

Original Article

Gingival Unit Graft Versus Free Gingival Graft for Treatment of Gingival Recession: A Randomized Controlled Clinical Trial

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Abstract

Objectives: Gingival recession can lead to root exposure and discomfort for patients. There are various techniques for root coverage. The aim of this study was to compare the use of gingival unit graft (palatal graft including the marginal gingiva and papillae) with free gingival graft for treatment of localized gingival recession.

Materials and Methods: In this randomized controlled clinical trial, 18 bilateral localized recessions of Miller class I and II were treated in nine systemically healthy patients. Recessions were randomly treated with gingival unit graft in one side and conventional free gingival graft in the other side. Clinical parameters including clinical attachment level, keratinized tissue width, probing depth and vertical recession depth (VRD) were recorded at baseline and at one, three and six months after surgery. The healing index and patient satisfaction were also evaluated. One-way and two-way repeated measures ANOVA and paired t-test were used for statistical analyses.

Results: Both techniques caused significant improvement in clinical parameters. Gingival unit graft produced higher satisfaction esthetically ($P=0.050$, 0.024 and 0.024 , respectively at the three time points), higher healing index ($P<0.001$), higher root coverage percentage at one month after surgery (34.04% , $P=0.011$) and greater reduction of recession width three months after surgery ($P=0.007$) but the reduction in VRD at this side was not significantly greater.

Conclusions: Gingival unit graft might be an acceptable modality in Miller Class I/II recession defects. This technique may have advantages over free gingival graft such as significantly superior clinical and esthetic results.

Keywords: Gingival Recession; Transplantation, Autologous; Tooth Root; Transplants

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INTRODUCTION

Marginal gingival recession refers to the location of the gingival margin apical to the cemento-enamel junction, which results in exposure of root surfaces and loss of attached gingiva [1]. The prevalence of gingival recession varies between 20% and 100% among adults [2-6]. Gingival recession may increase tooth hypersensitivity [7] and cause pain, root caries, unesthetic appearance of the gums, periodontal attachment loss and tooth loss [8], and make oral hygiene and plaque control difficult [9]. These clinical problems can be resolved by root coverage treatments, including free gingival graft [10],

pedicle graft [11], connective tissue graft [12], coronally positioned flap [13] and guided tissue regeneration [14]. Although most of these methods result in significant clinical improvement [6,15], selection of the method of choice depends on many factors, including factors related to the defect, patient and technique. Therefore, treatment of gingival recession is still a challenge and researchers are working to find new more efficient surgical techniques for this purpose [16]. Although non-submerged palatal grafts have shown predictable results for root coverage [17], some changes can be made to improve the

clinical conditions of grafts including submerging the graft [18,19], "butt joint" adaptation of the graft to the adjacent tissues at the recipient site, root planing with the aim of reducing its prominence, and special suturing techniques to improve adaptation of the graft and its blood supply at the recipient site. The only donor site modification that can promote success in root coverage with non-submerged grafts is to increase graft thickness to maintain its viability on the avascular root surface [10,16,17,20,21]. Use of site-specific donor tissue may increase graft survival at the recipient site, which does not have optimal blood perfusion.

Allen and Cohen [22] used a palatal graft containing marginal gingiva as a free gingival graft without increasing its thickness. They used this technique based on the data in a case report [23]. Gingiva has a unique structure and characteristics [24,25]. The vascular plexus of the gingiva is rich of horizontal anastomoses, which perfuse the marginal zone but do not extend to the interproximal area. Marginal and interdental gingival tissues can be used to benefit from better blood perfusion of the donor and recipient sites and, therefore, improve the chances for graft survival [26]. The vascular characteristics of the graft are probably important for rapid anastomosis of the capillaries of the recipient site with the injured vessels of the graft [16].

Gingival unit graft should be harvested from an area, which is not esthetically important [26]. There are only a few case studies and just one clinical trial in this respect [22,26]. Kuru and Yildirim [16] compared the gingival unit graft and the conventional free gingival graft in patients with Miller class I/II gingival recession. Reduction in vertical recession and attachment and keratinized tissue gain were significantly higher in gingival unit group. Other human studies have shown that in clinically healthy gingiva, there are significantly different vascular distributions in marginal, attached and interdental gingiva [16]. The involvement of marginal

gingiva and papillary tissue in the graft can improve defect coverage, healing process and color adaptation. The aim of the present study was to compare the use of gingival unit graft with the conventional free gingival graft (palatal graft) for treatment of localized gingival recession.

MATERIALS AND METHODS

In this randomized controlled clinical trial with split-mouth design, 18 localized recessions of Miller class I and II were treated in nine systemically healthy patients. Bilateral recessions with a vertical depth of ≥ 2 mm on the buccal aspects of mandibular premolars and incisors were selected. The exclusion criteria were (I) smoking, (II) pregnancy, (III) root surface restorations, (IV) endodontic treatment and (V) poor oral hygiene (O'Leary plaque score $>20\%$). All details of the two treatment modalities were explained to the participants and written informed consent was obtained from them.

The study protocol was approved by the Ethical Committee of Babol University of Medical Sciences and registered at <http://www.irct.ir> (registration number: IRCT 201403061760N33; date registered: April 27, 2014).

Study design

Scaling and root planing was performed for all patients and all patients were given oral hygiene instructions. Occlusal adjustment was performed when necessary. Next, the right or left side was randomly selected for treatment with gingival unit graft (palatal tissue involving marginal gingiva and interdental papilla) or conventional free palatal graft (palatal tissue only) to be placed supraperiosteally at the recipient sites. In this study, a simple randomization method was used. The two treatment methods were written on cards and placed in envelopes. Right before the surgery on one side, a clinician selected an envelope and performed the procedure written on the card. The other surgical method was performed on the other side. All the surgical procedures were performed by the same surgeon while the clinical

measurements and randomization were done by another clinician.

Clinical assessments

The clinical parameters were measured at baseline and at one, three and six months after the surgical procedures to the nearest 0.5 millimeter using a University of Michigan "O" probe with Williams marking. The measured clinical parameters were as follows:

- Keratinized tissue width (KTW): Distance between the most apical part of the gingival margin and mucogingival junction.
- Clinical attachment level (CAL): Distance between the cementoenamel junction (CEJ) and bottom of the pocket
- Probing depth (PD): Distance between the most apical part of the gingival margin and bottom of the pocket at three points of mesial, midbuccal and distal.
- Vertical recession depth (VRD): Distance

between the CEJ and the most apical part of the gingival margin.

-Recession width (RW): Width of exposed root 1mm apical to the CEJ [27].

On the 10th postoperative day (suture removal session), the patients were asked about post-surgical pain using a 10-point visual analog scale (VAS) in which, 0 indicated no pain and 10 represented the worst pain experienced.

In addition, patients were questioned about esthetics at one, three and six months after surgery. Patient satisfaction was assessed using a 10-point VAS (0 indicated "dissatisfied" and 10 indicated "fully satisfied").

The Landry's healing index was assessed according to Jankovic et al, [28] at 10 days and one month after the surgery (Table 1).

The coverage percentage was calculated using the following formula: [(baseline VRD–new VRD)/baseline VRD]×100 at one, three and six

Table 1: Healing index used in the present study

Score	Characteristics
1: Very poor	<ul style="list-style-type: none">• Tissue color: ≥50% of gingiva red• Response to palpation: bleeding• Granulation tissue: present• Incision margin: not epithelialized, with loss of epithelium beyond incision margin• Suppuration present
2: Poor	<ul style="list-style-type: none">• Tissue color: ≥50% of gingiva red• Response to palpation: bleeding• Granulation tissue: present• Incision margin: not epithelialized, with connective tissue exposed
3: Good	<ul style="list-style-type: none">• Tissue color: ≥25% and <50% of gingiva red• Response to palpation: no bleeding• Granulation tissue: none• Incision margin: no connective tissue exposed
4: Very good	<ul style="list-style-type: none">• Tissue color: <25% of gingiva red• Response to palpation: no bleeding• Granulation tissue: none• Incision margin: no connective tissue exposed
5: Excellent	<ul style="list-style-type: none">• Tissue color: all tissues pink• Response to palpation: no bleeding• Granulation tissue: none• Incision margin: no connective tissue exposed

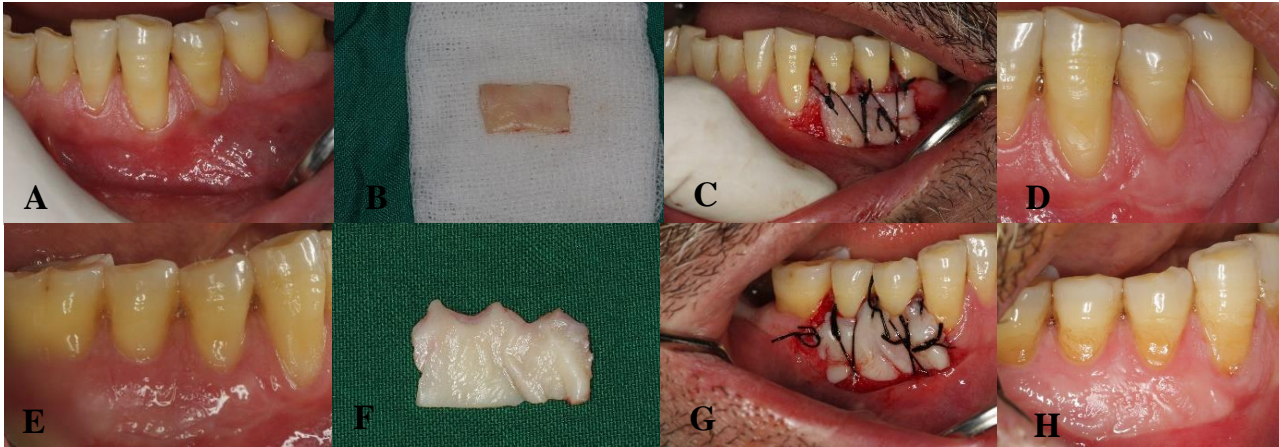


Fig. 1: Surgical procedure and follow-up (A) Initial clinical appearance at the free gingival graft side, (B) Free gingival graft, (C) Suturing, (D) After six months, (E) Initial clinical appearance at the gingival unit graft side, (F) Gingival unit graft, (G) Suturing, (H) After six months

months after surgery.

The CAL was measured at the midpalatal area of the relevant premolar tooth at baseline and at three and six months after gingival unit graft harvesting for clinical evaluation of donor site healing.

Surgical procedures

Recipient site

After achieving local anesthesia in both groups, the preparation of the recipient site was begun by making two vertical incisions extending to the adjacent teeth and about 3 to 4 mm beyond the mucogingival junction, as well as a horizontal incision at the mucogingival junction. A sharp split-thickness flap was reflected, and the surfaces between these incisions were de-epithelialized (Figs. 1A and 1E).

At the gingival unit side, the interdental papillae were removed; but, at the conventional graft side, the butt-joint incisions were made on the papillary tissue at the level of the CEJ. The exposed surface of the root was planed with hand instruments and rinsed with sterile saline.

Donor site

At the gingival unit side, the palatal tissue, including the marginal and interdental gingiva, was harvested from the palatal aspect of the maxillary premolars (Fig. 1F). On the other side (conventional graft), the palatal tissue was

harvested from the same area on the other side but about 2 mm away from the gingival margin (Fig. 1B). For the grafts at both sides, care was taken to obtain a thickness of about 1 mm. Next, the grafts were contoured, adapted and sutured at the level of the CEJ (Figs. 1C and 1G). Both the donor and recipient sites were covered with periodontal dressing. The sutures were removed after 10 days.

Postsurgical care

Patients were advised not to brush their teeth at the surgical site, avoid chewing hard food, and rinse once daily with 0.2% chlorhexidine digluconate mouthwash for three weeks. Systemic antibiotic (500 mg amoxicillin, tid) was prescribed for one week. After 10 days, the patients were instructed to resume gentle brushing, directed coronally, at the operated sites. Postsurgical recalls were scheduled every other week during the first month and at one, three and six months after the surgery (Figs. 1D and 1H).

Statistical analyses

Statistical analyses were performed using SPSS version 21 (SPSS Inc., IL, USA). Quantitative data of midbuccal measurements of the recession sites were recorded as mean±standard deviation. The data were analyzed using repeated measures two-way ANOVA and Tukey's post hoc test for intragroup comparisons. Paired *t*-test was used for intergroup comparisons.

RESULTS

Nine patients eligible for this study were treated in a split-mouth design. All patients completed the follow-up sessions. Healing was uneventful and there were no complications.

No significant differences in pre-surgical parameters were found between the two sides. Paired *t*-test found no significant difference between the two groups at each time point. The gingival unit graft produced significantly higher esthetic satisfaction at one, three and six months after the surgery ($P=0.050$, $P=0.024$ and $P=0.024$, respectively). Higher healing index (4.44 ± 0.88 at the gingival unit graft side and 3.44 ± 1.01 at the free gingival graft side, $P<0.001$) and lower level of pain (3.33 ± 1.50 for the gingival unit graft side and 5.22 ± 1.64 for the free gingival graft side, $P=0.020$) were noted at suture removal session. Higher root coverage percentage was obtained one month after surgery (30.04 ± 12.41 at the gingival unit graft side and 20.24 ± 16.78 at the free gingival graft side, $P=0.011$). Reduction of recession width was observed three months after surgery ($P=0.007$; Fig. 2)

The mean percentage of root coverage at three months after surgery was 44.04 ± 18.78 (range: 16.66 to 75) and 26.33 ± 24.28 (range: 0 to 66.66) in gingival unit graft side and free gingival graft side, respectively and this difference was not significant ($P=0.110$). The mean percentage of root coverage at six months after the surgery was 60.52 ± 21.22 (range: 28.57 to 100) and 45.52 ± 21.94 (range: 0 to 66.66) in gingival unit graft side and free gingival graft side, respectively and this difference was not significant ($P=0.062$, Table 2).

Complete defect coverage, defined as gingival margins at the level of the CEJ, was found in one of nine patients (11%) after six months in unit graft side; whereas, none of the patients showed complete coverage in the other side.

No complications were observed at the gingival unit donor site, and healing was uneventful. The CAL of the relevant maxillary premolar at the

gingival unit donor site was 2.22 ± 1.52 , 2.00 ± 1.54 and 2.00 ± 1.54 at baseline, three months and six months after surgery, respectively and there was no recession or attachment loss in this area but significant attachment gain was observed after three months ($P=0.00$).

There were no significant changes in midbuccal PD at each side. A significant reduction in VRD was observed after surgery at each side and the coronal movement of the gingival margin at the gingival unit side was greater than that in the other side; this difference was not significant between the two sides ($P>0.05$, Table 2).

DISCUSSION

This randomized controlled clinical trial lasted for six months and evaluated the treatment of localized gingival recessions (Miller class I/II) with an alternative technique (gingival unit graft) in which, palatal grafts include the marginal gingiva and the papillary tissue to use a site-specific vascular configuration.

Gingival arterioles are oriented in an apico-coronal direction. Capillaries in the marginal gingiva form repetitive networks and several small vessels form loops extending towards the marginal gingiva. Additionally, it has been demonstrated that predominant gingival vessels decrease in size and increase in number as they extend coronally [16]. Thus, in this modified

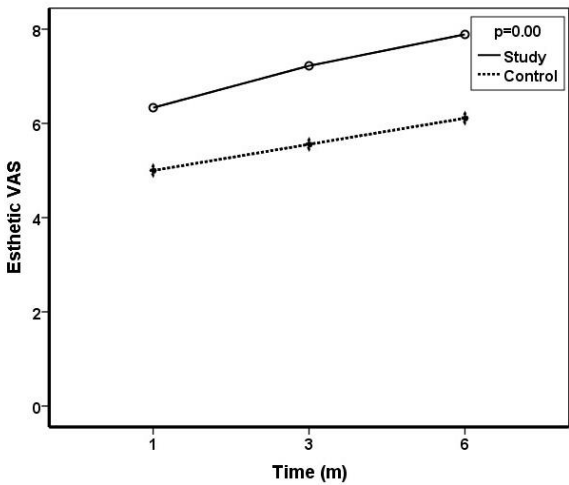


Fig. 2: Esthetic visual analog scale score changes

Table 2: Clinical parameters at baseline and at one, three and six months

Parameters		Baseline	One month	Three months	Six months	P-value ²
KTW (mm)	Study	2.44±1.52 ^a	5.94±1.07 ^b	5.33±1.03 ^c	5.05±1.01 ^d	<0.001
	Control	2.16±1.47 ^a	5.38±1.43 ^b	4.83±1.52 ^c	4.38±1.36 ^d	<0.001
	P-value ¹	Group= 0.273		Time= <0.001		Group*Time= 0.730
CAL (mm)	Study	5.33±1.85 ^a	3.72±0.83 ^b	3.50±1.14 ^b	2.66±1.56 ^c	<0.001
	Control	5.05±1.66 ^a	3.72±1.00 ^{bc}	3.72±1.12 ^b	3.00±1.17 ^c	<0.001
	P-value ¹	Group= 0.904		Time= <0.001		Group*Time= 0.499
VRD (mm)	Study	4.11±1.63 ^a	2.72±1.09 ^b	2.38±1.29 ^b	1.83±1.47 ^c	<0.001
	Control	3.72±1.46 ^a	2.83±0.93 ^b	2.61±1.08 ^b	2.00±1.11 ^c	<0.001
	P-value ¹	Group= 0.960		Time= <0.001		Group*Time= 0.315
RW (mm)	Study	3.00±1.19 ^a	2.50±0.82 ^b	2.11±0.82 ^{bc}	1.94±0.72 ^c	<0.001
	Control	3.16±1.54 ^a	2.50±1.17 ^b	2.66±1.19 ^b	2.44±1.21 ^b	0.011
	P-value ¹	Group= 0.205		Time= <0.001		Group*Time= 0.177
PD (mm)	Study	1.22±0.83 ^a	1.00±0.50 ^a	1.11±0.33 ^a	0.83±0.25 ^a	0.356
	Control	1.44±0.28 ^a	0.77±0.08 ^a	1.11±0.11 ^a	1.00±0.08 ^a	0.124
	P-value ¹	Group= 0.503		Time= 0.168		Group*Time= 0.136

In each row, the same superscripted letters show non-significant difference; 1: Repeated measures two-way ANOVA; 2: Repeated measures one-way ANOVA; KTW: Keratinized tissue width; CAL: Clinical attachment level; VRD: Vertical recession depth; RW: Recession width; PD: Probing

technique, the size and number of vessels and vascular configuration of donor tissue would better match those of the recipient site. In the current study, gingival unit side produced significantly greater esthetic satisfaction at one, three and six months, higher healing score, lower post-surgical pain score, higher root coverage percentage at one month and greater reduction in recession width at three months. These results indicate that this modified technique may improve these periodontal parameters. Proper plaque control, root surface biocompatibility, careful surgical manipulation and tissue thickness have been accepted as important factors affecting the outcome of graft procedures and should be controlled [16]. In the current study, one inclusion criterion was good oral hygiene during the study period. The exposed root surfaces were planed carefully with hand instruments to ensure root surface biocompatibility. Sutures were made without tension

over the graft to avoid graft displacement or impairment of blood supply. Special care was taken to obtain approximately 1mm thick grafts. Search of the literature yielded one previous comparative controlled clinical trial comparing gingival unit graft and conventional free gingival graft, which did not have a split-mouth design (eight patients in gingival unit graft, nine patients in free gingival graft) [16]. In the current study, in order to eliminate the inter-individual variability of the treatment effect, a split-mouth design was used. In the afore-mentioned previously published study [16], 50% of the sites in gingival unit group showed complete defect coverage at eight months, but none of the patients in the other group showed complete coverage. Allen and Cohen [22] in a case study reported that using gingival unit graft resolved the localized recession defects at three months. Another study used the modified gingival unit graft for treatment of Miller class III recession

defects. The results showed that gingival unit grafts provided better defect coverage than free gingival grafts [29].

In our study, 11% of the gingival unit sides showed complete coverage at six months. Similar to the above-mentioned comparative study [16], none of the free gingival graft sides showed complete coverage. The mean coverage percentage at the gingival unit side was 60.52% at six months compared to 45.52% at the free gingival graft side. The lower degree of coverage in the current study compared to the other study may be due to greater VRD and width at baseline. Probing depth did not change significantly, but keratinized tissue gain was significant in our study. These findings were similar to the findings of the afore-mentioned study [16]. Although reduction of VRD was significant at both sides and greater reduction was observed at the gingival unit side, this difference was not significant. The greatest reduction in VRD was observed during the first month, but then showed another significant reduction during three to six months. Migration of gingival margin in the coronal direction was probably because of creeping attachment. This phenomenon was first described by Goldman [30]. Creeping attachment may be observed one to 12 months after surgery and gives an average coverage of about 1mm in narrow recessions. The current study was limited to six months and a longer follow-up period is needed to draw definite conclusions regarding this phenomenon.

In the current study, esthetic satisfaction was assessed by asking patients their subjective opinion regarding the esthetic outcome. Significantly higher scores were obtained at one, three and six months after surgery for the gingival unit graft side compared to the free gingival graft side. Kuru and Yildirim [16] showed similar results regarding esthetic satisfaction. These findings, in addition to better root coverage, reveal that gingival unit graft can be used as a modification of free gingival graft, particularly in

areas where esthetics is important. Clinical healing at the gingival unit donor site was uneventful and without complications. It seems that some complications, such as necrosis of the primary palatal flap, which is commonly seen in subepithelial connective tissue grafts, do not occur in gingival unit grafts [31]. The recession of the donor site following denudation of the palatal bone, which is probably seen in laterally positioned flaps, does not seem to occur in this technique [32].

In the current study, care was taken to avoid significant changes in the location of the attachments. Even if the attachment apparatus was slightly injured, new attachment has always been observed to form quickly [33] without any problems.

Significant attachment gain was observed at three months, which was probably because of the formation of new attachments in this area. One of the benefits of this technique is its feasibility, even in presence of a thin palatal fibromucosa [16].

CONCLUSION

The present study showed that the use of gingival unit graft could be an acceptable treatment for Miller Class I/II recession defects. This technique may have advantages over the conventional free gingival graft technique such as significantly superior clinical and esthetic outcomes.

Additional studies with longer follow-up periods are required. More studies are needed to support the biological aspect of this technique by evaluating the blood flow and vasculature of the gingival unit graft.

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