compare the efficacy of albumin plateletrich fibrin (Alb-PRF) also referred to as(extended plateletrich fibrin (e-PRF))membranecompared to the de-epithelized free gingival graft (DFGG) in combination withvestibular incision subperiosteal tunnel access technique (VISTA) for treatment of localized gingival recession type 1 (RT1).

#### Methods:

Thirty patients aged above 18 were randomly allocated for localized gingival recession (GR) treatment into two main groups (VISTA and DFGG) or (VISTA and e-PRF). The primary outcome was to evaluate therecession depth (RD) improvement, while the secondary outcomes included the changes ofprobing depth (PD), clinical attachment level (CAL),plaque index (PI) and gingival index (GI).All the study outcomes were evaluated3 and 6 months postoperatively.

#### Results:

Both groups achieved highly significant clinical improvements from baseline to 6 months ( $P < 0.001$ ). While VISTA+DFGG demonstrated a statistically superior recession depth reduction at 3 months ( $P = 0.028$ ), both groups yielded statistically comparable results at the final 6-month follow-up for recession depth ( $0.40 \pm 0.27$  mm vs  $0.49 \pm 0.37$  mm;  $P = 0.390$ ) and clinical attachment level ( $1.59 \pm 0.55$  mm vs  $1.62 \pm 0.52$  mm;  $P = 0.749$ ). No significant inter-group differences were observed for probing depth, plaque index, and gingival index at any evaluation period ( $P > 0.05$ ).

#### Conclusion:

It was concluded thatDFGGremains the gold standard for root coverage, although e-PRF membrane combined with VISTA shows promising, less invasive potential for gingival augmentation and regeneration.

**Keywords:** Gingival recession, extended plateletrich fibrin, Albumin plateletrich fibrin,Vestibular Incision Subperiosteal Tunnel Access technique, de-epithelialized Free Gingival Graft

## 32 Introduction

33 Gingival recession (GR) is characterized by the displacement of the marginal gingival tissues  
34 in an apical direction relative to the cemento-enamel junction. Although this condition does not  
35 necessarily lead to tooth loss, it is often viewed as a significant aesthetic issue for patients. It is  
36 frequently linked to root surface exposure, which can result in dentin hypersensitivity and the  
37 formation of both carious and non-carious cervical lesions. In advanced stages, recession may cause  
38 substantial root abrasion and, in rare circumstances, may eventually contribute to tooth loss. (1) The  
39 causes of GR are diverse, ranging from aggressive tooth brushing and poor orthodontic management  
40 to malpositioned teeth, defective restorations, plaque-related inflammation, a thin biotype, and  
41 periodontal disease. (2)

42 To standardize diagnosis, several classifications for gingival recession have been  
43 proposed over the years. Currently, the Cairo classification is the most frequently utilized system  
44 because it includes interproximal attachment loss, offering improved diagnostic accuracy and  
45 prognostic value. (3) Cairo categorized gingival recession into three types, Recession type 1 (RT1) is  
46 defined by the absence of interproximal CAL, whereas in recession type 2 (RT2), interproximal CAL  
47 is equal to or less than buccal CAL. In recession type 3 (RT3), interproximal CAL exceeds buccal  
48 CAL. (1)

49 Various surgical techniques are available for GR management. The coronally advanced flap  
50 (CAF) is a widely utilized method, offering predictable results for GR. CAF is frequently combined  
51 with various grafting materials, including connective tissue graft (CTG), enamel matrix derivatives  
52 (EMD), platelet-rich fibrin (PRF), and xenogeneic collagen matrix (XCM). (4) Although the CAF  
53 remains the gold standard for GR treatment, advancements in surgical techniques have introduced  
54 new approaches for more predictable treatment such as the double papilla technique, tunnel  
55 technique (TUN) and its modifications. (5)

56 A significant advancement is the VISTA (vestibular incision subperiosteal tunnel access)  
57 method, a minimally invasive surgical procedure designed to protect blood supply and reduce tissue  
58 trauma, resulting in consistent root coverage and excellent aesthetics. Clinical study by Zadeh has  
59 demonstrated promising results with VISTA, establishing it as a valuable option in contemporary  
60 periodontal plastic surgery. (6) Although traditional connective tissue grafts are successful, they  
61 require a secondary donor site for harvesting, which often increases patient pain and surgical  
62 morbidity. (7)

63 To overcome the issues of traditional grafting, several alternative materials have been  
64 developed. However, substitutes like allografts and xenografts, while avoiding palatal harvesting,  
65 introduce significant drawbacks such as substantial financial cost. (8) In regenerative therapy,  
66 autologous platelet derivatives like PRP and PRF are frequently employed because they contain high  
67 concentrations of platelets, white blood cells, and multiple growth factors. (9) Even though PRF  
68 provides a more sustained release of growth factors than PRP, both materials usually resorb quickly  
69 (within 2-3 weeks), which can restrict their performance when used as sole regenerative barriers. (10)

70 Recent advancements have led to extended-PRF generation (e-PRF or Alb-PRF), which  
71 might significantly prolongs resorption times (4–6 months), thereby preserving longer regenerative  
72 potential and enhancing biological activity. Alb-PRF membranes show considerable promise as  
73 viable alternatives to collagen membranes in guided bone regeneration (GBR), with documented  
74 clinical applications including extraction site coverage, lateral sinus lifts, and treatment of multiple  
75 GR.(10)

76 Nevertheless, a demanding need still exists for optimal and regenerative approaches. While  
77 the VISTA technique offers excellent outcomes and Alb-PRF (e-PRF) shows superior biological  
78 properties, their combined synergistic effect for GR treatment is largely unexplored in previous  
79 literatures. Thus this study aimed to address this significant research gap by clinically evaluating and  
80 comparing the effectiveness of e-PRF membrane versus DFGG when combined with the VISTA  
81 technique for localized GR treatment.

82

## 83 **Materials and Methods**

### 84 **Study Design**

85 This clinical comparative study was designed as two-arm parallel randomized control trial  
86 (RCT). A total of thirty participants with localized gingival recession classified as Cairo  
87 recession type 1 (RT 1).(11) Participants were divided into study group VISTA + e-PRF and control  
88 group VISTA +DFGG.

89 Following a comprehensive discussion of all procedures, participants provided signed  
90 informed consent. The study protocol was approved by the ethics committee (Ref: A0801024OM)  
91 and registered on ClinicalTrials.gov (ID: NCT07079293).

### 92 **Eligibility criteria**

93 Inclusion criteria were patients with Cairo recession type 1 (RT1) (i.e. GR without  
94 interproximal attachment loss). Participants were systemically healthy, non-smokers with good oral  
95 hygiene and no history of mucogingival surgery. Exclusion criteria included interproximal bone loss,  
96 pregnancy or lactation, immunocompromised status, fixed prostheses, cervical caries or restorations  
97 at the treatment site, and poor oral hygiene.

### 98 **Sample size calculation**

99 The sample size was calculated using G\*Power 3.1.9.4, based on root coverage data from a  
100 previous study.(12) With an effect size of 0.95, an alpha of 0.05, and 85% power, a minimum of 13  
101 participants per group was required to compensate for potential dropouts.

102

103 **Intervention:**

104 **Pre-operative phase:**

105 Six weeks prior to surgery, all participants received professional nonsurgical periodontal  
106 therapy and individualized oral hygiene instructions.

107 **Surgical procedures:**

108 The VISTA technique was performed as described by Zadeh.(6)Following the administration  
109 of local anesthesia (4% articaine), a 15c surgical blade (Smi, Belgium) was used to perform a  
110 vestibular access incision.Specialized tunneling instruments (Helmut Zepf, Germany) were then  
111 utilized to elevate a subperiosteal tunnel.

112 For participants in the control group (figure 1), a free gingival graft (FGG) was obtained from  
113 the palatal donor site under local anesthesia.The graft was de-epithelialized extra-orally using a 15c  
114 blade to produce a DFGG with a thickness ranging from 1 to 1.5 mm.(13)The DFGG was then  
115 placed through the tunnel and secured with a horizontal suture. To prevent apical relapse, 5.0 Vicryl  
116 sutures (Ethicon, US) were anchored to each tooth by acid-etching the facial enamel, washing, and  
117 drying, then placing flowable composite (Estelite universal flow composite, Tokuyama America)  
118 over the knot. Finally, the access incision was closed with 5.0 Vicryl sutures.(6)

119 For the e-PRF group(10) (figure 2), 10 mL of whole blood was collected in additive-free tubes  
120 (Vacutest, Italy) and centrifuged at 700g for 8 minutes (DLAP DM0424, China).The resulting platelet-  
121 poor plasma (PPP) uppermost layer was collected and heated at 75°C for 10 minutes, then cooled for 2  
122 minutes to create an injectable albumin gel.

123 Concurrently, the liquid platelet-rich fibrin (liquid-PRF) and buffy coat were collected. This  
124 liquid-PRF was then mixed with the prepared albumin gel using a luer-lock connector. After  
125 approximately 5 minutes for fibrin polymerization, the resulting e-PRF membrane was obtained(14),  
126 placed through the tunnel, adapted, and secured with sutures.

127 **Post-operative phase:**

128 After the procedure, participants were advised to refrain from mechanical brushing at the  
129 surgical area for 14 days and to follow a soft diet. A 0.1% chlorhexidine rinse (OROVEX-H®,  
130 Egypt) was prescribed thrice-daily. Antibiotics (Augmentin® 1000 mg, UK) and analgesics  
131 (Brufen® 600 mg, Egypt) were administered twice daily for five days. At one week, the site was  
132 professionally cleaned, and palatal sutures were removed (control group). Gingival recession site  
133 sutures were removed at 7-14 days. Follow-up visits were scheduled at 1, 3, and 6 months for  
134 outcome assessment and oral hygiene reinforcement.

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136

137 **Clinical assessments**

138 Periodontal outcomes were assessed at baseline, three and six months post-surgery using a  
139 UNC15 probe (Sedradent, Egypt). The primary outcome was the recession depth  
140 reduction(15)whileSecondary outcomes included changes in probing depth(16), clinical attachment  
141 level(17), plaque index(18) and gingival index.(19)

142 **StatisticalAnalysis:**

143 Data were analyzed using IBM SPSS v.20.0. Categorical data were presented as numbers and  
144 percentages, while quantitative data were described by mean, standard deviation, median, and IQR.  
145 The Shapiro-Wilk test verified data normality. Statistical significance was set at the 5% level ( $p <$   
146  $0.05$ )

147

148 **Results**

149 Thirty participants, all over 18 years of age and suffering from Cairo RT1, were randomized  
150 to the DFGG (n=15) and e-PRF (n=15) groups. The DFGG group (Group I) and the e-PRF group  
151 (Group II) were both treated using the VISTA technique in conjunction with their respective  
152 materials.

153 The demographic data presented in Table 1 indicates that there were no statistically  
154 significant differences between the control group and the study group in terms of either sex  
155 distribution or mean age. The p-value;  $p > 0.05$  for both age and sex, indicating that the two groups  
156 were comparable in these baseline characteristics.

157 Table 2 reveals the primary andsecondary outcomes. Regarding the mean RD, the baseline  
158 was comparable between the VISTA+DFGG ( $2.63 \pm 0.77$  mm) and VISTA+e-PRF ( $2.54 \pm 0.38$ mm)  
159 groups, with no statistically significant difference ( $P=0.609$ ). Following the surgical intervention,  
160 both treatment modalities achieved a highly significant reduction in RD over the 6-month study  
161 period ( $P0 < 0.001$ ). Notably, the VISTA+DFGG group demonstrated superior clinical performance  
162 exhibiting significantly lower RD values at the 3-month interval ( $0.50 \pm 0.31$  mm vs.  $0.74 \pm 0.31$   
163 mm;  $P=0.028$ ) compared to the VISTA+e-PRF group. However, at the 6-month interval, there was no  
164 statistically significant difference between the two groups ( $0.40 \pm 0.27$  mm vs.  $0.49 \pm 0.37$  mm;  
165  $P=0.390$ ).

166 In terms of CAL, both groups showed highly significant longitudinal improvements from  
167 baseline to the final follow-up ( $P0 < 0.001$ ). The baseline CAL was comparable between the  
168 VISTA+DFGG ( $4.04 \pm 0.83$  mm) and VISTA+e-PRF ( $3.90 \pm 0.40$  mm) groups, with no statistically  
169 significant difference ( $P=0.462$ ). By 6 months, both groups achieved a gain in attachment, resulting  
170 in a mean CAL of  $1.59 \pm 0.55$  mm in the VISTA+DFGG group compared to  $1.62 \pm 0.52$  mm in the  
171 VISTA+e-PRF group, with no statistically significant difference between them ( $P=0.749$ ).  
172 Conversely, PD remained relatively stable throughout the study, with no statistically significant

173 differences observed between the two groups at any time point ( $P > 0.05$ ) or within either group over  
174 the three periods ( $P > 0.05$ ).

175 Regarding the PI and GI, inter-group comparisons revealed no statistically significant  
176 differences for either index at baseline, 3 months, or 6 months ( $P > 0.05$ ), indicating that both  
177 surgical protocols resulted in comparable outcomes for these parameters throughout the follow-up  
178 phases. However, longitudinal changes within the groups varied. For the PI, neither group exhibited  
179 statistically significant changes over the 6-month duration ( $P > 0.05$ ). For the GI, the  
180 VISTA+DFGG group showed no significant longitudinal changes ( $P > 0.05$ ), whereas the  
181 VISTA+e-PRF group exhibited a statistically significant change over time ( $P = 0.003$ ).

182

## 183 Discussion

184 This randomized controlled clinical trial aimed to evaluate the efficacy of VISTA  
185 combined with DFGG compared to VISTA with e-PRF for the treatment of localized gingival  
186 recession.(6) Our primary findings demonstrated that both modalities achieved highly significant  
187 longitudinal improvements in Recession Depth (RD) and Clinical Attachment Level (CAL) from  
188 baseline to 6 months ( $P < 0.001$ ). While the VISTA+DFGG group exhibited a statistically  
189 superior reduction in RD at the 3-month interval ( $P=0.028$ ), this difference was transient. By the  
190 final 6-month follow-up, both groups yielded statistically comparable results for RD ( $P=0.390$ )  
191 and CAL ( $P=0.749$ ). Furthermore, Probing Depth (PD), Plaque Index (PI), and Gingival Index  
192 (GI) showed no significant differences between the two treatment groups at any observation  
193 period.

194 Autologous grafts, such as connective tissue grafts (CTG) and DFGG, have traditionally  
195 been considered the gold standard for gingival recession coverage due to their robust clinical  
196 outcomes.(20) This is supported by recent studies, such as the randomized clinical trial by Abu-  
197 Ta'a (2023), which compared Coronally Advanced Flap (CAF) with either Advanced Platelet-  
198 Rich Fibrin (A-PRF) or CTG and concluded that CTG yielded significantly better reductions in  
199 recession depth and width.(21) Similarly, a systematic review by Balčiūnaitė et al. (2020)  
200 acknowledged that while standard PRF is an effective and less invasive alternative, autologous  
201 grafts remain the established gold standard for achieving predictable root coverage.(20)

202 While these referenced studies utilized different generations of platelet concentrates (such  
203 as A-PRF and standard L-PRF), they were specifically selected for our comparative analysis  
204 because they share the same fundamental biological rationale as e-PRF.(22) All these PRF  
205 concentrates are autologous biomaterials designed to entrap platelets and leukocytes within a  
206 fibrin matrix, thereby serving as reservoirs for the sustained release of essential growth factors,  
207 including Platelet-Derived Growth Factor (PDGF), Transforming Growth Factor-beta (TGF- $\beta$ ),  
208 and Vascular Endothelial Growth Factor (VEGF).(23) Our 3-month results partially align with the  
209 historical findings of these studies, as the DFGG group initially provided significantly lower RD  
210 values. This early superiority can be attributed to the immediate structural coverage provided

211 directly by the transferred autogenous donor tissue.

212 However, the comparable outcomes observed at 6 months in our study challenge the  
213 notion that autologous grafts are always statistically superior in the long term, highlighting the  
214 unique physical advantage of extended PRF (e-PRF). Although standard platelet concentrates  
215 (like PRF and A-PRF) successfully secrete growth factors, their primary limitation has  
216 historically been their rapid resorption rate, typically dissolving within a 2- to 3-week timeframe.  
217 This rapid degradation often limits their ability to act as a long-term barrier. In contrast, the e-  
218 PRF used in our study overcomes this limitation by incorporating a heated platelet-poor plasma  
219 (PPP) layer that denatures albumin, restructuring it into a denser tridimensional matrix. This  
220 specific modification dramatically extends the biodegradation period of the membrane up to 4 to  
221 6 months.(10)

222 Consequently, this prolonged stability allows for the slow, gradual release of the  
223 entrapped growth factors (PDGF, TGF- $\beta$ , VEGF) over an extended period, stimulating sustained  
224 cellular migration, fibroblast proliferation, and continuous collagen synthesis.(10, 24) This  
225 extended regenerative window effectively explains the progressive clinical improvement  
226 observed in the VISTA+e-PRF group. While autologous grafts like DFGG remain the  
227 unparalleled gold standard for providing robust structural volume, the sustained biological  
228 stimulation from e-PRF significantly narrowed the initial clinical gap observed at 3 months.  
229 Ultimately, e-PRF yielded highly satisfactory and statistically comparable outcomes for both RD  
230 and CAL by the 6-month mark.

231 The surgical technique itself also played a critical role in facilitating these favorable  
232 outcomes. The VISTA approach provides minimally invasive access through a vestibular  
233 incision, enabling the elevation of a subperiosteal tunnel that preserves the anatomical integrity of  
234 the interdental papillae and minimizes tension on the gingival margin during coronal  
235 advancement.(6, 21) By combining VISTA with e-PRF, the surgical site benefits from both  
236 minimized surgical trauma and extended biological stimulation. Crucially, this combination  
237 eliminates the necessity for a secondary surgical donor site, addressing one of the main  
238 drawbacks of DFGG and CTG, which is postoperative patient morbidity, pain, and  
239 discomfort.(20)

240 Regarding the secondary clinical parameters, our results demonstrated stable Probing  
241 Depths (PD) throughout the study, with no significant longitudinal changes for either group. This  
242 is in agreement with existing literature, such as the study by Abu-Ta'a (2023), indicating that both  
243 autologous grafts and PRF-based recession therapies successfully maintain periodontal health  
244 without inducing pathological pocket formation.(21) Furthermore, the Plaque Index (PI) remained  
245 stable and exceptionally low across all observation periods for both groups ( $P > 0.05$ ), confirming  
246 that the minimally invasive VISTA protocol effectively preserves the anatomical integrity of the  
247 interdental papillae, thereby not creating plaque-retentive areas or hindering patients' oral hygiene  
248 practices.

249 Interestingly, while the Gingival Index (GI) showed no significant inter-group differences,

250 the VISTA+e-PRF group exhibited a statistically significant longitudinal increase from baseline  
251 to 6 months ( $P=0.003$ ). However, the final mean GI value ( $0.59 \pm 0.15$ ) remains strictly within  
252 the clinical parameters of healthy, non-inflamed gingival tissues. Biologically, this slight  
253 longitudinal shift in visual gingival scores should not be interpreted as pathological plaque-  
254 induced inflammation. Rather, recent evidence by Kargarpour et al. (2021) suggests that liquid  
255 PRF and heated albumin gels actually possess potent in vitro heat-sensitive anti-inflammatory  
256 activity.(25)Therefore, this minor visual change in gingival erythema is more likely attributed to  
257 the robust angiogenesis and hypervascularization stimulated by the sustained and prolonged  
258 release of Vascular Endothelial Growth Factor (VEGF) from the e-PRF matrix, which naturally  
259 enhances local blood perfusion during the extended tissue remodeling phase.(24)

260

## 261 **Conclusion**

262 The use of e-PRF in conjunction with the VISTA technique provides a highly effective  
263 and minimally invasive alternative to DFGG for the treatment of localized gingival recession. By  
264 utilizing modified heating protocols, e-PRF shares the powerful growth factor release mechanism  
265 of previous PRF concentrates but overcomes their rapid resorption limitations(10). While it may  
266 not entirely replicate the absolute volumetric gain of an autologous graft, e-PRF offers sustained  
267 regenerative benefits that yield reliable long-term clinical outcomes for recession reduction and  
268 attachment gain, while significantly enhancing patient comfort by eliminating donor-site  
269 morbidity.

270

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277

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## 280 **Ethics approval and consent to participate:**

281 Written consent had been taken from each patient before performing any steps and approved  
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283 **Competing interests:**

284 Non

285

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## Tables & Figures

382

383 **Table (1): Demographic characteristics of study populations**

Demographic data	VISTA+DFGG (n = 15)	VISTA+e-PRF (n = 15)	p
<b>Sex, N (%)</b>			
<b>Male</b>	<b>6 (40.0%)</b>	<b>(8) 53.3%</b>	<b>0.464</b>
<b>Female</b>	<b>9 (60.0%)</b>	<b>(7) 46.7%</b>	
<b>Age (years)</b>			
<b>Mean ± SD</b>	<b>32.80 ± 10.63</b>	<b>35.20 ± 12.47</b>	<b>0.575</b>
<b>Range</b>	<b>21 – 53</b>	<b>21 – 64</b>	

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385 **Data was expressed using Mean ± SD. SD: Standard deviation**

386 **P: P value for Comparison between the two studied groups**

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401 **Table (2): Comparison between the two studied groups according to primary and**  
 402 **secondary outcomes in each period**

Outcomes	Periods	VISTA+DFGG (n = 15)	VISTA+e-PRF (n = 15)	P
Recession Depth (RD)	Baseline	2.63 ± 0.77	2.54 ± 0.38	0.609
	After 3M	0.50 ± 0.31	0.74 ± 0.31	0.028*
	After 6M	0.40 ± 0.27	0.49 ± 0.37	0.390
	P0	<0.001*	<0.001*	
Probing Depth (PD)	Baseline	0.89 ± 0.24	0.97 ± 0.39	0.870
	After 3M	0.80 ± 0.23	0.89 ± 0.28	0.980
	After 6M	0.79 ± 0.22	0.87 ± 0.27	0.939
	P0	0.268	0.073	
Clinical Attachment Level (CAL)	Baseline	4.04 ± 0.83	3.90 ± 0.40	0.462
	After 3M	1.76 ± 0.70	1.95 ± 0.52	0.345
	After 6M	1.59 ± 0.55	1.62 ± 0.52	0.749
	P0	<0.001*	<0.001*	
Plaque Index (PI)	Baseline	0.17 ± 0.12	0.19 ± 0.11	0.695
	After 3M	0.21 ± 0.12	0.23 ± 0.11	0.597
	After 6M	0.24 ± 0.14	0.29 ± 0.15	0.270
	P0	0.186	0.055	
Gingival Index (GI)	Baseline	0.35 ± 0.26	0.36 ± 0.15	0.907
	After 3M	0.42 ± 0.30	0.46 ± 0.20	0.849
	After 6M	0.46 ± 0.39	0.59 ± 0.15	0.346
	P0	0.536	0.003*	

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404 Data was expressed using Mean ± SD. SD: Standard deviation

405 P: P value for Comparison between the two studied groups

406 P<sub>0</sub>: P value for comparison between the three studied periods in each group

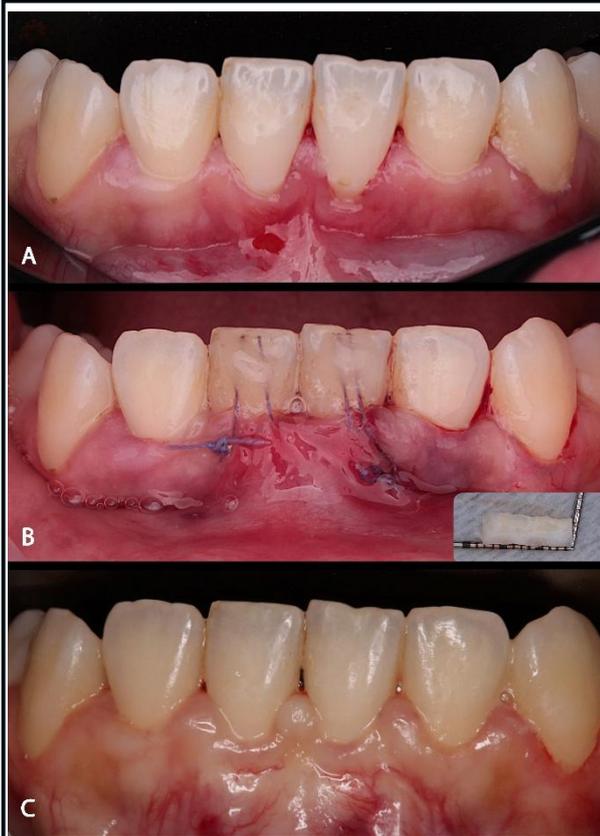
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**Figure 1:** show clinical view of the surgical procedures in the control group (VISTA + DFGG) (A) preoperative view of RT1 defect (B) immediately after surgery with DFGG in the corner of picture (C) six months follow up

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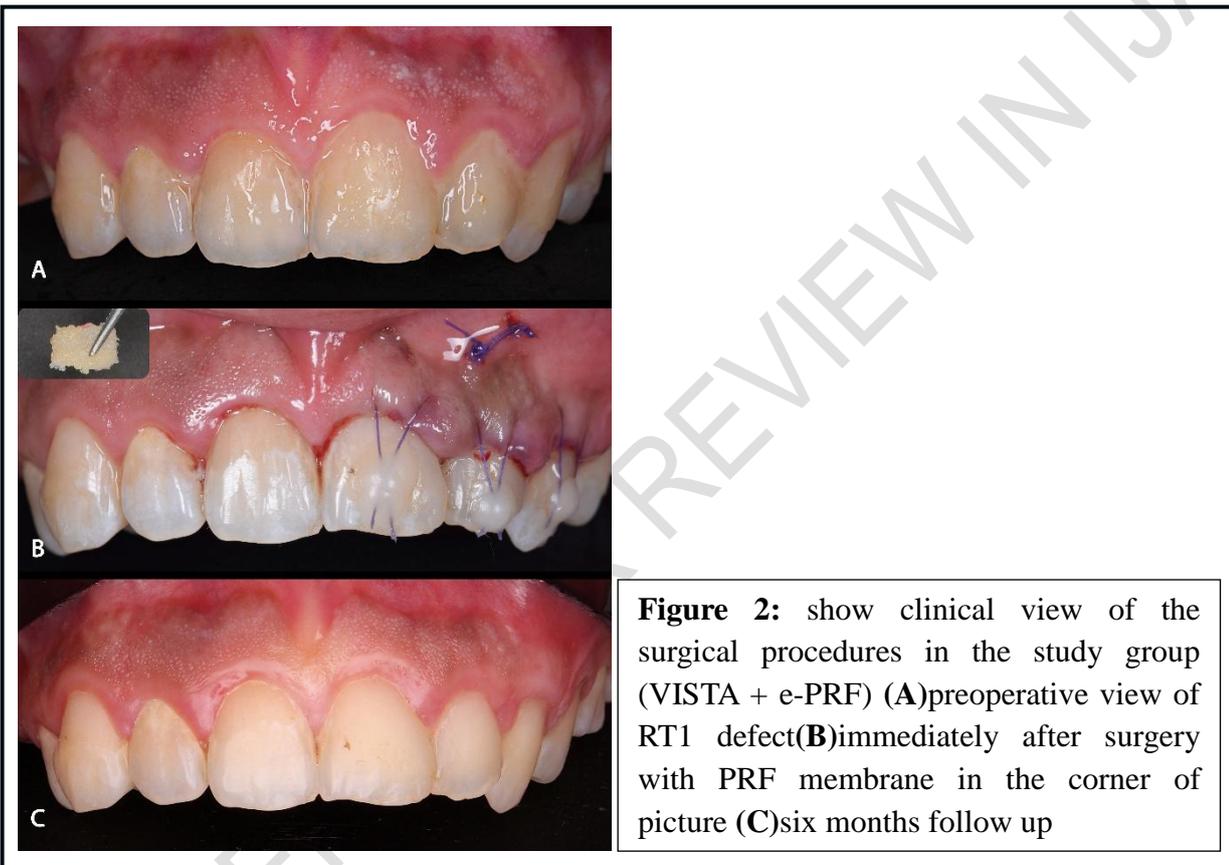
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