

Use of flowable composite as intermediary layer in non-carious cervical lesions restored with composite resin: 48-month follow-up

Avaliação clínica de uma resina flow como uma camada intermediária em restaurações de lesões cervicais não cariosas: 48 meses

Abstract

Purpose: In this case report, the clinical performance of a microhybrid resin composite placed with or without a flowable resin composite was compared, over a 48-month period.

Case description: The patient of this case report presented 2 pairs of equivalent cervical abfraction lesions, under occlusion. Four restorations were placed in teeth 34, 35, 44 and 45. The restorations were divided into groups (Single Bond + Filtek-Flow + Filtek Z250 or Single Bond + Filtek Z250) and the materials were applied according to the manufactures instructions. Two previously calibrated operators placed the restorations and two other independent examiners evaluated the restorations at baseline and after 48 months, according to the USPHS criteria and modified criteria for color match.

Conclusion: After 48 months of evaluation the lesions restored with Filtek-Flow as a liner under Filtek Z250 did not show better clinical performance than the restorations without Filtek-Flow. All restorations showed a trend toward dark yellowing after 48 months.

Key words: Clinical evaluation; adhesive systems; composite resin; flowable composite

Resumo

Objetivo: Este relato de caso compara o desempenho clínico após 48 meses de restaurações de lesões cervicais não cariosas com uma resina composta microhíbrida associada ou não a uma camada de resina *flow* como um agente intermediário.

Descrição do caso: O paciente do presente caso apresentava 2 pares de lesões cervicais não cariosas ocasionadas por abfração sob oclusão. Nos elementos dentários 35 e 44 as restaurações foram feitas com Single Bond + Filtek-Flow + Filtek Z250 e nos elementos 45 e 34 com Single Bond + Filtek Z250, sendo os materiais empregados de acordo com as recomendações do fabricante. Dois operadores previamente calibrados colocaram as restaurações e dois outros examinadores avaliaram as restaurações no período imediato (*baseline*) e após 48 meses, de acordo os critérios USPHS modificado para o critério cor.

Conclusão: Após 48 meses as lesões restauradas com a resina *flow* como uma camada intermediária não demonstraram melhor desempenho clínico em relação às restaurações sem a resina *flow*. Todos os grupos apresentaram uma tendência à descoloração após 48 meses de acompanhamento clínico.

Palavras-chaves: Avaliação clínica; sistemas adesivos; resina composta; resina *flow*

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Received: October 27, 2009

Accepted: March 1, 2010

Conflict of Interest Statement: The authors state that there are no financial and personal conflicts of interest that could have inappropriately influenced their work.

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Introduction

Hard tissue loss of non-carious origin in the cervical region is a very common clinical condition, and its prevalence and severity increases with increasing age (1). According to De Munck et al. (2), non-carious cervical lesions are preferred for evaluating adhesive systems due to several factors, such as: 1) cervical lesions are completely expulsive and therefore, loss the of restoration can only be caused by a bond failure; 2) they always present margins in enamel and dentin; 3) as they are more common on the vestibular face of anterior and pre-molar teeth, they provide easy access for all restorative procedures and assessment procedures; 4) previous preparation of the restoration is minimum and/or dispensable, and relatively easy to perform, thus reducing the operator-related variables; 5) their prevalence is high, and generally several lesions are found in one patient, thus facilitating patient selection and development of the study design; and 6) despite the variability of cavity configuration factors and the consequent result, i.e., generation of excessive stresses at the bond interface, the properties of the materials used for restoration seem to be less important than the bonding procedure itself (3,4).

Furthermore, non-carious cervical lesions present dentin with a high degree of sclerosis, as well as high mineral content when compared with intact or caries-affected dentin. Hybrid layer formation in a region of sclerotic dentin, such as in a cervical lesion is difficult because this substrate does not favor the formation of a lasting bond (5).

The performance of several adhesive systems has been tested and the retention rate of conventional systems was clearly superior to that of previous generations of systems (2,6). However, the retention rate of conventional systems varies significantly over a period from 1 to 3 years (6). The material used for the restorative procedure has been partly responsible for premature failures. A clinical study with previous generations of adhesive systems has shown that retention of restorations in non-carious cervical lesions was not influenced by the modulus of elasticity of the resin composite (7).

The theory behind this concept is that: when a material has a high modulus of elasticity, it is considered inflexible when the tooth structure is deformed under the action of loads and therefore, it is capable of being displaced more easily from the cavity. On the other hand, a material with a low modulus of elasticity is capable of flexing/bending with the tooth structure under the action of loads, and consequently, the restoration can remain in position (3,7).

It has been proposed that microparticle composites show better performance in comparison with hybrid composites in abfraction defects due to the lower modulus of elasticity microparticle resins have (1,7). Based on this hypothesis, flowable composites that also present a low modulus of elasticity could minimize the development of stresses during function (8).

Generally, the flowable composites present reduced mechanical properties, such as the modulus of elasticity,

due to the smaller number of load particles disposed in the organic matrix (9). Consequently, some researchers have proposed the use of this material between the cavity walls and the final restoration, in order to absorb the stresses generated during polymerization shrinkage of the latter, usually performed with a material that has a high modulus of elasticity.

According to the above description, Unterbrink and Liebenberg (10) recommended the use of a thin radiopaque layer of flow resin on the adhesive to provide better sealing of the cavity margins. However, laboratory and clinical studies with the aim of evaluating the use of flowable composite resin as an intermediate layer between the final restorative materials have shown controversial results (8).

Thus, the aim of this case report is to describe the performance of resin composite restorations in non-carious cervical lesions either a using flowable composite resin, or not, as an intermediate layer over a period of 48 months.

Description of the Case

The patient, a 50-year old man, sought treatment due to the existence of non-carious cervical lesions in all the mandibular pre-molars that were causing him esthetic discomfort, as may be seen in Figure 1. The patient did not present a condition of hypersensitivity in any of these teeth, did show a good condition of oral hygiene, confirmed by the absence of carious lesions, existent restorations in good condition and good periodontal health.

The cervical lesions in teeth 34, 35, 44 and 45 showed significant loss of enamel and dentin, indicating an initial diagnosis of abfraction associated with wear caused by excessive tooth brushing. The degree of dentinal sclerosis in all the pre-molars was classified as type 3, which means teeth with moderate amounts of dentinal sclerosis according to Swift Jr et al. (11).



Fig. 1. Vestibular view of two cervical lesions of non-carious origin (34 and 35) diagnosed as resulting from an abfraction associated with abrasion. Note the lack of enamel at the cervical margin and degree of sclerosis grade 3.

The restorative proposal was to perform restorations of teeth 35 and 44 with the use of flowable composite resin as an intermediate layer (Single Bond + Filtek Flow + Filtek Z250) and of teeth 45 and 34 without the use of flowable composite resin as a layer between the adhesive and microhybrid resin (Single Bond + Filtek Z250).

Restorative Procedure

The restorative procedures were as follows: anesthesia, cleaning with pumice stone and water with a rubber cup, followed by rinsing and drying; selection of the Filtek Z-250 microhybrid resin shade by means of the color scale provided by the manufacturer; all restorations were performed under absolute isolation. No additional retention or bevel was made. All materials used in the restorative procedures were applied in accordance with the manufacturer's recommendations.

Restorations without Filtek-Flow: In teeth 44 and 36, the Single Bond adhesive system was applied in the following way: a – acid etching (15 s); b – washing (15 s); c – drying with an air stream (30 s); d – re-wetting dentin with water (humid technique); e – a layer of adhesive was applied (10 s) by rubbing on the surface; f – air stream at a distance (20 s); g – application of another layer of adhesive by rubbing (10 s); h – air stream at a distance (20 s); i – light activation for 10 s with a VIP appliance at 600mW/cm² (Bisco, Schaumburg, IL, USA). The lesions were filled with Filtek Z-250 in increments (± 3 increments). Each increment was light polymerized with the VIP halogen light appliance at 600 mW/cm² (Bisco, Schaumburg, IL, USA) for 40 s.

Restorations with Filtek-Flow: Teeth 34 and 45 were restored in a similar manner to teeth 44 and 35, with the exception that the Filtek-Flowable composite resin was used. After the adhesive was light activated, a thin layer of Filtek-Flowable composite resin (± 1.5 mm) was inserted and light polymerized for 40 s.

The restorations were finished and polished with diamond burs 1190F and 2135F (KG Sorensen, Barueri, SP, Brazil) with the aid of a spatula to protect the marginal gingiva (Fig. 2) and Sof-Lex Pop-On abrasive discs (3M ESPE, St. Paul, MN, USA).

The patient was followed-up periodically, after 1 week, 18, 36 and 48 months (Fig. 3) and all the restorations were clinically evaluated in accordance with the USPHS criteria (United States Public Health Service – U.S. Public Health Service) (12) modified for the color criterion. The following criteria were assessed: retention, anatomic shape, marginal discoloration, marginal desadaptation, secondary caries and post-operative sensitivity. The criterion for color combination used was that of Reusens et al. (13).



Fig. 2. Vestibular view of restoration of teeth 34 (restored with Single Bond + Filtek Z250, without Filtek-Flowable composite) and 35 (restored with Single Bond + Filtek Z250, with the use of Filtek-Flow as an intermediate layer). Note care taken when removing excesses in the cervical margin of the restoration.

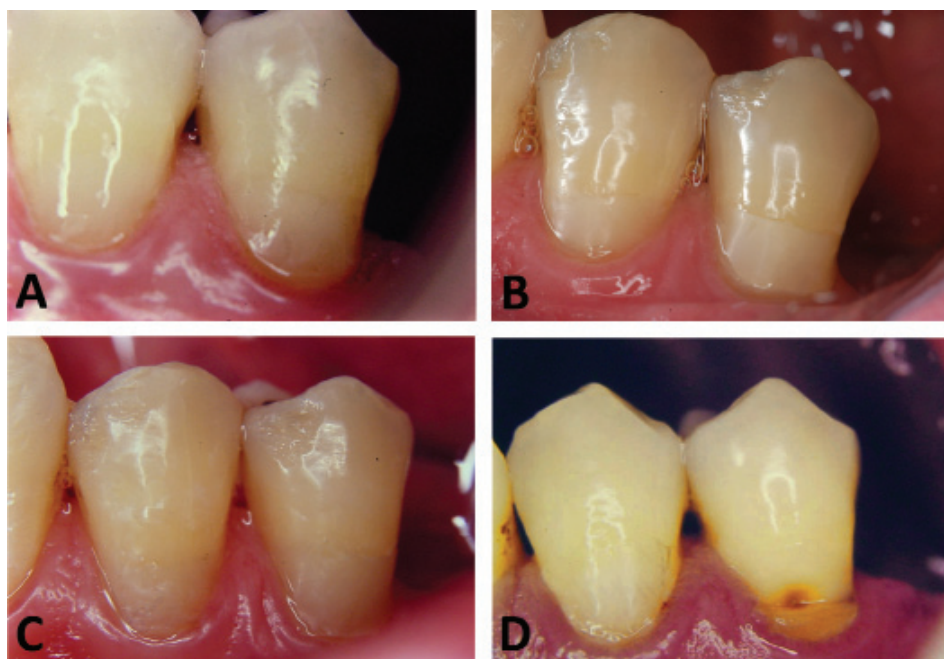


Fig. 3. Vestibular view of restoration of teeth 34 and 35 after 1 week (A), 18 months (B), 36 months (C) and 48 months (D). Note that after 18 months (D), there is a clear marginal discrepancy at enamel margin. After 36 and 48 months (C and D), restorations show loss of surface texture and after 48 months, loss of retention of the restoration of tooth 35 (restored with Single Bond + Filtek Z250, using Filtek-Flow as an intermediate layer).

In this criterion, the authors reclassified Score A of the USPHS system as A1 and A2. Traditionally, criterion A indicates that the restoration shows to be the same color as the tooth, and if there are differences in the color match and translucence between the restoration and the tooth, it is clinically acceptable. Reusens et al. (13) proposed the score of: A1 in which the restoration has an excellent color match to the point of not being perceptible, and A2 when the color match is good, but the difference with regard to the color between the tooth and the restoration is clinically perceptible.

Results

Up to 36 months of evaluation, there was no loss of retention, recurrent caries and loss of anatomic shape. After 18 months, the restorations began to present lack of marginal adaptation classified as "B". The patient did not indicate any spontaneous tooth sensitivity or sensitivity to an air stream.

All restorations showed a tendency to body discoloration, such as a type of yellowing, as from the period of 24 months (score A2). After 48 months, the restoration of tooth 35 was lost.

Discussion

Non-cariou cervical lesions are considered the model for clinical evaluation of adhesive systems, in accordance with the recommendation of the ADA (6). To obtain partial approval from the ADA, the adhesive systems need to present less than 5% of marginal discoloration after six months, and no loss of any restoration. Whereas to obtain final approval, the failure by loss of the restoration must not attain 10% after 18 months and less than 10% of the restorations may show marginal discolorations. The results of this clinical evaluation indicate that both groups tested would receive the seal of full approval.

One of the most important factors in the retention of non-cariou cervical lesions is the bond to the cavity walls, since this type of cavity does not present any type of mechanical retention of the bond, it is provided exclusively by the adhesive system. Although a series of other factors may directly influence the retention of Class V restorations, such as: occlusion; degree of dentinal sclerosis; and patient's age (1,6), correct diagnosis is the most essential factor.

It is well known that etiology of non-cariou cervical lesions is of a multifactorial nature (1). Patients with a history of bruxism or clinical evidence of other forms of traumatic occlusion usually generate high occlusal stress on these teeth. This creates an increase in flexure in the cervical region due to the high occlusal stress that may result in the restoration falling out.

An alternative used to maximize retention rate of Class V restorations is using a material with a low modulus of elasticity. These materials can serve as type of cushion, because they are flexible enough to resist the stresses

generated by polymerization shrinkage and facilitate the dissipation of these stresses produced by thermal variations, water sorption and occlusal load on the interface.

Some studies have shown an improvement in the performance of resin composite restorations when an additional layer of an intermediate material was placed between the resin composite and dentin substrate. Better dissipation of polymerization stresses (14), lower microleakage (14) and improved marginal adaptation (7) have been recorded.

As mentioned in the introduction, flowable composite resins have a low modulus of elasticity (8) and thus they may be used in cavities that undergo dental flexion (1). In addition, the flow capacity of this material becomes a desirable property due to good wetting, thus promoting better adaptation of the restorative material to the cavity walls.

However the nomenclature "flowable composite resin" is used for materials of very different characteristics (8). Flowable composite resins are more fluid materials, and this decrease in viscosity is always attributed to the reduction in the volume of inorganic material. In reality, the increase in fluidity may be achieved by modifying the monomer composition of the material, not necessarily indicating a reduction in the modulus of elasticity of the material (8).

Nevertheless, recent findings have concluded that the use of a flowable composite resin as an intermediate layer did not increase the retention rate of class V restorations after 12 and 24 months (15), indicating that other factors may be associated, irrespective of the presence of an intermediate layer absorbing stresses, to evaluate the retention rate of class V restorations.

Some studies have proposed the use of flowable composite resin as the only material to restore non-cariou cervical lesions (9). However, this technique has some disadvantages: 1) flowable composite resin present reduced mechanical properties when compared with microparticle and microhybrid resins (9); 2) lower availability of shades to match the tooth structure (8); microparticle and microhybrid resins offer a variety of shades to restore dental elements providing an excellent esthetic results; 3) it is more difficult to perform the sculpture of the restorations with flowable composite resins (9); 4) the high organic content of the flowable composite resins allow higher water sorption and greater discoloration over time, as has been shown for the microparticle resins that have a higher organic content than the microhybrid resins (13,15).

Whereas in comparison with Adper Single Bond, the 48-month findings of the present case only confirm the good results of this material in recent systematic review of literature (6)

It was concluded that: the use of Filtek-Flow as an intermediate layer did not improve clinical performance in comparison with restorations in which Filtek-Flowable composite resin was not used, after 48 months of clinical evaluation in non-cariou cervical lesions. In the clinical case presented, the restoration that was lost was lined with the flowable composite resin.

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