Comparative study to evaluate shear bond strength of RMGIC to composite resin using different adhesive systems

Manoj G. Chandak, Navdheeraj Pattanaik, Ayan Das

Abstract

**Aim:** The aim of the study is to compare and evaluate the role of new dental adhesives to bond composite to the resin-modified glass ionomer cement (RMGIC). **Materials and Methods:** Thirty specimens were prepared on acrylic blocks, with wells prepared in it by drilling holes, to retain the RMGIC. The specimens were randomly divided into three groups of ten specimens each. In Group a thin layer of selfetch adhesive (3M ESPE) was applied between the RMGIC and the composite resin FILTEK P60 (3M SPE). In Group II, total etch adhesive (Adeper Scotch bond 2, 3M ESPE) was applied, and in Group III, there was no application of any adhesive between RMGIC and the composite resin. After curing all the specimens, the shear bond strength was measured using an Instron universal testing machine. **Results:** The results were drawn and tabulated using ANOVA-fishers and Dunnet D statistical tests. The maximum shear bond strength values were recorded in Group I specimens with self-etch adhesive showing a mean value of 2.74 when compared to the Group II adhesive (Total etch) showing a mean shear strength of value 1.89, where no adhesive was used, showed a minimum mean shear bond strength of 1.42. There was a great and significant difference between Group I and Group II (P value 0.05) whereas, both Group I and Group II showed a vast and significant difference from Group III (P value = 0–001). **Conclusion:** Hence, this present study concludes that application of self-etch adhesive (3M ESPE, U.S.A) in between RMGIC and composite resin increases the shear bond strength between RMGIC and the resin composites, as compared to the total-etch type adhesive (Adeper Scotch bond 2,3M ESPE, U.S.A) as well as without application of the adhesive agent.

**Keywords:** Sandwich technique, self-etch adhesive, total-etch adhesive

Introduction

One of the major challenges in dentistry has been to find an ideal restorative material that has physical properties similar to those of tooth structure, adhesion to dentin and enamel, in addition to resistance to degradation in the oral cavity. In attempt to reach these characteristics, glass ionomer cement (GIC) was developed and first presented by Wilson and Kent[1] in 1972. A remarkable improvement of this class of material occurred approximately 15 years ago, with the introduction of the resin-modified glass ionomer cement (RM-GIC). This material is characterized by the addition of photo-activated methacrylate, and a small amount of resin, such as 2-HEMA or bisphenol-A-glycidyl methacrylate (Bis-GMA), to the conventional glass ionomer cement (GIC) liquid or powder. Presently, RM-GICs present two or three setting reactions: (a) acid–base reaction, typical of conventional GICs (initiated when the powder and liquid are mixed, occurring without light); (b) photo initiated setting reaction through the methacrylate groups (initiated when the powder/liquid mixture is exposed to light and occurs only where the light penetrates); (c) free-radical methacrylate curing without light (initiated when the powder and liquid are mixed without necessity of light).[3] A strong bond between RMGIC and the composite resin is an important factor for the quality of bilayered restoration. Etching of RMGIC is not required prior to the bonding of the composite resin.[4] However, application of resin bonding agents promotes the adhesion of the resin composite to both conventional GIC and RMGIC. There is limited literature on the bond strength of RMGIC to composite resin with adhesive agents in between. However, bonding agents have been seen to improve the wettability of GIC to help it adhere to the composite resin.[5]

Studies have shown that bonding agents who demonstrate a high degree of wettability, low viscosity, and low contact angle achieve a better union between GIC and the resin composite. Newer adhesive agents have undergone various modifications, such as, changes in viscosity, modification of primers, addition of nanofillers, and so on, to improve the bond strength between the tooth and composite resin.[6] Hence, the present study was conducted to evaluate and...
compare the shear bond strength of RCMGC to composite resin, using different generations of bonding systems applied on RCMGC.

Materials and Methods

Resin modified GIC (Vitrebon 3M ESPE, St. Paul USA) was bonded to a resin composite (Filtek™ F-60 3M ESPE, St.Paul USA) by using two different bonding agents, a total-etch adhesive (Adper™ Scotch Bond 2-3M ESPE, St.Paul USA) and a self-etch adhesive (Adper™ Prompt™ I, Pop™ -3M ESPE, St. Paul USA) and followed the curing with light (LED) curing light, Rotex, Taiwan.

Preparation of the specimens

The thirty specimens used in this investigation were prepared in 12 mm/25 mm dimensions, which was polished with 220, 320, and 400 grit carbide polishing paper. In each block, a well of 6 mm diameter and 2.5 mm depth was prepared by drilling holes in it to retain the RCMGC. Groove was placed on the walls for increasing the retention. The wells were then filled with light cure GIC by mixing it according to the manufacturer’s instructions and covering them with glass plates to produce a smooth surface and to permit light for curing the material. It was then cured with a blue light-emitting diode (LED) curing light (Rotex, Taiwan) for 40 seconds, according to the manufacturer’s recommendation, to produce a final set. The glass plate was carefully removed to ensure that the glass ionomer surface was smooth and not pitted.

Specimens were randomly divided into three groups of 10 specimens each, the groups were as follows.

Group A: To the light cure GIC (Vitrebon™) a thin layer of self-etch adhesive (3M ESPE) was applied according to the manufacturer’s instructions and cured, and then a cylinder of composite resin (Filtek™ F60, 3M ESPE, St. Paul USA) was added and cured over the specimen.

Group B: Same as group A, but a thin layer of total-Etch adhesive (Adper Scotchbond 2) was applied and cured over RCMGC.

Group C: Acts as a control group, no adhesive agent was applied between the light cure GIC and resin composite.

Immediately after following this procedure, a transparent plastic ring, 5 mm in height, with a 3 mm internal diameter, was centered over the resin-treated GIC, in the templates. The composite resin was condensed into a transparent plastic ring, using an incremental curing technique, above the RCMGC substrate, and all sides of the specimen were cured to ensure complete curing of the material. Following the curing the plastic ring was removed. All the procedures were conducted at room temperature. The bonded specimens were stored in distilled water at 37° for 24 hours, to simulate the conditions of oral cavity, until the specimens underwent the shear bonding test. The shear bond strength was measured by shearing of the bonded specimens on an Instron universal testing machine (model 4406), using a cross head speed of 3 mm/minute [Figure 1]. The shearing apparatus was constructed to grip the acrylic block, and a wedge blade system was designed to apply a shear force of approximately 0.1 mm on the adhesive interface. The readings were tabulated and subjected to statistical analysis using ANOVA, Dunnet D test. Mean and standard deviation were calculated for each group by using the ANOVA test and intergroup comparison was done by the multiple comparison test—Dunnet D test, which revealed a statistical significance among the groups.

Results

The mean shear bond strength and standard deviations were calculated for each group [Table 1]; and analyzed using the ANOVA test.

| N  | Mean | Std. deviation | Std. error | 95% Confidence Interval for Mean | Minimum | Maximum |
|----|------|----------------|------------|--------------------------------|---------|---------|
|    |      |                |            | Lower bound | Upper bound |         |         |
| Group A | 10 | 2.74 | 0.03 | 0.01 | 2.70 | 2.78 | 2.72 | 2.79 |
| Group B | 10 | 1.89 | 0.10 | 0.04 | 1.76 | 2.02 | 1.73 | 1.99 |
| Group C | 10 | 1.42 | 0.06 | 0.02 | 1.34 | 1.50 | 1.34 | 1.51 |

Table 2: Intergroup comparison and its statistical significance

| Mean difference (I-J) | Std. error | P-value | 95% Interval | Confidence |
|-----------------------|------------|---------|--------------|------------|
|                       |            |         | Lower bound  | Upper bound |
| Group A                | Group C    | 1.32    | 0.04         | 0.000 S, P<0.05 | 1.20 | 1.43 |
| Group B                |            | 0.47    | 0.04         | 0.000 S, P<0.05 | 0.35 | 0.58 |
The present study is in agreement with other studies where the effect of surface treatments and storage methods on composite/GIC were evaluated, where it was established that an application of a silane coupling agent over a non-etched GIC surface, followed by application of a bonding agent, demonstrated maximum shear bond strength. [9,10] In the present study, RMGIC was used over the conventional GIC under composite resin restoration because RMGIC sets by an acid–base reaction and exhibits a command set when activated by light or chemical agents via the methacrylate group. RMGIC has also demonstrated a better bonding to composite resin than the conventional GIC. [11,12] This is due to a similar chemistry between RMGIC and the composite resin, which allows the strong bonding of RMGIC to composite resin. Both RMGIC and the resin composite are cured by a free radical initiator system, which provides a potential for the chemical bonding between these two materials. 

**Conclusions**

From the results of the present study it can be concluded that application of bonding agents improves the wettability of glass ionomer cement to adhere to composite resin, thus promoting a strong shear bond between RMGIC and the resin composite. 

Moreover, the application of the self-etch adhesive between RMGIC and the composite resin showed significantly higher shear bond strength compared to the total-etch adhesive.

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