

## Research Article



# Microhybrid versus nanofill composite in combination with a three step etch and rinse adhesive in occlusal cavities: five year results

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### Conflict of Interest

No potential conflict of interest relevant to this article was reported.

### Author Contributions

Conceptualization: Tuncer S, Demirci M; Data curation: Tuncer S, Demirci M, Öztaş E, Tekçe N; Formal analysis: Uysal Ö; Funding acquisition: Tuncer S; Investigation: Tuncer S, Demirci M, Tekçe N; Methodology: Tuncer S, Demirci M; Project administration: Tuncer S,

## ABSTRACT

**Objectives:** The aim of the study was to evaluate the 5-year clinical performance of occlusal carious restorations using nanofill and microhybrid composites, in combination with 3-step etch-and-rinse adhesives, in patients who were going to commence orthodontic treatment.

**Materials and Methods:** A total of 118 restorations for occlusal caries were conducted prior to orthodontic treatment. Occlusal restorations were performed both with Filtek Supreme XT (3M ESPE) and Filtek Z250 (3M ESPE) before beginning orthodontic treatment with fixed orthodontic bands. Restorations were clinically evaluated at baseline and at 1, 2, 3, 4, and 5-year recalls.

**Results:** None of the microhybrid (Filtek Z250) and nanofill (Filtek Supreme XT) composite restorations was clinically unacceptable with respect to color match, marginal discoloration, wear or loss of anatomical form, recurrent caries, marginal adaptation, or surface texture. A 100% success rate was recorded for both composite materials. There were no statistically significant differences in any of the clinical evaluation criteria between Filtek Z250 and Filtek Supreme XT restorations for each evaluation period.

**Conclusions:** The composite restorations showed promising clinical results relating to color matching, marginal discoloration, wear or loss of anatomical form, recurrent caries, marginal adaptation, and surface texture at the end of the 5-year evaluation period.

**Keywords:** Occlusal restoration; Clinical performance; Dental composite; Orthodontics

## INTRODUCTION

Caries, a common dental disease, is more prevalent in adolescents than in adults [1]. All patients should be screened for caries before and throughout orthodontic treatment. If a lesion is detected, it should be permanently restored and any pulpal involvement addressed before initiating treatment [1]. Therefore, prior to commencement of orthodontic treatment, it is necessary to treat all existing tooth caries. Caries can occur in pits and fissures and account for more than 60% of all lesions [2,3].

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Operative dentistry has seen much progress in recent years in posterior teeth restoration, including a gradual movement towards resin composites, rather than dental amalgam, which has allowed the adoption of minimal intervention approaches that conserve and preserve remaining tooth tissues and structures [4]. There are no reasons why posterior composite restorations should not have long survival rates as long as patient, operator, and materials factors are all considered at the time of restoration. Many of the improvements achieved in today's materials have been realized through continuously reducing the size of filler particles incorporated in the resin matrix of commercial dental composites to the present nano-composite materials [5,6].

Nanofill resin composite that contains nanosized fillers and/or nanofiller clusters has been developed [7], while the microhybrid composites are based on the particles averaging about 0.4–1.0 µm in size. These materials are recommended for both anterior and posterior restorations based on their combination of strength and polishability [7,8].

To the best to our knowledge there are no data available on the clinical performance of occlusal carious restorations completed prior to the initiation of orthodontic treatment. Therefore, the aim of study was to evaluate the clinical performance of restorations for occlusal caries over 5 years, which were performed using nanofill and microhybrid composites in combination with 3-step etch-and-rinse adhesive and completed before orthodontic treatment. The null hypothesis tested was that there was no difference in the clinical performance between the 2 composites after 5 years.

## MATERIALS AND METHODS

### Study design

The study protocol was approved by the local ethics committee (Ethical Committee of Istanbul University; Reference number: 2008/696). Details of the materials used in the study (etchant, adhesive, and composites) are shown in **Table 1**. The restorations were placed between April 2008 and December 2009 at the Department of Restorative Dentistry, Istanbul University Faculty of Dentistry. A total of 28 patients (8 males and 20 females), who were going to commence orthodontic treatment and aged between 11 and 28 years (mean age: 16.8 years)

**Table 1.** The brand names, chemical compositions, application procedures, and manufacturers of the materials used in the study

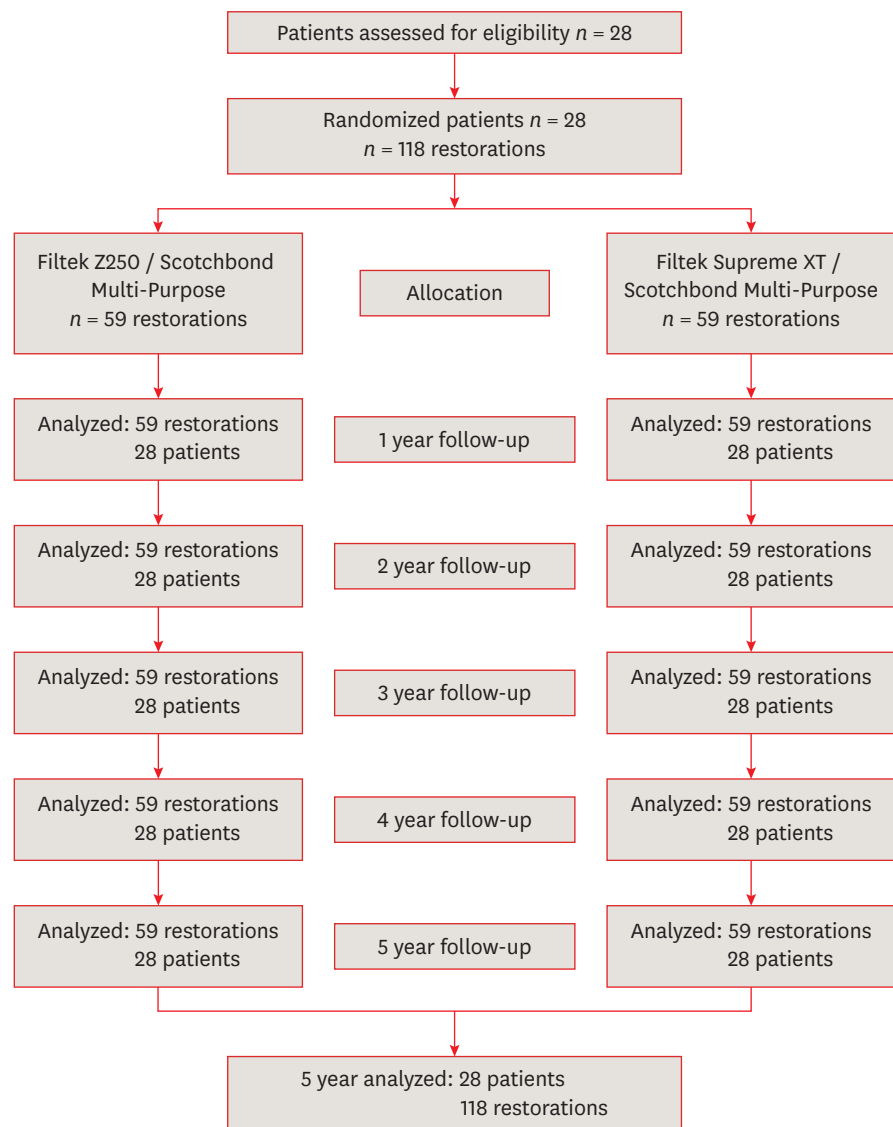
Adhesive	Component	Procedure	Manufacturer
Scotchbond Multi-Purpose	Etchant: 35% H <sub>3</sub> PO <sub>4</sub> , water, and silica Primer: 2-hydroxyethylmethacrylate, polyalkenoic acid, copolymer, and water Adhesive: 2-hydroxyethylmethacrylate, Bis-GMA, and photoinitiator	Apply the etchant to dentin and enamel for 15 sec. Rinse the surface for 15 sec and dry the surface slightly leaving a visible moist dentin surface. Apply the primer and dry gently for 5 sec. Apply the bond and light cure for 20 sec.	3M ESPE, St. Paul, MN, USA
Filtek Supreme XT	Bis-GMA, UDMA, TEGDMA, Bis-EMA, silica filler, zirconia filler, and aggregated zirconia/silica cluster filler	Tooth color to be restored was selected using the corresponding composite guide or custom composite samples before isolating the tooth. The corresponding body shade was selected. In 2 mm layers or less increments of body shade were applied. Each increment was light cured 20 sec.	3M ESPE, St. Paul, MN, USA
Filtek Z250	Bis-GMA, UDMA, Bis-EMA, silica filler, and zirconia filler	Tooth color to be restored was selected using the corresponding composite guide or custom composite samples before isolating the tooth. Place 3M Filtek Z250 restorative in increments less than 2.5 mm. Light cure each increment for 20 sec.	3M ESPE, St. Paul, MN, USA

H<sub>3</sub>PO<sub>4</sub>, phosphoric acid; Bis-GMA, bisphenol A glycidyl dimethacrylate; UDMA, urethane dimethacrylate; TEGDMA, triethylene glycol dimethacrylate; Bis-EMA, ethoxylated bisphenol A glycol dimethacrylate.

were included in the study. All persons gave their informed consent prior to their inclusion in the study. A total of 118 occlusal caries restorations completed prior to orthodontic treatment were evaluated (**Figure 1**). Each patient received between 2 to 8 occlusal restorations. Patient inclusion and exclusion criteria are presented in **Table 2** [7,9,10]. All teeth had opposing and adjacent tooth contacts. Patients were required to receive treatment for all existing tooth caries on the occlusal surfaces of posterior teeth before beginning orthodontic treatment. Therefore, the interaction of the 2 disciplines was managed in the diagnosis and clinical treatment of these patients.

### Treatment protocol

According to the faculty's patient treatment protocol, the patients were first examined in the Department of Oral Diagnosis and Radiology, Istanbul University. After periapical and/or panoramic radiographs were taken, patients were referred to related departments. First, the teeth were cleaned using pumice-water slurry and a rubber cup to remove the pellicle and



**Figure 1.** Flow diagram showing evaluation history of restorations.

**Table 2.** Inclusion and exclusion criteria

Criterion
Inclusion criteria
Patients received 2 to 8 restorations for primary caries on the occlusal surface
Occlusal contact with the antagonist tooth
Displayed good oral hygiene
Having no active periodontal or pulpal diseases
Patients were required to have received orthodontic treatment with fixed appliances
Willing to return for follow-up examinations as outlined by the investigators
Exclusion criteria
Patients with uncontrolled parafunction
Presenting insufficient oral hygiene
Pulp exposure during caries removal or cavities with imminent risk of pulp exposure
Spontaneous pain or sensitivity to percussion
Patients were pregnant or nursing
Patients had periodontal or gingival disease

residual plaque. Lesions were diagnosed macroscopically under a dental operating light using plain dental mirrors, air syringes, and World Health Organization (WHO) periodontal probes to check for surface discontinuity. The lesions were identified as being cavitated, if they had reached the dentin (International Caries Detection and Assessment System [ICDAS] codes 3–5), but the lateral spread was limited to the dentin.

For the patients who received 2 occlusal restorations, one was performed with a nanofill resin composite (Filtek Supreme XT, 3M ESPE, St. Paul, MN, USA), while the other was performed with a microhybrid composite (Filtek Z250, 3M ESPE). The microhybrid and nanofill resin composites were applied in combination with a 3-step etch-and-rinse adhesive (Scotchbond Multi-Purpose, 3M ESPE) (**Table 1**). The microhybrid composite and tooth number were randomly selected by flipping a coin. This approach was first used for the patients with 2 restorations and the same randomized approach was then used to select the nanofill and microhybrid resin composites, and tooth number, respectively, for the patients with more than 2 restorations. After randomization, the number of restorations with Filtek Supreme XT per patient was equal to the number of Filtek Z250 restorations.

Cavity preparation was performed only for removal of caries and the cavity margins were not beveled. From each occlusal carious lesion, unsupported enamel rods were removed with minimal invasion using a cylindrical diamond bur (Komet, Gebr. Brasseler, Lemgo, Germany) in a high-speed handpiece under water cooling until the carious dentin was exposed. All of the cavities were prepared using round and cylindrical tungsten carbide burs (Komet, Gebr. Brasseler) in a low-speed contra-angle handpiece (maximum speed of 1,500 rpm) under air/water spray. Caries removal was ended, when a hard cavity floor was felt upon gentle pressure with a blunt dental explorer [11]. Cavity margins were not in occlusal contact. The cavities were categorized by depth as shallow (49 cavities), medium (31 cavities), or deep (38 cavities). The depth of the prepared cavity was measured against the mesial and distal marginal ridges using a graduated periodontal probe. According to this, > 2.5 mm cavities were classified as deep cavities, while < 2.5 mm as medium cavities and the dentin that neighbors the enamel as shallow cavities [12]. The average facio-lingual width of the cavities was between 1/3 and 2/3 of the intercuspal width. In deep cavities, a small amount of calcium hydroxide (Dycal, DeTrey/Dentsply, Konstanz, Germany) was placed on the deep portion of the cavity and then cavity floor was lined using a resin-modified glass ionomer cement (Vitrebond, 3M ESPE). Cavities considered as medium or shallow depth did not require lining. Cavity treatment and material application was performed according to the

manufacturer's instructions by the same experienced practitioner (ST) who was familiar with both materials.

The shade was chosen using the corresponding guide or custom samples before isolation. Cavities were isolated using cotton rolls and saliva ejectors [13]. Scotchbond Multi-Purpose was applied to etched enamel and dentin in accordance with manufacturer's instructions (Table 1). The adhesive was light cured for 20 seconds using a halogen light curing unit (VIP, Bisco Inc., Schaumburg, IL, USA).

Either a microhybrid composite (Filtek Z250) or a nanofill resin composite (Filtek Supreme XT) was then applied in layers (maximum 2 mm) using an oblique incremental placement technique. Each increment of both composites was cured for 20 seconds. Halogen light intensity was checked with a radiometer (Hilux Curing Light Meter, Dental Benlioglu Inc., Ankara, Turkey) prior to and after curing to ensure that the output was at least 600 mW/cm<sup>2</sup>. After checking the occlusion and articulation, removal of excess material, contouring, and finishing were performed using microfine finishing diamonds ranging from fine to superfine (8368.204.023, Komet, Gebr. Brasseler). The restorations were completed by polishing using Sof-Lex abrasive disks (3M ESPE). After that, the patients received orthodontic treatment with fixed orthodontic appliances.

### Evaluations

Two experienced calibrated examiners from the Department of Restorative Dentistry of Istanbul University evaluated the restorations using a dental explorer and mirror, in accordance with the modified United States Public Health Service (USPHS) criteria (Tables 3 and 4) [13-16]. The evaluations were made 1 week after restorations placement (baseline)

**Table 3.** Direct clinical evaluation criteria (modified USPHS criteria) using visual inspection

Criterion	Rating	Aspect
Color match	Alpha	There is no a mismatch in color, shade and/or translucency between the restoration and the adjacent tooth structure.
	Bravo	There is a mismatch in color, shade and/or translucency between the restoration and the adjacent tooth structure, but the mismatch is within the normal range of tooth color, shade and/or translucency.
	Charlie	The mismatch is between the restoration and the adjacent tooth structure outside the normal range of tooth color, shade and/or translucency.
Cavosurface marginal discoloration	Alpha	There is no discoloration anywhere on the margin between the restoration and the tooth structure.
	Bravo	There is discoloration anywhere on the margin between the restoration and the tooth structure, but the discoloration has not penetrated along the margin of the restorative material in an enamel direction and can be polished away.
	Charlie	The discoloration has penetrated along the margin of the restorative material in an enamel direction.
Wear/anatomic form	Alpha	The restoration is not under-contoured, that is, the restorative material is not discontinuous with existing anatomic form.
	Bravo	The restoration is under-contoured, that is, the restorative material is discontinuous with existing anatomic form, but sufficient restorative material is not missing so as to expose the enamel or base.
	Charlie	Sufficient restorative material is missing so as to expose the enamel or base.
Caries	Alpha	There is no evidence of caries contiguous with the margin of the restoration.
	Bravo	There is evidence of caries contiguous with the margin of the restoration.

USPHS, United States Public Health Service.

**Table 4.** Direct clinical evaluation criteria (modified Ryge criteria) using explorer

Criterion	Rating	Aspect
Marginal adaptation	Alpha	There is no visible evidence of a crevice along the margin into which the explorer will penetrate.
	Bravo	There is visible evidence of a crevice along the margin into which the explorer will penetrate. The enamel or base is not exposed.
	Charlie	There is visible evidence of a crevice along the margin into which the explorer will penetrate. The enamel or base is exposed.
	Delta	The restoration is fractured or missing in part or <i>in toto</i> .
Surface texture	Alpha	Surface of restoration is smooth.
	Bravo	Surface of restoration is slightly rough or pitted, can be refinished.
	Charlie	Surface deeply pitted, irregular grooves (not related to anatomy), cannot be refinished.
	Delta	Surface is fractured or flaking.

and after 1, 2, 3, 4, and 5 years. The examiners were not involved in the restoration phase and were fully blinded to the experimental protocol. The 2 examiners studied a set of reference photographs to illustrate each score for each criterion. They then clinically evaluated 20 occlusal restorations together at 2 day intervals. The restorations are not presented in the current study. When a minimum 85% intra- and inter-examiner agreement was achieved in the calibration phase, the evaluation phase was commenced [16]. At baseline and 1, 2, 3, 4, and 5-year recalls, color match, wear or loss of form, discoloration, caries formation, marginal adaptation, and surface texture were evaluated and scored as Alpha, which represented perfect clinical form; Bravo, acceptable; Charlie, unacceptable and the restoration had to be replaced; Delta, which represented restoration fracture, mobility of the restoration, or the restoration was missing and required immediate replacement. Scoring conflicts were resolved through consensus.

### Statistical analysis

All analyses were performed using SPSS for Windows version 20.0 (SPSS, Chicago, IL, USA). Data obtained from each composite were statistically analyzed using the Friedman test to examine changes that occurred during the 5 years evaluation period. When a statistically significant difference was identified for any criterion assessed, the Dunn test was used for multiple comparisons between each recall time for each composite. The Mann-Whitney test was used to evaluate the differences between the 2 different composite materials. Kaplan-Meier survival analysis was used to determine the probability of the clinical survival of the 2 composites for a given time period. A *p* value less than 0.05 was considered statistically significant. Cohen's kappa was used to check for inter- and intra-examiner agreement.

## RESULTS

The duration of orthodontic treatments ranged between 14 and 48 months (mean: 29.7 months). The distribution of occlusal restorations according to composite material types and teeth numbers are presented in **Table 5**. For the lower first molars, a few number of occlusal restorations were done. However, 69.5% of the restorations were done on the second molars. All patients attended the 1, 2, 3, 4, and 5-year recall visits, which resulted in a 100% recall rate for each evaluation period (**Figure 1** and **Table 6**). Cohen's kappa (0.87) exhibited strong agreement between the examiners and no statistically significant difference was found between their evaluations ( $p > 0.05$ ). The cumulative failure and retention (success) rates according to the Kaplan-Meier survival analysis are provided in **Table 6**. At each of the 5-year recalls, none of the restorations had failed, resulting in a 100% success rate for each evaluation period and a 100% cumulative success rate at the end of 5 years. Friedman test showed no significant differences between all of the evaluation periods of Filtek Z250 and Filtek Supreme XT restorations with respect to color match, marginal discoloration, wear or loss of anatomic form, caries, marginal adaptation, and surface texture.

Direct clinical evaluation results at baseline, and at the 1, 2, 3, 4, and 5-year recalls are shown

**Table 5.** Distribution of occlusal restorations according to composite material type and tooth number

Composite/dentin adhesive	No.	Tooth number							
		#16	#17	#26	#27	#36	#37	#46	#47
Filtek Z250/Scotchbond Multi-Purpose	59	6	8	7	14	2	10	2	10
Filtek Supreme XT/Scotchbond Multi-Purpose	59	6	11	5	11	2	9	6	9
Sum of restorations	118	12	19	12	25	4	19	8	19

**Table 6.** Results of clinical evaluation of 2 different composite restorations using modified USPHS criteria

Observation time/composite	Recall rate	Survival rate	Retention			Color match			Marginal discoloration			Wear/anatomic form			Caries			Marginal adaptation			Surface texture					
			A	C	A	B	C	A	B	C	A	B	C	A	B	C	A	B	C	A	B	C	D			
<b>Baseline</b>																										
Filtek Z250	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	-	-	
Filtek Supreme XT	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	-	-	100.0 (59)	-	-	98.3 (58)	1.7 (1)	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-
<b>1 year</b>																										
Filtek Z250	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	1.7 (1)	-	98.3 (58)	1.7 (1)	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	98.3 (58)	1.7 (1)	-	-	-	-
Filtek Supreme XT	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	1.7 (1)	-	98.3 (58)	1.7 (1)	-	98.3 (58)	1.7 (1)	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-
<b>2 year</b>																										
Filtek Z250	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	3.4 (2)	-	96.6 (57)	3.4 (2)	-	98.3 (58)	1.7 (1)	-	100.0 (59)	-	-	100.0 (59)	-	-	98.3 (58)	1.7 (1)	-	-	-	-
Filtek Supreme XT	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	3.4 (2)	-	96.6 (57)	3.4 (2)	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-
<b>3 year</b>																										
Filtek Z250	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	5.0 (3)	-	88.1 (52)	11.9 (7)	-	98.3 (58)	1.7 (1)	-	100.0 (59)	-	-	100.0 (59)	-	-	98.3 (58)	1.7 (1)	-	-	-	-
Filtek Supreme XT	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	5.0 (3)	-	89.8 (53)	10.2 (6)	-	96.6 (57)	3.4 (2)	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-	100.0 (59)	-	-
<b>4 year</b>																										
Filtek Z250	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	6.8 (4)	-	93.2 (55)	6.8 (4)	-	96.6 (57)	3.4 (2)	-	100.0 (59)	-	-	100.0 (59)	-	-	98.3 (58)	1.7 (1)	-	-	-	-
Filtek Supreme XT	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	10.2 (6)	-	89.8 (53)	10.2 (6)	-	95.0 (56)	5.0 (3)	-	100.0 (59)	-	-	100.0 (59)	-	-	98.3 (58)	1.7 (1)	-	-	-	-
<b>5 year</b>																										
Filtek Z250	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	15.3 (9)	-	84.7 (50)	15.3 (9)	-	96.6 (57)	3.4 (2)	-	100.0 (59)	-	-	100.0 (59)	-	-	98.3 (58)	1.7 (1)	-	-	-	-
Filtek Supreme XT	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	100.0 (59)	11.9 (7)	-	88.1 (52)	11.9 (7)	-	95.0 (56)	5.0 (3)	-	100.0 (59)	-	-	100.0 (59)	-	-	96.6 (57)	3.4 (2)	-	-	-	-

Five-year data of recall rate, survival rate, retention, color match, marginal discoloration, wear or loss of anatomic form, caries, marginal adaptation, and surface texture showed no statistically significant difference ( $p > 0.05$ ).

USPHS, United States Public Health Service; A, Alpha; B, Bravo; C, Charlie; D, Delta.

in **Table 6**. None of the restorations were clinically unacceptable with regard to any evaluation criteria. Mann-Whitney test revealed there were no statistically significant differences in any of the clinical evaluation criteria between Filtek Z250 and Filtek Supreme XT restorations in each evaluation period. Regarding the color match, at the 5-year recall, 84.7% and 88.1% of the Filtek Z250 and Filtek Supreme XT restorations, respectively, were clinically ideal (Alpha). After 5 years, 28.8% of Filtek Z250 and 18.6% of Filtek Supreme XT restorations exhibited clinically acceptable marginal discoloration (Bravo). However, this discoloration was superficial, located on anywhere along the margin, did not penetrate towards the pulp along the margin of the restorative material, and could be polished away. With respect to wear and anatomic form, 96.6% of Filtek Z250 and 95% of Filtek Supreme XT restorations were clinically ideal (Alpha) after 5 years. Regarding the marginal adaptation rates, 98.3% of Filtek Z250 and 96.6% of Filtek Supreme XT restorations were clinically ideal (Alpha) at the 5-year recall. After 5 years, 98.3% of Filtek Z250 and 100% of Filtek Supreme XT restorations were clinically ideal (Alpha) with respect to surface texture.

## DISCUSSION

In the present study, 2 types of resin composites were used in patients who were going to commence orthodontic treatment. Filtek Supreme XT nanofill resin composite contains nanometer sized particles (1–100 nm) throughout the resin matrix. Larger primary particles are not present. In microhybrid resin composite (Filtek Z250), the fine particles with an average particle size of 0.6  $\mu\text{m}$  are blended with microfine silica [17,18]. The 5-year survival rates of Filtek Z250 and Filtek Supreme XT restorations were 100%. No restorations failed, which meant an overall success rate of 100%. The change in performance between Filtek Z250 and Filtek Supreme XT restorations was only from clinically ideal (Alpha) to a clinically acceptable (Bravo) (not significant, Mann-Whitney test). In agreement with this finding, a nanofilled composite (Filtek Supreme XT) exhibited a success rate of 100% in occlusal and posterior approximal restorations after 5 years [19].

In contrast, a greater failure rate (6.4%) was reported for nanofill (Filtek Z350) and microhybrid (Filtek Z250) composites after 54 months in occlusal restorations [7]. However, in a 10-year retrospective study that investigated the longevity of posterior approximal restorations using 4 similar microhybrid resin composites, the overall survival rate was 97.9% and an improved performance was observed with Filtek Z250 restorations (99.1%) [20]. In contrast to this study, the present study evaluated 2 composites in occlusal cavities for 5 years. Cavity size played an important role on composite restoration survival. When compared with 1 surface restorations, the relative risk of failure has been reported to be approximately 2.3 times greater for 2 surface restorations and 3.3 times greater for multi-surface restorations [21]. It has also been documented that a reduction in cavity size protected the restoration from chewing forces [22]. Therefore, the size and type of cavity may have contributed to the high success rates in the present study. In accordance with this assumption, a review of 34 clinical studies performed over periods of at least 5 years found that 90% reported an annual failure rate between 1% and 3% for occlusal and posterior approximal composite restorations [23]. The variation in failure rates were dependent on several factors including tooth type and location, operator, and socioeconomic, demographic, and behavioral elements [23].

In the current study, 9 (15.3%) Filtek Z250 restorations and 7 (11.9%) Filtek Supreme XT restorations showed a change in color match after 5 years. This color change was clinically



acceptable (Bravo) and did not necessitate restoration replacement. In contrast, another study reported no color change for a nanofill composite (Filtek Supreme XT) after 5 years in occlusal and posterior approximal restorations [19]. However, in support of the findings from the present study, 12.5% of microhybrid (Filtek Z250) and 6.5% of nanofill (Filtek Z350) composite restorations demonstrated acceptable color change after 54 months in occlusal cavities [7]. Staining would probably require a longer duration due to the transient nature of staining, and because saliva and other fluids dilute substances that stain and restorations are regularly cleaned through brushing. Therefore, color change may be due to pigment absorption or to dietary and oral hygiene habits [24].

In regards to marginal adaptation, 1.7% of Filtek Z250 and 3.4% of Filtek Supreme XT restorations exhibited Bravo scores, which showed visible evidence of a crevice along the margin into which the explorer could penetrate. Marginal discoloration also recorded Bravo scores in 28.8% of Filtek Z250 and 18.6% of Filtek Supreme XT restorations after 5 years. Most of Bravo ratings were shown in deep cavities. Deep cavities (39.4%) showed marginal discoloration and this was higher than medium (19.4%) and shallow (14.3%) cavities. Compared with the present study, a lower proportion of marginal discoloration and a reduced marginal deterioration rate (Bravo) have been reported after 54 months and after 5 years [7,19]. In contrast, a 10 year retrospective study of posterior composites demonstrated that marginal quality decreased with time because of physiologic and chemical reactions in the oral cavity, and degradation might indicate issues related to adhesive or resin composite [20].

After 5 years, only 3.4% of Filtek Z250 and 5% of Filtek Supreme XT restorations showed clinically acceptable (Bravo) wear or anatomic form. However, in a previous study, anatomic form deficiencies were recorded as Charlie in 6.5% of microhybrid (Filtek Z250) and nanofill (Filtek Z350) composites after 54 months [7]. However, this wear was more frequent and severe (Charlie) than in the present study. In contrast, another study reported that 2.2% and 2.7% of the posterior and hybrid composite restorations, respectively, were clinically acceptable (Bravo) anatomically after 4 years in occlusal and posterior approximal cavities [25]. It was indicated that anatomic form deficiencies in cases of 3 surface posterior restorations were greater than in 2 surface restorations, independent of the materials. This was accounted for by the greater resin composite surface, which wore after abrasive attack and led to material loss [20]. Therefore, in the present study, size and cavity type may have been contributing factors to the lower wear rates. In support of this, small-to-moderate sized posterior composite restorations were reported to have been used successfully for up to 20 years [26]. Seventy-six percent of the 85 restorations in their study were recalled after 17 years and were clinically acceptable. In addition, the effect of impact wear was stated to be limited owing to the moderate-to-narrow width of the restorations, which guaranteed that there was occlusal contact on tooth structure in almost every case [26].

In the present study, after 5 years only 1.7% of Filtek Z250 restorations was slightly pitted and exhibited rough surfaces (Bravo) that could be restored by polishing. In contrast, all Filtek Supreme XT restorations were clinically ideal (Alpha) with regards to surface texture. However, the incidence of clinically acceptable (Bravo) surface texture was previously reported greater for microhybrid and nanofill composites after 54 months and 5 years [7,19]. In agreement with our study, 97.4% of Filtek Z250 restorations showed no change and were clinically ideal in respect to surface roughness after 10 years [20]. In addition, 3 types of universal composites have been shown to exhibit a very high incidence of clinically ideal surface luster after 20 years [27]. Two previous studies also reported similar results after 17

and 22 years for posterior composites [21,28]. Furthermore, in accordance with the findings of the present study, it was reported that fine-hybrid and nanohybrid composites exhibited ideal surface texture for all restorations (100%) in occlusal and posterior approximal cavities after 4 years [29]. Modern particulate resin composites have smooth surface characteristics after polishing [10]; therefore, reduced particle dimension and increased filler loading may have contributed to the higher incidences of clinically ideal surface texture in the present study. In addition, size and type of cavity may be factors that contribute to improvements in surface texture.

## CONCLUSIONS

After 5 years, none of the microhybrid (Filtek Z250) and nanofill (Filtek Supreme XT) composites failed; the success rate was 100% for both composite materials. The composite restorations were clinically acceptable for all parameters at the end of the 5 years evaluation period. However, acceptable restoration rates (Bravo) were higher for marginal discoloration, followed by color match, compared with the other evaluation criteria. Modern composites such as the microhybrid and nanofill composites can result in high-quality restorations and produce positive long-term outcomes in occlusal cavities. In addition, orthodontic treatment did not affect clinical performance of 2 different composite restorations.

## REFERENCES

1. Uribe FA, Chandhoke TK, Nanda R. Individualized orthodontic diagnosis. In: Nanda R, editor. *Esthetics and biomechanics in orthodontics*. 2nd ed. St. Louis (MA): Elsevier Saunders; 2015. p1-32.
2. Bourzgui F, Sebbar M, Hamza M. Orthodontics and caries. In: Naretto S, editor. *Principles in contemporary orthodontics*. Rijeka: INTECH; 2011. p309-326.
3. Chaussain C, Opsahl Vital S, Viallon V, Vermelin L, Haignere C, Sixou M, Lasfargues JJ. Interest in a new test for caries risk in adolescents undergoing orthodontic treatment. *Clin Oral Investig* 2010;14:177-185. [PUBMED](#) | [CROSSREF](#)
4. Lynch CD, Opdam NJ, Hickel R, Brunton PA, Gurgan S, Kakaboura A, Shearer AC, Vanherle G, Wilson NHAcademy of Operative Dentistry European Section. Guidance on posterior resin composites: Academy of Operative Dentistry - European Section. *J Dent* 2014;42:377-383. [PUBMED](#) | [CROSSREF](#)
5. Ferracane JL. Current trends in dental composites. *Crit Rev Oral Biol Med* 1995;6:302-318. [PUBMED](#) | [CROSSREF](#)
6. Ilie N, Hickel R. Resin composite restorative materials. *Aust Dent J* 2011;56 Supplement 1:59-66. [PUBMED](#) | [CROSSREF](#)
7. de Andrade AK, Duarte RM, Medeiros e Silva FD, Batista AU, Lima KC, Monteiro GQ, Montes MA. Resin composite class I restorations: a 54-month randomized clinical trial. *Oper Dent* 2014;39:588-594. [PUBMED](#) | [CROSSREF](#)
8. Ferracane JL. Resin composite--state of the art. *Dent Mater* 2011;27:29-38. [PUBMED](#) | [CROSSREF](#)
9. Gresnigt MM, Kalk W, Ozcan M. Randomized controlled split-mouth clinical trial of direct laminate veneers with two micro-hybrid resin composites. *J Dent* 2012;40:766-775. [PUBMED](#) | [CROSSREF](#)
10. Wolff D, Kraus T, Schach C, Pritsch M, Mente J, Staehle HJ, Ding P. Recontouring teeth and closing diastemas with direct composite buildups: a clinical evaluation of survival and quality parameters. *J Dent* 2010;38:1001-1009. [PUBMED](#) | [CROSSREF](#)
11. Neves Ade A, Coutinho E, De Munck J, Van Meerbeek B. Caries-removal effectiveness and minimal-invasiveness potential of caries-excitation techniques: a micro-CT investigation. *J Dent* 2011;39:154-162. [PUBMED](#) | [CROSSREF](#)

12. de Souza Costa CA, Teixeira HM, Lopes do Nascimento AB, Hebling J. Biocompatibility of resin-based dental materials applied as liners in deep cavities prepared in human teeth. *J Biomed Mater Res B Appl Biomater* 2007;81:175-184.  
[PUBMED](#) | [CROSSREF](#)
13. Demirci M, Uysal O. Clinical evaluation of a polyacid-modified resin composite (Dyract AP) in Class I cavities: 3-year results. *Am J Dent* 2006;19:376-381.  
[PUBMED](#)
14. Barnes DM, Blank LW, Gingell JC, Gilner PP. A clinical evaluation of a resin-modified glass ionomer restorative material. *J Am Dent Assoc* 1995;126:1245-1253.  
[PUBMED](#) | [CROSSREF](#)
15. Ryge G. Clinical criteria. *Int Dent J* 1980;30:347-358.  
[PUBMED](#)
16. Cvar JF, Ryge G. Reprint of criteria for the clinical evaluation of dental restorative materials. 1971. *Clin Oral Investig* 2005;9:215-232.  
[PUBMED](#) | [CROSSREF](#)
17. Sakaguchi RL, Powers JM. *Craig's restorative dental materials*. 13th ed. Philadelphia (PA): Elsevier Mosby; 2012. p166-167.
18. 3M Dental Products Laboratory (US). *Filtek™ Z250 universal restorative system: technical product profile*. St. Paul (MN): 3M; 1998.
19. Cetin AR, Unlu N, Cobanoglu N. A five-year clinical evaluation of direct nanofilled and indirect composite resin restorations in posterior teeth. *Oper Dent* 2013;38:E1-E11.  
[PUBMED](#) | [CROSSREF](#)
20. Lempel E, Tóth Á, Fábrián T, Krajczár K, Szalma J. Retrospective evaluation of posterior direct composite restorations: 10-year findings. *Dent Mater* 2015;31:115-122.  
[PUBMED](#) | [CROSSREF](#)
21. da Rosa Rodolpho PA, Cenci MS, Donassollo TA, Loguercio AD, Demarco FF. A clinical evaluation of posterior composite restorations: 17-year findings. *J Dent* 2006;34:427-435.  
[PUBMED](#) | [CROSSREF](#)
22. Manhart J, Chen H, Hamm G, Hickel R. Buonocore Memorial Lecture. Review of the clinical survival of direct and indirect restorations in posterior teeth of the permanent dentition. *Oper Dent* 2004;29:481-508.  
[PUBMED](#)
23. Demarco FF, Corrêa MB, Cenci MS, Moraes RR, Opdam NJ. Longevity of posterior composite restorations: not only a matter of materials. *Dent Mater* 2012;28:87-101.  
[PUBMED](#) | [CROSSREF](#)
24. Bagheri R, Burrow MF, Tyas M. Influence of food-simulating solutions and surface finish on susceptibility to staining of aesthetic restorative materials. *J Dent* 2005;33:389-398.  
[PUBMED](#) | [CROSSREF](#)
25. Manhart J, Chen HY, Hickel R. Clinical evaluation of the posterior composite Quixfil in class I and II cavities: 4-year follow-up of a randomized controlled trial. *J Adhes Dent* 2010;12:237-243.  
[PUBMED](#)
26. Wilder AD Jr, May KN Jr, Bayne SC, Taylor DF, Leinfelder KF. Seventeen-year clinical study of ultraviolet-cured posterior composite Class I and II restorations. *J Esthet Dent* 1999;11:135-142.  
[PUBMED](#) | [CROSSREF](#)
27. Baldissera RA, Corrêa MB, Schuch HS, Collares K, Nascimento GG, Jardim PS, Moraes RR, Opdam NJ, Demarco FF. Are there universal restorative composites for anterior and posterior teeth? *J Dent* 2013;41:1027-1035.  
[PUBMED](#) | [CROSSREF](#)
28. Da Rosa Rodolpho PA, Donassollo TA, Cenci MS, Loguercio AD, Moraes RR, Bronkhorst EM, Opdam NJ, Demarco FF. 22-Year clinical evaluation of the performance of two posterior composites with different filler characteristics. *Dent Mater* 2011;27:955-963.  
[PUBMED](#) | [CROSSREF](#)
29. Schirrmeyer JE, Huber K, Hellwig E, Hahn P. Four-year evaluation of a resin composite including nanofillers in posterior cavities. *J Adhes Dent* 2009;11:399-404.  
[PUBMED](#)