Original Research Article

Coronally advanced flap with and without hyaluronic acid (HYALOSS) for the treatment of gingival recession — a randomized clinical trial

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Abstract – Objectives: To compare and evaluate the surgical outcome of gingival recession treatment using CAF alone or in conjunction with hyaluronic acid (HA) on Miller’s Class I and class II gingival recession defects. Materials and methods: After randomisation, the control group (15 patients) received CAF alone and the test group (15 patients) received HA as an adjunct to CAF technique for an isolated gingival recession accompanied by an adequate width of keratinized gingiva on maxillary canines and premolars. All patients were evaluated at 7, 15, 30, 60 and 90 days interval post-operatively. Data obtained was subjected to statistical analysis and p value <0.05 was considered as statistically significant. Results: At 90 days interval statistically highly significant root coverage was obtained with mean gain of 66.1% in control group and 86.6% in test group. Statistically significant reduction in depth of gingival recession and gain in clinical attachment level was found for control and the test groups, and intergroup comparison showed statistically significant differences for root coverage and clinical attachment level between the groups (p=0.000), however no significant differences were found for width of attached gingiva, keratinized gingiva and gingival thickness between the groups at 90 days (p > 005). Conclusion: HA has proven to be beneficial in the field of regenerative therapy. Our analysis suggested that HA can be used as a promising adjunct with CAF for root coverage, however further studies are required to imply the results on larger population.

Introduction

The ultimate goal of dental and periodontal care is to maintain the health, comfort, function, and aesthetics of the natural dentition. Gingival recession is most common periodontal condition characterized by root exposure as a result of displacement of gingival margin apical to cemento enamel junction and has become aesthetic concern due increase in clinical crown height. Incidence of GR has been found to increase with age, with prevalence rate >88% in individuals above 65 years of age [1]. Patients are usually asymptomatic but may experience pain due to exposed root surfaces and dentinal hypersensitivity. In addition, it is associated with root caries, cervical abrasion, and greater dental plaque accumulation. Early treatment of gingival recession is necessary to prevent progressive marginal tissue recession that may adversely affect prognosis of concerned tooth [2].

Periodontal plastic surgery procedures such as free gingival grafts, pedicle flaps, (laterally or coronally displaced) and subepithelial connective tissue grafts (SCTG) have long been used to treat gingival recession defects. Reports have shown conflicting success rates of these procedures and long term stability of root coverage is still questionable [3]. Connective tissue (CT) grafts have been demonstrated to be the most predictable surgical procedure for obtaining complete root coverage and is currently considered as the “gold standard” treatment for gingival recession [4,5]. However, CT grafts require a second surgical site to harvest the tissue and therefore are associated with undesirable side effects such as postsurgical pain, discomfort and potential postoperative bleeding. Coronally advanced flap (CAF) is a modality of root coverage surgery that does not involve a palatal donor site and has an advantage over palatal grafts by reducing several postoperative complications [6]. However, CAF does not always result in the regeneration of attachment apparatus such as cementum, periodontal ligament (PDL), and alveolar bone, which are major risk factors in the recurrence of gingival recession. Therefore, modifications to the
CAF technique are required, with success and predictability rates similar or superior to the CT graft. Several regenerative materials such as guided tissue regeneration membranes (GTR) [7], enamel matrix proteins derivatives (EMD) [8], acellular dermal matrix, alloderm (ADM) [9], platelet rich fibrin (PRF) and living tissue-engineered human fibroblast derived dermal substitute [10], have been used with CAF in the treatment of gingival recession.

Hyaluronic acid (HA) also known as hyaluronan or hyaluronate is relatively a new biomaterial that has got the potential to augment the results of periodontal therapy. It is a linear polysaccharide found in extracellular matrices of skin and is an indispensable component of intact, healthy gingiva. HA has many structural and physiological functions within tissues, including extracellular and cellular interactions, growth factor interaction, regulation of osmotic pressure and tissue lubrication, which helps in the maintenance of the structural and homeostatic integrity of tissues [11,12]. It has many other properties that makes it a potentially ideal molecule for assisting wound healing by inducing early granulation tissue formation, inhibiting inflammation, promoting epithelial turnover and connective tissue angiogenesis [12].

The purpose of the present study was to compare the clinical predictability of coronally advanced flap alone with that of coronally advanced flap+Hyaluronic acid in the treatment of Miller’s Class I and II gingival recession. The level of recession coverage, increase in width of attached gingiva, gain in clinical attachment level and thickness of keratinized tissue were assessed for a follow up period of 90 days.

Materials and methods

Study design

A single blinded randomized clinical trial was conducted in the Department of Periodontology after approval of the study protocol by the ethics committee of the institute. Informed consent was taken from each patient after explaining the benefits of the treatment.

Inclusion criteria

Out of 50 initially screened patients, 30 sites in 30 healthy adult patients (25 to 50 years) were included based on following criteria:

- Miller Class I and II gingival recession with ≥2 mm depth.
- Presence of identifiable CEJ and adequate width of keratinized tissue apical to recession.
- Patients who gave written informed consent to participate in the study.

Exclusion criteria

- Patients with relevant systemic condition or disease and, on drug therapy.
- Patients with presence of occlusal interferences such as mispositioned or rotated teeth.
- Habitual smokers and tobacco chewers.
- Patients with a history of previous mucogingival or periodontal surgery at an experimental site in last 6 months.
- Presence of gingival recession with caries, restoration or with pulpal pathology.

The sites of gingival recession were chosen on maxillary second premolar, first premolar, canine, lateral incisor and central incisor. GR was treated by CAF alone in control group and combination of CAF + HA (HYALOSS-meta-advanced medical technology, Italy) in the test group.

The primary endpoint of the study was percentage of root coverage at 90 days. All patients were examined under good illumination by using mouth mirror, William’s graduated periodontal probe, tweezer and pellets of cotton. The following clinical parameters (i) gingival recession depth, (ii) width of keratinized gingiva, (iii) probing depth, (iv) width of attached gingiva, (v) gingival thickness, and (vi) gain in clinical attachment level (CAL) were measured by single investigator by using cementoenamel junction (CEJ) as a fixed point. All the patients received scaling and root planning (SRP) before surgical therapy. A thorough plaque control programme was initiated for each patient 2 weeks prior to surgery including instructions in brushing technique (Modified Stillman) and supervised tooth brushing with soft tooth brush. 1 week after initial preparation, the patients were re-examined in order to ascertain their periodontal status. The degree of gingival inflammation was monitored by Gingival Index [13], and oral hygiene effectiveness was assessed by Plaque Index [14].

Sample size estimation

The sample size for the study was analysed using the following formula

\[ n = \left( \frac{Z_{a/2} + Z_{b}}{2} \right)^2 \times \frac{\sigma^2}{d^2} \]

where \( n \) is the minimum sample size, \( d \) is expected mean difference, \( \sigma \) pooled standard deviation (SD), \( Z_{a/2} \) is 95% confidence interval, \( Z_{b} \) is standard normal variate power for 80% power it is 0.84.

This clinical trial was powered to detect a minimum clinically significant difference in root coverage (recession reduction) of 0.5 mm using \( \alpha = 0.05 \), a power \((1 - \beta)\) of 80%, and a hypothesized within-group sigma of 0.4 mm. Considering possible dropouts, the number of patients was increased by 15% for each arm. On the basis of these data, the minimum number of patients needed to be enrolled in this study was 30 in total, 15 for the test (CAF + HA) and 15 for the control group (CAF only).

Randomisation, allocation concealment and blinding

Randomization was done using computer generated sequence wrapped in the sealed envelopes. The patients were operated by single surgeon with more than 10 years of experience in periodontal plastic surgery. The surgeon who was allotted to perform the surgery was given an envelope which
contained the information regarding the tooth on which interventional material was to be used. It was decided using coin toss by another surgeon who didn’t operate or assisted during the surgery. These envelopes were opened by the surgeon right before the surgery and accordingly the patients either received test material that is CAF + Hyaloss \((n = 15)\) or CAF alone \((n = 115)\). This was a single blinded study as the investigator was unknown about the type of intervention (CAF + HA or CAF only) the patient received and only the operator and the patient were familiar with this information.

**Surgical protocol**

One hour prior to surgery oral antibiotics (capsule amoxicillin 500 mg) were commenced. The area of surgery was disinfected using Povidone Iodine swab and was anesthetized using 2% xylocaine infiltration with 1: 50,000 epinephrine on both the facial and lingual aspect. After local anaesthesia (LA) and before the elevation of the flap, the exposed root surfaces were planed to reduce root convexity and washed for 60 s with water spray. Intraseulcular incisions were made on the buccal aspect of the involved tooth, followed by two horizontal incisions in the interdental papillae at the level of CEJ on the mesial and distal side of the defect extending to the adjacent teeth. Two apically directed vertical releasing incisions were given at the line angles of adjacent teeth extending beyond mucogingival junction (MGJ) into the alveolar mucosa. Trapezoidal split thickness flap was elevated by sharp dissection which was then followed by full thickness flap reflection by periosteal elevator up to MGJ. Beyond MGJ the trapezoidal flap was again reflected as split thickness by sharp dissection leaving the underlying peristomeum in place and extended as far as necessary to allow for flap advancement to the CEJ without tension. This necessitated a mesio-distal and apical extension of the partial thickness dissection in order to release any residual muscle tension and allow a passive coronal displacement of the flap. The vestibular epithelium of the adjacent interdental papillae was de-epithelized. The root surface was re-examined for any irregularities. Finally the flap was positioned at the level or slightly coronal to the CEJ and fixed with mattress suture. The two vertical incisions were sutured with interrupted sutures. Abundant saline irrigation was performed during the procedure. Figures 1 and 2 depict pre-surgical, surgical and post-surgical recipient site treated by CAF for Control Group.

For test group, CAF was identical to the one previously described in the control group, except for the addition of Hyaloss. After LA and root planning, the root surface was
washed for 60 s with water spray and dried. HA (Hyaloss) was applied to the exposed root surfaces, starting at the base of the recession using the applicator tip. The flap was then displaced coronally without tension at the CEJ level, fully covering the recession defects. A non-resorbable, number 4–0 vicryl suture was used to secure the CAF at the CEJ level by using interrupted sutures. Both vertical incisions were closed with interrupted sutures and the operated area was covered with periodontal pack (coe-pack). No pressure was applied to the flap after suturing. Figures 3 and 4 depict pre-surgical, surgical and post-surgical recipient site treated by CAF + HYALOSS, for test group respectively.

Subjects were prescribed analgesics (Ibuprofen 400 mg BD) for 3 days, capsule amoxicillin 500 mg, BD for 5 days and were instructed to use 0.2% chlorhexidine mouthwash twice daily. All the parameters were recorded post surgically at 1st, 2nd and 3rd month. The photographs were taken pre and post-operatively to assess the change in color and contour.

Results

Table I shows comparative analysis of parameters at baseline between the two groups.

Gingival recession

For control group, the mean depth of gingival recession at baseline was 2.67 ± 0.72 mm which reduced to 0.87 ± 0.52 mm at 90 days interval (p = 0.000) (Tab. II). For test group, the mean depth of gingival recession at baseline 3.13 ± 0.74 mm reduced to 0.47 ± 0.64 mm (p = 0.000) (Tabs. I and III). Inter group comparison showed that the mean gingival root coverage in the test group was more (86.6%) as compared to the control group (66.1%) and the mean difference of 0.87 ± 0.06 mm was found to be statistically significant (p = 0.004) (Tab. IV).

Clinical attachment level

Statistically significant gain in CAL was observed for both the groups at 90 days. Mean depth of CAL for control group at baseline was 4.53 ± 0.83 mm and test group was 5.00 ± 0.76 mm. At 90 days, CAL reduced to 2.60 ± 0.83 mm in control group and 2.20 ± 0.86 mm in test group (p = 0.000) (Tabs. I–III).

Keratinized gingival

For control group as well as test group, the mean width of keratinized gingiva was increased at 90 days (p = 0.000) (Tabs. I–III). However, intergroup comparison showed non-significant results between the groups (p = 0.607) (Tab. IV).
Fig. 3. Pre-surgical, surgical and post-surgical recipient site treated by CAF + HYALOSS, for Test Group, Case 3. (A) Pre-operative gingival recession defect with 13. (B) Horizontal and vertical incisions performed. (C) Split-Full – Split thickness flap reflected. (D) HYALOSS Gel. (E) Hyaloss applied at recipient site. (F) Tin Foil placed. (G) Vertical Incisions closed with Interrupted Sutures. (H) Periodontal Pack Placed. (I) Post-operative view at 15 days. (J) Post-operative view at 60 days. (K) Post-operative view at 90 days.

Fig. 4. Pre-surgical, surgical and post-surgical recipient site treated by CAF + HYALOSS, for Test Group, Case 4. (A) Pre-operative gingival recession defect with 14. (B) Hyaloss applied at recipient site. (C) Coronally advanced flap sutured over Hyaloss. (D) Post-operative view at 15 days. (E) Post-operative view at 90 days.
For both control group and test group, the mean width of attached gingiva was increased at 90 days and mean difference was statistically highly significant (p = 0.000) (Tabs. II and III). But, intergroup comparison showed that the mean difference of 0.13 ± 0.07 mm between test group and control group was statistically non-significant (p = 0.651) (Tab. IV).

**Attached gingival**

Attached gingival

The gingival tissue thickness was measured by using a file with rubber stopper midway between the gingival margin and MGJ and measured with a digital vernier caliper with 0.05 mm resolution. For control group, the mean thickness of gingival tissue at baseline, 1.48 ± 0.45 mm increased to 1.58 ± 0.41 mm at 90 days (p = 0.571) (Tabs. I and II). Similarly, for test group the mean thickness of gingival tissue at baseline,
has FDA approval [18]. It does not need cutaneous sensitivity testing and interference. Clinical trials have shown that HA could be used with no drug allergy reaction or discomfort, during the period of healing. HA was well tolerated by all the patients without any allergic reaction. Complications were successfully by the patients with no signs of infection or pain. A statistically significant difference in GR (root coverage) was observed between the groups (p = 0.004) and clinical attachment level (p = 0.000) between the groups (p = 0.004) between the groups. All the grafts appeared to have been taken up successfully by the patients with no signs of infection or complications i.e. pain, bleeding, suppuration, gangrene, swelling or discomfort, during the period of healing. HA was well tolerated by all the patients without any allergic reaction. Clinical trials have shown that HA could be used with no drug interference. It does not need cutaneous sensitivity testing and has FDA approval [18].

### Table IV. Shows intergroup comparison of parameters at 90 days after surgery for both control (CAF) and test (CAF+Hyaloss) group.

<table>
<thead>
<tr>
<th>S. No.</th>
<th>Parameters</th>
<th>Group</th>
<th>N</th>
<th>Mean</th>
<th>SD</th>
<th>‘t’ Value</th>
<th>p value</th>
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<td>Gingival sulcus depth</td>
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<td>0.35</td>
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<td></td>
<td></td>
<td>CAF + Hyaloss</td>
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<td>Gingival recession</td>
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<td>3.17</td>
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<td></td>
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<td>0.72</td>
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<tr>
<td>3.</td>
<td>Clinical attachment level</td>
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<td>1.93</td>
<td>0.79</td>
<td>2.86</td>
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<td></td>
<td>CAF + Hyaloss</td>
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<td>2.80</td>
<td>0.86</td>
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<td></td>
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<td>4.</td>
<td>Keratinized gingival</td>
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<td>0.87</td>
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<tr>
<td>5.</td>
<td>Attached gingival</td>
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<td>0.651</td>
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<tr>
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<td>CAF + Hyaloss</td>
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<td>0.83</td>
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<tr>
<td>6.</td>
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<td>1.04</td>
<td>0.309</td>
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<td></td>
<td></td>
<td>CAF + Hyaloss</td>
<td>15</td>
<td>0.09</td>
<td>0.21</td>
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*Significant p value.

1.51 ± 0.42 mm increased to 1.60 ± 0.42 mm at 90 days (p = 0.125) (Tabs. I and III). Intergroup comparison also showed non-significant results for mean gingival thickness between the groups (p = 0.309) (Tab. IV).

### Discussion

The present study compared the treatment effects on control sites (CAF alone) to that of test sites (CAF+ HA) on Miller’s class I & II gingival recession. HA has shown promising results in the field of regenerative therapy. It has been used extensively in the field of dentistry for its various beneficial applications. Exogenous application of HA has been shown to have anti-edema, anti-inflammatory and good wound healing properties [15]. In periodontics it has been used to treat gingivitis, periodontitis and periodontal intrabony defects [16,17]. Considering the study design and homogeneity of groups at baseline, differences in clinical outcomes were totally attributed to the techniques employed. Plaque index and gingival index scores were the supporting measurements that were recorded which established the degree of compliance by the patient and condition of periodontal soft tissue. Plaque index score was maintained ≤10% throughout the study.

Both the groups showed improved clinical outcome in terms of root coverage, clinical attachment level, width of keratinized gingiva and width of attached gingiva at 90 days compared to baseline with statistically significant difference in GR (root coverage, p = 0.004) and clinical attachment level (p = 0.000) between the groups. All the grafts appeared to have been taken up successfully by the patients with no signs of infection or complications i.e. pain, bleeding, suppuration, gangrene, swelling or discomfort, during the period of healing. HA was well tolerated by all the patients without any allergic reaction. Clinical trials have shown that HA could be used with no drug interference. It does not need cutaneous sensitivity testing and has FDA approval [18]. In the control group the mean depth of gingival recession was reduced significantly compared to that of baseline at 90 day interval. This is in accordance with the previous studies conducted by Zingale [19], Egli et al. [20] and Bernimoulin et al. [21]. Mean percentage of root coverage obtained in the control group was 66.1% which is in accordance with the values previously reported after the use of CAF, that ranged from 56.1% to 98.8% after 6 months follow up. Three sites out of 15 in the control group achieved 100% root coverage which was comparable to the results of other studies in the literature [22,23]. In the test group, the mean depth of gingival recession at baseline was reduced significantly at 90 days interval. The mechanism of action for root coverage by HA may be explained on the basis of its anti-inflammatory effect, which may be due to its action as a scavenger by draining prostaglandin, metalloprotei¬nases and the other bioactive molecules. It also scavenges reactive oxygen species such as superoxide radical (O2−), hydroxyl radical (OH·), thus preventing periodontal destruction [24]. HA has an anti-edematous effect which may be related to the osmotic activity. In dentistry HA has showed a positive effect on reduction of plaque and sulcus bleeding index of patients with plaque induced gingivitis [24]. In very few studies HA has been applied as an adjunct to SRP in nonsurgical treatment of periodontitis [25]. Johnson et al. reported significant reduction of bleeding on probing and periodontal destruction after the use of subgingivally applied 0.8% HA gel immediately post SRP and after 1 week [25]. This implies that reduction in gingival inflammation and edema due to anti-inflammatory and anti-edematous properties of HA might have prevented further apical migration of gingival margin leading to GR coverage in the test group. The mean root coverage in the test group at 90 days was comparable to the results reported by Cortes et al. (CAF+ Acellular Dermal Matrix, ADM) [26], Pilloni et al. (CAF+ Enamel matrix Derivative, EMD) [27], Gupta et al. (CAF+ Platelet Rich Fibrin, PRF) [28], and Kumar et al. (CAF+Hyaluronan) [29].
In the present study, nine sites in test group achieved 100% root coverage which was favourably comparable, and mean percentage of root coverage (88.6%) in the test group was much higher than reported by Kumar et al. (68.33%) [29]. This might be attributed to the variability in the observation period of the studies. But the mean root coverage in our study was comparable to the results reported by Jagannathachary et al. (82.2% using CAF + ADM) [30], and Jaiswal et al. (86.3% using CAF + EMD) [31]. At 90 days the mean amount of root coverage in control group was 1.80 ± 0.78 mm and in the test group was 2.67 ± 0.72 mm, with statistically significant difference between the groups (p = 0.004). The difference noted between the groups could be further explored and explained by analysing the soft tissue changes during the early healing phases. Both the procedures were performed with a clear goal of providing complete root coverage of treated roots. To reach this objective the best clinical skills was applied in trying to obtain a tension free pedicle flap, to position the flap margins coronal to CEJ and to provide flap stability with suturing technique.

Ballini et al. studied biological properties of HA that was seen to promote periodontal regeneration through space preservation and surface protection. It is said that the success of the root coverage procedure depends on the intimacy between the recipient bed and the covering tissues. A thin and stable clot assures better healing. HA promotes wound healing by cell migration and proliferation, it facilitates white blood cell infiltration and improves tissue hydration [32]. This could contribute to better stability of root coverage procedure that is seen with the use of HA.

This study showed statistically significant (p = 0.000) increase in the width of keratinized gingiva in both the groups could be due to influence of the granulation tissue derived from the periodontal ligament or the tendency of the MG line to regain its original position. The exact role of HA in stimulating the overlying mucosa to become keratinized is still not clear. However, it may be correlated with its role in keratinocyte proliferation [33]. HA is preferred over GTR membrane as the use of GTR is technique sensitive, moreover there is instability of GTR membrane under the pedicle flap, and bacterial contamination of membrane leads to compromised results. Use of non-resorbable membrane may cause perforation of overlying soft tissue and second surgical procedure [34]. At 90 days the mean increase in width of attached gingiva in control group was 1.00 ± 0.76 mm and in the test group was 1.13 ± 0.83 mm. The mean difference of 0.13 ± 0.07 mm was found to be statistically non-significant. The increase in the width of attached gingiva in the test group (1.13 mm) was in accordance with the results reported by Kuis et al. [35], and Jagannathachary et al. [30].

At 90 days, the mean gain in CAL in control group was 1.93 ± 0.79 mm and in the test group was 2.80 ± 0.86 mm. The mean difference of 0.87 ± 0.07 mm was found to be statistically highly significant (p = 0.000). Kumar et al. reported 2.3 mm of gain in CAL in the test group after the use of HA + CAF; but did not find any significant difference between the groups (CAF and CAF + HA) [29]. The increase in CAL in the test group could be attributed to embryologic and would healing properties of HA including the facilitation of cell migration and differentiation during tissue formation and repair [36]. HA shares bone induction characteristics with osteogenic substrates such as calcitonin gene related peptide (CGRP) and bone morphogenic protein [37,38]. Recent studies demonstrate that HA aids in the repair process of both soft and hard tissue (bone) [39]. During healing in the later granulation stage, HA synthesis stops and the existing HA depolymerized by host enzymes hyaluronidase, results in the formation of low molecular compounds and alteration of granulation tissue composition. This indicates that low molecular hyaluronic fragments formed after subsequent hyaluronidase activity promotes angiogenesis in the lesion that provides the necessary components required for rapid regeneration [39]. As per the existing literature few clinical studies have assessed thickness of gingival tissue with the use of HA and results showed significant increase in the thickness of gingiva [30]. Similarly, in this study at 90 days, the mean increase in the thickness of gingiva in the control group was 0.02 ± 0.14 mm and in the test group was 0.09 ± 0.21 mm but the mean difference of 0.07 ± 0.7 mm was found to be statistically non-significant.

HA has been found to be associated to the collagen molecules providing extracellular matrix elasticity, resistance and lubrication, and has an important function in areas with rapid cell proliferation, since it facilitates cell displacement. It is also a key component of chronic wound during the series of stages associated with the wound healing process in both mineralized and non-mineralized tissues. Though HA is mostly advocated in dentistry for topical application some recent studies were done on hyaluronic acid in biomaterials assisting regeneration [40]. Vanden Bogaerde demonstrated the osteoinductive effect of esterified low molecular weight HA preparation in the treatment of infrabony defects [41]. It was used as a co-adjunct with autogenous bone for grafting procedure. At 9 months from procedure, the dental parameters were virtually stable with a mean CAL gain of 2.6 mm. Radiographic evolution after 24 months showed defect fill. In another randomised controlled trial, Ni Jing et al. injected the HA gel into the base of the deficient papilla, and it was found that the height of the gingival papilla increased to 0.311, 0.45, and 0.4 mm from baseline at 3, 6, and 12 months, respectively, after treatment (p < 0.05) [42]. Shirakata et al. conducted an experimental study on dogs in which they found successful periodontal regeneration of gingival recession defects after treatment with CAF + HA. Statistically significant differences were observed in CAL (p < 0.05), width of gingival recession (p < 0.01), in formation of cementum and connective tissue attachment (p < 0.05) at 10 weeks in HA + CAF group [43]. From these observations we can assume some amount of true regeneration to take place under CAF with connective tissue attachment [41].

The histologic evaluations of the attachment obtained with laterally positioned flap, CAF, free gingival graft and connective tissue graft for the purpose of root coverage were most often reported in case reports [36,37]. Results of the study by
Shirakata et al. on dogs concluded that use of cross-linked hyaluronic acid with or without a collagen matrix has a potential to promote periodontal wound healing in Class III furcation defects [44]. Some reports showed long junctional epithelium, some connective tissue attachment while others showed small amount of regeneration. Often many reports showed more than one mode of attachment [36,37]. In the present study the tissue was firmly attached to the tooth in all the patients, and showed resistance to probing. The mean sulcus depth at 90 days in control group was 1.40 ± 0.51 and in test group was 1.20 ± 0.41 mm with no bleeding on probing. This type of attachment couldn’t be determined without human histology. Moreover, it requires removal of successfully treated tooth. This study showed promising results for HA in promoting periodontal regeneration but more clinical trials on larger population should be carried out to validate our results and long term analysis is necessary to see the stability of the outcomes.

Conclusion

Hyaluronic acid is a relatively newer biomaterial, the use of which is extending in dentistry for the treatment of mucogingival problems, reconstruction of interdental papilla, furcation involvement and periodontal regeneration. Our analysis suggested that HA may be used for successful recession coverage with coronally advanced flap but further studies are required to imply the results on larger sample size.

Conflict of interest

Authors declare no conflict of interest.

Funding

No funding.

Ethical approval

Ethical clearance has been obtained from ethics committee of the institute.

Informed consent

Has been obtained from the patients after explaining the aim and procedure.

Authors’ contribution


References


