



Marginal Adaptation and Depth of Cure of Flowable versus Packable Bulk-fill Restorative Materials: An *In Vitro* Study

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Abstract

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AIM: The aim of the study was to investigate the marginal adaptation and depth of cure of a flowable bulk-fill giomer (BEAUTIFIL Flow Plus X [BFP]), a flowable bulk-fill resin composite (PALFIQUE BULK FLOW) bulk-fill resin composite, a packable bulk-fill giomer (BEAUTIFL-Bulk Restorative [BBR]), and two packable bulk-fill resin composites (X-tra fil) and (Filtek™ One Bulk Fill Restorative).

MATERIALS AND METHODS: Twenty-five standardized Class II cavities were prepared in the occlusomesial surfaces of maxillary premolars. A self-etching dental adhesive was used. All restorative materials were applied, and light cured according to their manufacturer's instructions. The teeth were subjected to 2500 thermal cycles between 5°C and 55°C. Epoxy resin replicas were obtained to examine the marginal by calculating the percentage of the continuous margin over the total margin length using scanning electron microscopic at 200× magnification. For assessing the depth of cure, fifty specimens with 4 mm height were prepared. Vickers microhardness testing was used to assess the depth of cure was calculating the bottom-to-top ratio of each specimen. If this ratio reaches 0.80 or more, an adequate depth of cure is achieved.

RESULTS: Regarding marginal adaptation, there was no significant difference between different groups before ($p = 0.398$) and after ($p = 0.644$) thermocycling. Within all groups, there was a significant decrease in marginal adaptation after thermocycling ($p < 0.001$). Regarding the depth of cure, all restorative materials achieved the required 0.8 bottom-to-top ratio. There was a significant difference between different groups ($p < 0.001$). The highest value was found in BFP group (0.97 ± 0.02), while the lowest value was found in BBR group (0.81 ± 0.11).

CONCLUSIONS: The marginal adaptation and depth of cure of bulk-fill giomer restorative materials are acceptable. Therefore, their use in restoration of 4-mm deep Class II cavities is appealing.

Introduction

Resin composite has become the material of choice for direct posterior restorations due to the increased demand for tooth-colored and mercury-free restorations [1], [2], [3]. Since the innovation of resin composites, they have undergone tremendous developments in their chemistry and application techniques producing restorative materials with excellent esthetics, relative stability, and wear resistance [4], [5]. Several clinical studies reported good long-term clinical performance and acceptable survival with an average annual failure rates ranging 1–3% [2], [6], [7], [8], [9]. Despite these continuous improvements in the field of resin composite technology, they still have some disadvantages, including high coefficient of thermal expansion than that of the tooth structures, water sorption, limited depth of cure, technique sensitivity and more importantly, and polymerization shrinkage [4], [10]. The shrinkage stresses, which inevitably occur, may be transmitted to the adhesive interface [11]. This, in turn, may compromise the quality of marginal adaptation of the restoration leading to marginal leakage with subsequent sequela including postoperative

sensitivity, marginal discoloration, caries adjacent to restoration (CAR), and pulpal irritation [12], [13], [14]. To overcome these catastrophic consequences and to ensure sufficient polymerization, incremental layering technique has been recommended for the placement of resin composite restorations [5], [15]. However, in deep posterior cavities, placing the restoration incrementally is time-consuming and increases the risk of contamination between successive layers or air bubbles entrapment [16].

Bulk-fill resin composites were introduced as a new class of resin composites to simplify handling, decrease restorative time, and improve clinical performance [17], [18]. They were developed by increasing the translucency and inclusion of special modulators and more sensitive photoinitiators allowing greater light to pass through the material to ensure uniform polymerization and proper degree of conversion. These modifications make the placement of 4–5-mm-thick increments feasible [19], [20]. Several bulk-fill resin composite materials are available on the market. They are classified according to their viscosity into two categories: packable (high-viscosity) and flowable (low-viscosity) [21], [22], [23], [24]. Packable

bulk-fill resin composites contain more inorganic fillers and are much more resistant to slumping, whereas flowable bulk-fill resin composites adapt better on the cavity walls [23].

Marginal leakage accounts for over 50% of failures of resin composite restorations [25]. Thus, achieving good marginal seal is climacteric for the long-term clinical success [26]. The evaluation of marginal adaptation in Class II restorations is a common procedure for evaluating the long-term stability of resin composite restorations [27]. Several direct or indirect methods have been proposed for this purpose by using clinical or laboratory tests. Among the *in vitro* tests, most studies are conducted using dye penetration, and/or scanning electron microscopic (SEM) analyses [28]. The leakage test with dye penetration represents the most frequently used method due to its simplicity. However, standardization of this process is not possible [23]. Therefore, SEM analysis is considered the gold standard for observation of marginal adaptation under a wide range of magnifications especially for indirect evaluation of restorations using replicas [26], [28], [29], [30]. These replicas are usually made of epoxy resin [23] and they allow accurate investigation and comparison of approximately the same marginal segments of the restoration after applying different ageing methods [31], [32].

One of the characteristics of bulk-fill resin composites is their improved depth of cure [22]. The depth of cure along with degree of conversion of the restorative material may influence the development of stresses, which are also likely to affect the integrity of tooth-restoration interface. If the depth of cure of the resin composite is limited, it is likely to induce less polymerization shrinkage stresses around the walls and margins of the cavity which, in turn, disguise an improved marginal adaptation due to poor polymerization [33]. The depth of cure of resin composites can be assessed using a variety of different methods. In general, they can be divided in two groups. First, depth of cure can be evaluated based on the degree of conversion. Second, it can be assessed indirectly by surface hardness either by scraping method according to ISO 4049 standard or in terms of the actual microhardness value. In the latter case, the values are expressed in percent, for example, bottom-to-top hardness ratio. A ratio of 0.80 was reported to be clinically acceptable [21], [22], [34], [35], [36].

A newly emerging trend in the dental industry is the development of hybrid materials combining the advantages of glass ionomers (anticariogenic and self-adhesive properties) and resin composites (esthetics, good mechanical strength, and high bond strength) [37]. Despite the controversy regarding the classification of these materials, they are sometimes called "bioactive" due to their ionic release property [37], [38]. Gionomers were introduced by incorporation of prereacted glass-ionomer (PRG) fillers into the resinous matrix of resin composite [39], [40]. Due to the great popularity

achieved by bulk-fill resin composites, "gionomer" bulk-fill restorative materials were commercially introduced [41]. Surface PRG fillers are currently incorporated in bulk-fill technologies as a high viscosity bulk-fill gionomer material (BEAUTIFL-Bulk Restorative [BBR], SHOFU INC., Kyoto, Japan) and more recently, a low viscosity bulk-fill gionomer material (BEAUTIFIL Flow Plus X [BFP], SHOFU INC., Kyoto, Japan). At present, the data available for these newly introduced restorative materials are limited and further laboratory investigations are required. Therefore, the aim of this *in vitro* study was to evaluate the marginal adaptation and depth of cure of five bulk-fill restorative materials. The two null hypotheses tested were that there would be no significant differences in (1) the marginal adaptation and (2) the depth of cure of the materials under investigation.

Materials and Methods

The marginal adaptation and depth of cure of five different bulk-fill restorative materials; two flowable and three packable from different manufacturers were investigated and compared. BFP (SHOFU INC., Kyoto, Japan), PALFIQUE BULK FLOW (PBF) (Tokuyama Dental Corporation, Tokyo, Japan), BBR (SHOFU INC., Kyoto, Japan), X-tra fil (XF) (VOCO GmbH, Cuxhaven, Germany), and Filtek™ One Bulk Fill Restorative (FOB) (3M ESPE Dental Products, MN, USA) were used in this study. The restorative materials and their specification, composition, shade, lot number, and manufacturers are described in Table 1.

Marginal adaptation assessment

According to the results of a study by Campos *et al.* [42], sample size was determined at five teeth for each restorative material considering alpha (α) level of (5%) and Beta (β) level of (20%), that is, power = 80%; therefore, a total of 25 teeth were included in the study. Sample size calculation was performed using G*Power Version 3.1.9.2. Twenty-five human maxillary premolars, which were extracted for orthodontic or periodontal reasons, with almost similar buccolingual and mesiodistal dimensions were selected from patients ranging from 18 to 40 years old. The teeth were obtained after the acquisition of patients informed content. The inclusion criteria were absence of cavities, restorations, cracks, or structural defects. Each tooth was washed with running water, brushed, scrubbed, and cleaned from any blood or soft tissue deposits. The teeth were submerged in distilled water at room temperature till their use.

Specially designed cylindrical Teflon molds of 20 mm height and 17 mm internal diameter were constructed and filled with self-curing acrylic resin

Table 1: Materials investigated in the study and their specification, composition, shade, lot number, and manufacturers

Product	Abbreviation	Specification	Composition		Filler Wt% (Vol%)	Shade	Polymerization time (s)	Lot Number	Manufacturer
			Resin matrix	Fillers					
BEAUTIFIL Flow Plus X	BFP	Low viscosity bulk-fill giomer	Bis-GMA TEGDMA Bis-MPEPP	S-PRG based on fluoroboroaluminosilicate glass	72.5 (51%)	A2	10	102036	SHOFU INC., Kyoto, Japan
PALFIQUE BULK FLOW	PBF	Low viscosity bulk-fill resin composite	Bis-GMA TEGDMA Bis-MPEPP	Silica-Zirconia filler, Composite filler	70% (56%)	A2	10	064E59	Tokuyama Dental Corporation, Tokyo, Japan
BEAUTIFIL-Bulk Restorative	BBR	High viscosity bulk-fill giomer	Bis-GMA UDMA TEGDMA Bis-MPEPP	S-PRG	87% (74.5%)	U	10	031931	SHOFU INC., Kyoto, Japan
X-tra fil	XF	High viscosity bulk-fill resin composite	Bis-GMA UDMA TEGDMA	Barium boron aluminum Silicate	86% (70.1%)	U	10	2044565	VOCO GmbH, Cuxhaven, Germany
Filtek™ One Bulk Fill Restorative	FOB	High viscosity bulk-fill resin composite	AUDMA AFM Diurethane-DMA, 1,12-Dodecane-DMA	Silica, Zirconia, Silica-Zirconia cluster, Ytterbium trifluoride	76.5% (58.5%)	A2	10	NA51014	3M ESPE Dental Products, St. Paul, MN, USA

BIS-GMA: Bisphenol A dimethacrylate, TEGDMA: Triethylene glycol dimethacrylate, Bis-MPEPP: Bisphenol A polyethoxy methacrylate, UDMA: Urethane dimethacrylate, AUDMA: Aromatic urethane dimethacrylate, AFM: Addition fragmentation monomers, 1,12-Dodecane-DMA.

(Acrostone Cold Cure, Acrostone, Cairo, Egypt). All the teeth were embedded vertically in the Teflon molds up to 2 mm under the cemento-enamel junction. Standardized occlusomesial Class II cavities were prepared by single operator. All cavities were prepared above cemento-enamel junction. The buccolingual width of each preparation was 2 mm at the occlusal part and 3.5 mm at the proximal part. The occlusopulpal depth was 2 mm, while the occlusogingival depth was 4 mm. The width of the gingival seat was 1 mm. All enamel margins were not beveled. The accuracy of the dimensions was checked using a digital caliper (Aluminum Caliper 4", IOS Ortho, Stafford, USA). The cavities were prepared using #330 and #245 carbide burs (Komet®, Gebr. Brasseler GmbH & Co. KG, Lemgo, Germany) rotating in high speed handpiece (COMFORTdrive™ 200 XD, KaVo Dental, Fruehauf, Germany) with copious amount of water coolant. After every five preparations, a new set of burs was used.

The teeth then were randomly divided into five groups (n = 5). The prepared cavities were thoroughly rinsed and air dried. A self-etching dental adhesive (BeautiBond, SHOFU INC., Kyoto, Japan) was used. The adhesive was applied, left undisturbed for 10 s, air dried with gentle air for 3 s and then dried with stronger air until a thin and uniform bonding layer was obtained. The adhesive layer was light cured using LED light curing unit (Dr's light AT CL-AT24, Good Doctors Co., Ltd., Incheon, Republic of Korea) for 5 s with light intensity of 1400 mW/cm² and wavelength 400–490 nm. Light intensity was checked with a hand-held radiometer (Model 100, Kerr Corporation, California, USA). Before application of restorative materials, ultra-thin celluloid band (Omni-Matrix™, Ultradent Products, Inc., St. Louis, USA) was placed to provide proper packing and contouring. Each restorative material was applied according to the manufacturer's instructions. Packable bulk-fill restorative materials were applied into the prepared cavities using gold plated composite applicator. Each restoration was contoured with fine grit finishing diamond stones (Komet®, Gebr. Brasseler GmbH and Co. KG, Lemgo, Germany) and was polished with fine (24 μm) and superfine (8 μm) Al₂O₃

discs (Sof-Lex™ discs, 3M ESPE Dental Products, MN, USA) mounted in a low speed hand piece (FX22, NSK, Tochigi, Japan) for 20 s.

For evaluation of marginal adaptation, impressions of the restored teeth were taken using polyether impression material (Impregum™ Soft Polyether Impression Material, 3M ESPE Dental Products, MN, USA) and then poured with epoxy resin (Kemapoxy 150, CMB International, Giza, Egypt) to obtain epoxy replicas. All cavity margins were analyzed by SEM (Model Quanta™ 250 FEG, FEI Company, Oregon, USA) attached with EDX Unit (Energy Dispersive X-ray Analyses), with accelerating voltage 30 K.V. at a magnification of 200×. The values are expressed as a percentage of the continuous margin over the total margin length for the occlusal and proximal margins (Figures 1 and 2). The marginal analyses were carried out by one evaluator experienced with quantitative margin analysis who was blinded to the groups.

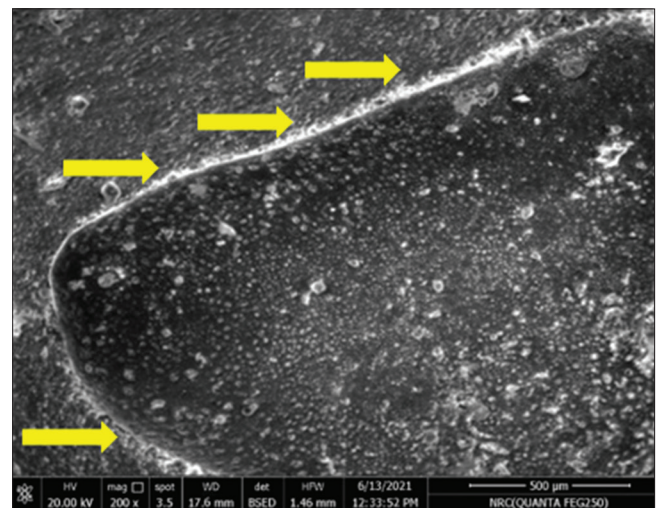


Figure 1: Representative scanning electron microscopic 200× image of continuous margins at the occlusal surface (yellow arrows)

All teeth were artificially aged by thermal cycling. The teeth were subjected to 2500 cycles in a thermocycling device (THE-1100 Thermocycler, SD

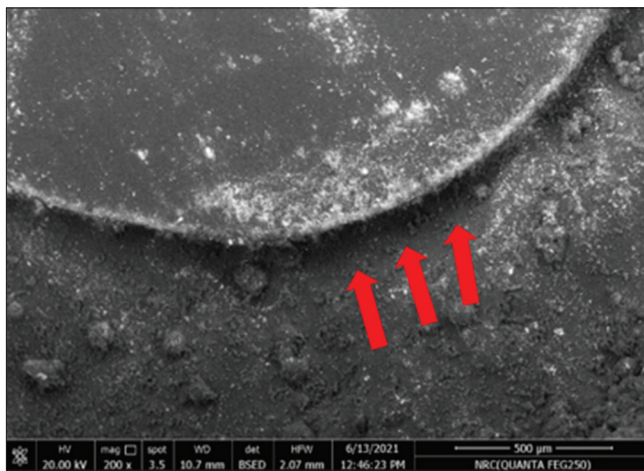


Figure 2: Representative scanning electron microscopic 200 \times image of discontinuous margins at the proximal surface (red arrows)

Mechatronik, Feldkirchen-Westerham, Germany) with water baths between 5°C and 55°C with a dwell time of 30 s in each bath and a transfer time of 5 s between each bath. After thermocycling, new epoxy replicas were obtained and examined as abovementioned.

Depth of cure by Vickers microhardness

Sample size was calculated based on the results of a study by Jang *et al.* [43]. Using alpha (α) level of (5%) and Beta (β) level of (20%), that is, power = 80%; the minimum estimated sample size was 10 disks per group. A split Teflon ring with an external diameter of 40 mm and internal diameter of 4 mm was used to prepare disks of restorative materials with 4 mm height. The Teflon ring was placed on top of a glass slab, and then each restorative material was applied inside the ring. A celluloid strip was placed on the Teflon ring and another glass slab was placed with slight pressure to produce a flat and smooth top surface of the disks. The disks were light cured by placing the tip of the light curing unit directly perpendicular on the celluloid strip. The upper and lower surfaces of each disk were polished with 2000-grit SiC after being removed from the split ring. The disks were then stored for 24 h at room temperature in dry and dark conditions.

A microhardness tester (Wilson[®] Tukon[™] 1102/1202, BUEHLER, Illinois, USA) was used to measure Vickers microhardness of the top and bottom surfaces of each disk. For these measurements, a 500 g force was applied for 5 s and three points were measured for each surface. The three readings were averaged to obtain a single value for each surface. The bottom-to-top surface microhardness ratio was

calculated. A 0.80 ratio indicates adequate depth of cure.

Statistical methods

Numerical data were represented as mean and standard deviation (SD) values. Shapiro–Wilk's test was used to test for normality. Homogeneity of variances was tested using Levene's test. Data were parametric and showed variance homogeneity. One-way analysis of variance test followed by Tukey's *post-hoc* test was used to study different intergroup comparisons. The significance level was set at $p < 0.05$ within all tests. Statistical analysis was performed with IBM[®] SPSS[®] Statistics Version 26 for Windows.

Results

Marginal adaptation

Results of inter and intragroup comparisons for marginal adaptation are presented in Table 2 and Figure 3. The results showed that there was no significant difference between different groups before ($p = 0.398$) and after ($p = 0.644$) thermocycling. Before thermocycling, the highest value was found in BFP group (94.81 ± 2.13), followed by PBF (94.24 ± 3.21), then BBR (92.69 ± 3.00), and FOB (92.52 ± 3.33), while the lowest value was found in XF group (91.63 ± 2.30). After thermocycling, the highest value was found in BFP group (80.45 ± 2.87), followed by XF (79.41 ± 2.51), then FOB (78.33 ± 3.98), and PBF (78.32 ± 2.94), while the lowest value was found in BBR group (77.39 ± 3.95). Within all groups, there was a significant decrease in marginal adaptation after thermocycling ($p < 0.001$).

Depth of cure

Results of intergroup comparison for bottom-to-top ratio are presented in Table 3 and Figure 4. The results showed that there was a significant difference between different groups ($p < 0.001$). The highest value was found in BFP group (0.97 ± 0.02), followed by PBF (0.93 ± 0.04), then XF (0.89 ± 0.05), and FOB (0.86 ± 0.08), while the lowest value was found in BBR group (0.81 ± 0.11). *Post hoc* pairwise comparisons showed

Table 2: Inter and intragroup comparisons for marginal adaptation (%) before and after thermocycling

Thermocycling	Marginal adaptation (%) (Mean \pm SD)				p-value
	(BFP)	(PBF)	(BBR)	(XF)	
Before	94.81 ± 2.13	94.24 ± 3.21	92.69 ± 3.00	91.63 ± 2.30	0.398
After	80.45 ± 2.87	78.32 ± 2.94	77.39 ± 3.95	79.41 ± 2.51	0.644
p-value	<0.001*	<0.001*	<0.001*	<0.001*	<0.001*

*significant ($p < 0.05$). BFP: BEAUTIFIL flow plus X, PBF: PALFIQUE BULK FLOW, BBR: BEAUTIFL-bulk restorative, XF: X-tra fill, FOB: Filtek[™] one bulk fill restorative.

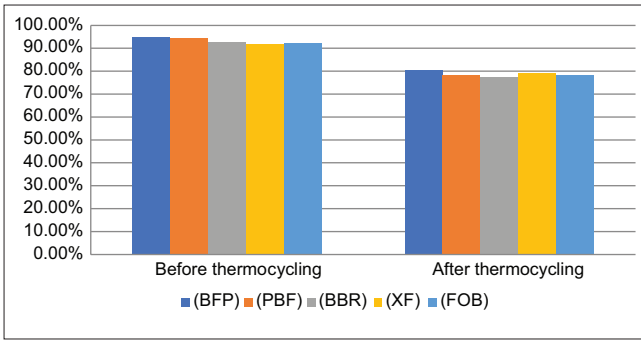


Figure 3: Bar chart showing average marginal adaptation (%) before and after thermocycling

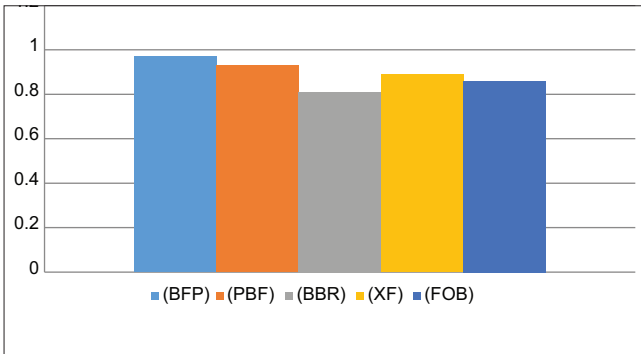


Figure 4: Bar chart showing average depth of cure (bottom-to-top ratio)

BFP to have a significantly higher value than all other groups except for (PBF) ($p < 0.001$).

Table 3: Intergroup comparisons for depth of cure (Bottom-to-top ratio)

Depth of cure (Bottom-to-top ratio) (Mean \pm SD)					p-value
(BFP)	(PBF)	(BBR)	(XF)	(FOB)	
0.97 \pm 0.02 ^A	0.93 \pm 0.04 ^{AB}	0.81 \pm 0.11 ^B	0.89 \pm 0.05 ^B	0.86 \pm 0.08 ^B	<0.001*

Different superscript letters indicate a statistically significant difference within the same horizontal row; *significant ($p < 0.05$), BFP: BEAUTIFIL flow plus X, PBF: PALFIQUE BULK FLOW, BBR: BEAUTIFIL-bulk restorative, XF: X-tra fil, FOB: Filtek™ one bulk fill restorative.

Discussion

Bulk-fill resin composites were introduced to shorten application time and to overcome the complications associated with incremental application technique [3]. The polymerization effectiveness of these materials has been questioned to guarantee adequate biocompatibility and mechanical properties and to decrease the risk of postoperative sensitivity to avoid early restoration failure [44]. This study focused on two of the aspects related to polymerization effectiveness: marginal adaptation and depth of cure. Proper marginal adaptation and absence of marginal leakage are core issues for the longevity of any restoration especially in the cervical margins of Class II cavities where the problem of microleakage becomes more pronounced [23], [45], [46], [47].

The results of present study showed that all investigated materials exhibited satisfactory marginal

adaptation before thermal cycling. Unfortunately, the level of marginal adaptation was not maintained after thermal cycling. There were no significant differences between different groups before and after thermocycling. Both flowable and packable bulk-fill resin composites showed similar marginal adaptation; therefore, the first null hypothesis was accepted. It could be argued that no significant difference was observed due to small sample size which is insufficient to detect differences between investigated materials. However, the sample size was chosen based on the results of a previous study by Campos *et al.* [42]. The marginal adaptation was expressed as a percentage of the continuous margins for the total judgeable margin length [24], [42]. Measuring the total length of marginal gap, rather than the width, is a valid method for analyzing the quality of marginal adaptation of resin composite restorations [28]. Based on SEM results, it was noticed that none of the groups could provide 100% continuous margins, regardless of the type of restorative material used. Discontinuous restoration margins may lead to CAR or loss of retention [24]. Two factors related to the resin composite are of key importance to marginal adaptation: polymerization shrinkage stress and elastic modulus. The lower elastic modulus of bulk-fill resin composites improves their adaptation to cavity walls [4]. The marginal adaptation of bulk-fill resin composites has been reported to be material-dependent [24], [46]. However, shrinkage stress is not a material's property. C-factor is among other several factors affecting the shrinkage stress [11]. In this study standardized Class II cavities were prepared with nearly similar dimensions; therefore, the C-factor was almost the same in all restorations. As previously described; marginal gaps may develop as a sequelae of polymerization shrinkage stresses if the shrinkage forces of the resin composite exceed the bond strength [24]. In this study, the self-etching dental adhesive (BeautiBond) was selected because it showed reliable results in both *in vitro* [48], [49] and clinical studies [50].

Reduction of shrinkage stress is an important feature of bulk-fill resin composites [51]. Though advertised as a new material class, most bulk-fill resin composites available in the market seem to have no significant modification in their chemical composition from conventional microhybrid and nanohybrid resin composites. They contain monomers such as Bisphenol A dimethacrylate (Bis-GMA), Triethylene glycol dimethacrylate (TEGDMA), and Urethane dimethacrylate in their organic matrix in addition to regular filler systems [51], [52]. Manufactures adopted different strategies to achieve sufficient polymerization and to reduce shrinkage stress [53]. These strategies involve using additional or more efficient photoinitiators [54] and addition of low shrink monomers, pre-polymerized fillers, or stress-relieving additives [14]. Other strategies focus

on increasing light transmission by using fillers and monomers with similar refractive index and reducing filler content [55], [56].

The results of the present study coincide with conclusion of a recent systematic review by Gerula-Szymańska *et al.*, [23] who concluded that the marginal adaptation of flowable and packable bulk-fill resin composites are similar when used for Class II restorations. In contrast to these findings, Paganini *et al.*, [24] reported that packable bulk-fill resin composites showed higher margin integrities compared with the flowable ones. The inconsistency between the results of the abovementioned study and the present study can be attributed to the differences in investigated bulk-fill restorative materials and testing protocol variations. The aging procedures were different between the two studies. The teeth were subjected to 1000 thermal cycles (between 5°C and 50°C) and 400,000 mechanical loading cycles while in our study the teeth were subjected 2500 thermal cycles (between 5°C and 55°C). It is noteworthy that elastic modulus of restorative materials determines their behavior under stress [42]. The elastic modulus was reported to be more important than the amount of shrinkage in determining the stress [57], [58]. Consequently, the high volumetric shrinkage produced by flowable bulk-fill resin composites can be compensated by the low elastic modulus, thus the stress buildup is reduced, and the marginal integrity is maintained [59].

A major concern regarding bulk-fill resin composites is whether the resin composite cures properly in the deeper portions of the restoration [42]. Inadequate curing of resin composites results in increased free monomers which are cytotoxic to the pulp and deteriorate the physicomechanical properties of the restoration [60]. According to the conclusions of the systematic review by Lima *et al.*, [44] most of the studies investigating the depth of cure of bulk-fill resin composites reported material-dependent results. Several factors may affect the depth of cure of bulk-fill resin composites. These factors include curing light time and intensity as well as resin chemical composition and most importantly fillers size, type, and volume [60]. The shade of bulk-fill resin composites influences their depth of cure [61]. In the present study, a light shade was selected for each material (shade A2, except for BBR and XF, which was a universal shade) to confirm that the effect of shading pigments would not be a confounding variable and to guarantee optimal light penetration through the restorations.

Depth of cure can be judged by determining microhardness ratio. If the bottom-to-top hardness ratio is 0.8 or more, an acceptable curing depth is achieved [20], [34]. In the present study, the bottom-to-top hardness ratio exceeded 0.8 in all bulk-fill restorative materials. However, the second null hypothesis was rejected because the depth of cure

values showed significant differences between different groups. Flowable bulk-fill restorative materials (BFP and PBF) recorded higher values in comparison with packable ones (BBR, XF, and FOB). These results are in line with the results of the previous studies [20], [21], [61] that reported greater depth of cure in flowable bulk-fill resin composites compared to packable ones. This could be attributed to the lower percentage of inorganic fillers in investigated flowable bulk-fill restorative materials, ranging between 70 and 72.5 wt%/51 and 56 vol%. In contrast to packable bulk-fill restorative materials which range between 76.5 and 87 wt%/58.5 and 74.5 vol%. The lower filler content in flowable bulk-fill restorative materials results in higher resin to filler ratio and a reduction in filler–matrix interface and light scattering leading to better light penetration and higher degree of conversion [20], [34]. Furthermore, the greater amount of organic matrix in flowable bulk-fill materials is an important factor causing higher depth of cure. It should be emphasized that low-molecular weight monomers such as TEGDMA and Bisphenol A polyethoxy methacrylate (Bis-MPEPP), which are the main constituents of the organic matrix, have higher reactivity and flexibility allowing increased formation of binding sites during light curing process, thus, enhancing the degree of conversion and increasing polymerization efficiency [44].

The resin matrix of giomers does not seem to differ from that contained in the resin composite being mainly constituted from Bis-GMA, TEGDMA, Bis-MPEPP and other monomers [37], [62]. Therefore, their polymerization efficiency is nearly like resin composites. A similar conclusion was reached by Ilie and Fleming [39]. They, too, concluded that flowable and packable giomer bulk-fill restorative materials showed an adequate depth of cure of >4 mm. On the other hand, some studies [60], [63], [64] reported that giomer bulk-fill restoratives were not able to achieve a depth of cure of 4 mm. Variations in results can be related to differences in study design regarding hardness testing conducted (Vickers vs. Knoop), hardness testing protocols and specimen mold material. Yap *et al.* [60] in their study assessed the depth of cure using Knoop hardness number (KHN). KHN was reported to be lower at or near the mold walls than at the center due to non-uniform distribution of KHN within molds [65]. Although Singla *et al.* [63] measured the depth of cure using Vickers hardness number as done in the present study, they reported different findings. This could be attributed to the difference in mold material (metallic versus Teflon). Metallic molds do not accurately reflect the depth of cure obtained clinically because they have different optical properties than tooth structure. Light transmittance through translucent tooth structure, may enhance the depth of cure of resin composites like the increased depth of cure reported with Teflon molds [54]. Tsujimoto *et al.* determined the depth of cure by the ISO 4049 scraping method. The reliability of scraping method was criticized because it is a subjective method

which may be affected by inter-operator differences in addition to difficulty to standardize the procedure of scraping off the uncured restorative material [20], [21].

Although *in vitro* testing of restorative materials provides an important initial screening for their properties, the obtained results can not accurately predict the clinical performance of restorations [66]. To mimic the influence of oral environment, the restorations in the present study were subjected to thermal ageing. Thermocycling is a valid *in vitro* method to simulate the thermal changes caused by drinking, eating, and breathing. However, there is no one standard protocol for thermocycling. Thermocycling regimens differ greatly. There is no agreement on the optimum number of cycles, temperature range, and dwell time [67]. Furthermore, a great contradictory exists between studies assessing microhardness regarding the indentation load and speed [68]. Hence, further laboratory and randomized clinical trials of bulk-fill restorative materials remain mandatory while overcoming the abovementioned limitations in the present study.

Conclusions

Based on the conditions and limitations of this *in vitro* study, it is possible to conclude that:

1. All investigated materials showed accepted marginal adaptation in Class II restorations
2. 100% perfect margins could not be achieved in any group
3. Thermocycling had detrimental effects on marginal adaptation
4. The 4-mm depth of cure of all investigated materials was acceptable (>0.80 bottom-to-top ratio)
5. Flowable bulk-fill restorative materials (BFP and PBF) showed statistically significant higher depth of cure values than packable ones (BBR, XF, and FOB)
6. The marginal adaptation and depth of cure of bulk-fill comonomers are acceptable and comparable to their resin composites counterparts.

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