Influence of light-exposure methods and depths of cavity on the microhardness of dual-cured core build-up resin composites

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ABSTRACT

bjective: The purpose of this study was to evaluate the Knoop hardness number (KHN) of dual-cured core build-up resin composites (DCBRCs) at 6 depths of cavity after 3 post-irradiation times by 4 light-exposure methods. Material and Methods: Five specimens each of DCBRCs (Clearfil DC Core Plus [DCP] and Unifil Core EM [UCE]) were filled in acrylic resin blocks with a semi-cylindrical cavity and light-cured using an LED light unit (power density: 1,000 mW/cm²) at the top surface by irradiation for 20 seconds (20 s), 40 seconds (40 s), bonding agent plus 20 seconds (B+20 s), or 40 seconds plus light irradiation of both sides of each acrylic resin block for 40 seconds each (120 s). KHN was measured at depths of 0.5, 2.0, 4.0, 6.0, 8.0, and 10.0 mm at 0.5 hours, 24 hours, and 7 days post-irradiation. Statistical analysis was performed using repeated measures ANOVA and Tukey's compromise post-hoc test with a significance level of p<0.05. Results: For both DCBRCs, at 0.5 hours post-irradiation, the 20 s and 40 s methods showed the highest KHN at depth of 0.5 mm. The 40 s method showed significantly higher KHN than the 20 s method at all depths of cavity and post-irradiation times, except UCE at depth of 0.5 mm (p<0.05). The 120 s method did not result in significantly different KHN at all depths of cavity and post-irradiation times (p>0.05). In DCP, and not UCE, at 24 hours and 7 days post-irradiation, the B+20 s method showed significantly higher KHN at all depths of cavity, except the depth of 0.5 mm (p<0.05). Conclusion: KHN depends on the light-exposure method, use of bonding agent, depth of cavity, post-irradiation time, and material brand. Based on the microhardness behavior, DCBRCs are preferably prepared by the effective exposure method, when used for a greater depth of cavity.

Keywords: Composite resins. Exposure time. Hardness.

INTRODUCTION

Core build-up materials are often required to reconstruct and provide an ideal morphology to severely damaged teeth prior to their preparation for indirect foundation restorations. Despite substantial documented evidence of the long-term success of large amalgam restorations^{26,40}, resin composites, since the early days of self-cured materials, have also been used for this purpose. More recently, light-cured core build-up materials that are more convenient to use than chemically cured composites have been widely used³³. Both these materials,

however, have their disadvantages^{12,29}. While chemically cured materials do not allow clinicians to adjust the setting time individually, light-cured resin composites do not ensure adequate polymerization in areas with limited access to the curing light.

However, resin-based composites are associated with polymerization shrinkage, causing stress development under confined conditions³⁶. Several strategies have been used to overcome the limitations of polymerization shrinkage. Various layering techniques have been suggested to minimize shrinkage stress^{20,23}. Nevertheless, time limitations when placing core build-up materials

restrict clinicians from using elaborate multilayering techniques. Therefore, dual-cured resin composites, in which polymerization is chemically initiated in the deeper portions of the canal or preparations, have been developed for use as core build-up materials; this has allowed clinicians to build extended foundation restorations quickly, and in bulk.

At the coronal region, dual-cured core build-up resin composites are mainly polymerized through photo-initiated reactions, whereas, in the apical region, polymerization is chemically initiated. However, the incorporation of self- and light-curing modes in the same material does not ensure maximal curing of the material. Due to incomplete compensation for deficient light activation, lower hardness values of dual-cured core build-up resin composites have been observed with increased depth of cavity¹. It has been speculated that a delay in light activation would be beneficial in enhancing the degree of conversion of dual-cured resin cements, since immediate exposure to light could interfere with the chemical-curing mechanism²⁴. On the other hand, it has been reported that time delay and duration of light exposure does not increase microhardness at different depths of a dual-cured core build-up resin composite 2 weeks after light irradiation³⁹. Moreover, light-activation delayed by 5 minutes after seating the fiber-reinforced post did not affect the microhardness of dual-cured resin cements at 3 regions of the root after 3 months of storage in water²⁸. Although dual-cured core build-up resin composites have been recently used to prepare prefabricated posts and core or coronalradicular build-ups, their hardness behavior at greater depths of cavity is unknown. Moreover, no information is available in the literature regarding the effect of applying a bonding agent on the cavity wall on polymerization of the dual-cured core buildup resin composite.

Based on these considerations, the purpose of the present study was to investigate the influence of light-exposure durations (20, 40, and 120 seconds) and application of a bonding agent on the extent of polymerization by measuring the microhardness of 2 dual-cured core build-up resin composites at different depths of cavity without prefabricated posts and various post-irradiation times. The following null hypotheses were tested: (1) an increase in light-exposure time results in no difference in hardness, regardless of the depth of cavity and type of dual-cured resin composite; (2) hardness is not affected by depth of cavity and post-irradiation time; and (3) application of the bonding agent does not improve hardness.

MATERIAL AND METHODS

Specimen preparation

Forty semi-cylindrical cavities with a diameter of 3 mm and a depth of 11 mm were prepared in 5x10x16 mm acrylic resin blocks (Figure 1). Two acrylic resin blocks, with or without a semicylindrical cavity, were placed in a silicon impression material mold (15x15x20 mm). Either 1 of 2 dualcured core build-up resin composite pastes [Clearfil DC Core Plus (DCP) or Unifil Core EM (UCE) (Figure 2)] was filled directly in the cavities using automixing tips, being sure to avoid entrapment of air, according to the manufacturer's instructions. The upper surface in the resin composite material was covered with a plastic strip and pressed with a thin cover glass to remove any excess resin. Light-irradiation was provided by placing the tip of the LED light unit (power density: 1,000 mW/ cm²; Pencure, J. Morita MFG Corp., Kyoto, Japan) on the plastic strip. Power output was verified with a curing radiometer (Cure Lite; Dentsply Caulk, Milford, CT, USA) immediately before every lightactivation throughout the study. The core build-up resin composites were light-cured for different durations using 1 of the 4 following light-exposure methods: (1) light-irradiation for 20 seconds on the plastic strip after filling the core build-up resin composite (20 s method); (2) light-irradiation for 40 seconds (40 s); (3) application of the bonding agent on the cavity wall with a brush, followed by air-drying, and light-irradiation for 10 seconds on the top of the cavity before the 20 s method (B+20 s method); (4) light-irradiation for 40 seconds on the plastic strip, followed by removal of the acrylic blocks from the silicon impression material mold, and irradiation of both sides of each acrylic resin block for 40 seconds each (120 s method). After irradiation of all the specimens, the acrylic resin blocks were removed from the silicon mold and separated. Five specimens of each dual-cured core build-up resin composite were irradiated by each of the 4 methods.

Hardness measurements

Hardness was measured at the following depths from the light-irradiated surface of the cavity: 0.5, 2.0, 4.0, 6.0, 8.0, and 10.0 mm. For each specimen, the Knoop hardness number (KHN) was measured 5 times at each depth using a microhardness tester (FM-700: Future-Tech Corp., Kawasaki, Japan) at 0.5 hours, 24 hours, and 7 days post-irradiation. A Knoop diamond indenter was applied under a load of 25 g for a dwell time of 15 seconds; the load was then removed, and the long diagonal of the indentation was measured under 400x magnification. KHN, which is inversely proportional to the square of the long diagonal,

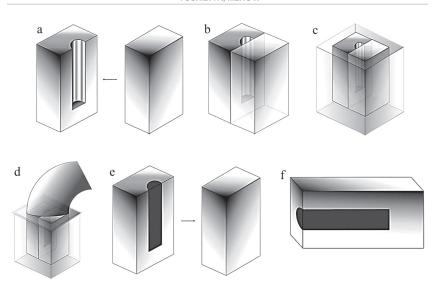


Figure 1- Schematic illustration of the preparation of specimens for measurement of Knoop hardness: a: acrylic resin blocks (5x10x16 mm) with or without semi-cylindrical cavity (diameter of 3 mm and depth of 11 mm); b: two acrylic resin blocks joined; c: two acrylic resin blocks placed in a silicon impression material mold (15x15x20 mm); d: dual-cured core build-up resin composite filled in the cavity and irradiated on the upper surface using LED light unit; e: two acrylic resin blocks separated; f: Knoop hardness measurement for the surface of resin composite filled in the semi-cylindrical cavity

Materials	Manufacturer	Composition			
Core material					
Clearfil DC Core Plus (DCP)	Kuraray Noritake Products	A paste: Bis-GMA, hydrophilic aliphatic dimethacrylate,			
(Batch # 0002AA)	Corp., Tokyo, Japan	hydrophobic aliphatic dimethacrylate,hydrophobic			
		aromatic dimethacrylate, silanized barium glass filler,			
		silanized colloidal silica, colloidal silica, chemical-initiator,			
		photo-initiator, pigments			
		B paste: TEGDMA, hydrophilic aliphatic dimethacrylate,			
		hydrophobic aromatic dimethacrylate, silanized barium			
		glass filler, silanized colloidal silica, aluminum oxide filler,			
		photo-accelerator , chemical-accelerator			
Bonding material					
Clearfil S ³ Bond Plus	Kuraray Noritake Products	Bis-GMA,HEMA, MDP, hydrophilic dimethacrylate,			
(Batch # 00024B)	Corp., Tokyo, Japan	hydrophobic dimethacrylate, colloidal silica, ethanol,			
		water, photo-initiators, photo- and chemical-accelerator,			
		sodium fluoride			
Core material					
Unifil Core EM (UCE)	GC Corp., Tokyo, Japan	Base: UDMA, dimethacrylates, fluoro-alumino-silicate			
(Batch # 1107011)		glass, silicon dioxide, photo-initiator, accelerator			
		Catalyst: UDMA, dimethacrylates, fluoro-alumino-silicate			
		glass, silicon dioxide, chemical-initiator, pigment			
Bonding material					
Self-etching bond A	GC Corp., Tokyo, Japan	dimethacrylates, 4-META, silicon dioxide, ethanol, water,			
(Batch # 1205181)		photo-initiator			
Self-etching bond B	GC Corp., Tokyo, Japan	ethanol, accelerator			
(Batch # 1206011)					

Bis-GMA: bis-phenol-A-glycidyldimethacrylate; TEGDMA: triethyleneglycol dimethacrylate; UDMA: urethane dimethacrylate; HEMA: 2-hydoxyethyl methacrylate; MDP: 10-methacryloyloxydecyl dihydrogen phosphate; 4-META: 4-methacryloyloxyethyl trimellitate anhydride

Figure 2- Dual-cured core build-up resin composites used in this study

was thus calculated. All specimens were stored under dry and dark conditions in a box, which was placed in a biochemical incubator at 37°C to avoid exposure to light, and was accessed only to obtain measurements.

Statistical analysis

The KHN data were statistically analyzed by the repeated measures three-way ANOVA test. The independent variables analyzed were depths of cavity and post-irradiation times for within-subject analysis, and light-exposure methods and type of dual-cured core build-up resin composite for between-subject analysis. A one-way ANOVA with the *post-hoc* Tukey's compromise test was used to establish specific differences in KHN values between the groups (α =0.05).

RESULTS

The results of this study are summarized in Tables 1 and 2, which show the mean KHN and standard deviation of all the experimental groups in DCP and UCE, respectively, at 6 depths of cavity after 3 different post-irradiation durations by 4 different light-exposure methods.

For the DCP, the KHN was effected by the light-exposure method (p=0.0001; F=1652.29), post-irradiation time (p=0.0001; F=606.09),

depth of cavity (p=0.0001; F=1184.99), and all the interactions between all the aforementioned factors (p=0.0001). For UCE, significant differences were found between the light-exposure methods (p=0.0001; F=201.27), post-irradiation times (p=0.0001; F=857.15), and depth of cavity (p=0.0001; F=488.76); in addition, significant interactions were also found between all the aforementioned factors (p=0.0001).

For both resin composites, at 0.5 hours post-irradiation, the 20 s and 40 s methods resulted in the highest KHN values at the depth of 0.5 mm; these values gradually decreased with increasing depths of cavity (p<0.05). On the other hand, at 24 hours or 7 days post-irradiation, the KHN values of DCP or UCE were not significantly different between the depths of 2.0 mm, 4.0 mm, and/or 6.0 mm, or 8.0 mm and 10.0 mm, respectively (p>0.05).

The 40 s method resulted in significantly higher KHN values than the 20 s method at all depths of cavity and post-irradiation times for both resin composites, with the exception of UCE at the depth of 0.5 mm (p<0.05). With the 120 s method, the KHN values of both resin composites were not significantly different among the 6 depths of cavity at all post-irradiation times (p>0.05), but they were significantly higher than those of the 40 s method, with the exception of that at the depth of 0.5 mm at all post-irradiation times (p<0.05).

Table 1- Mean KHN±SD for Clearfil DC Core Plus (DCP) at 6 depths of cavity after 3 post-irradiation times by 4 light-exposure methods

The same lower-case superscript Greek characters indicate no statistically significant differences between exposure methods at same post-irradiation time in the same depth of cavity; the same lower-case superscript letters indicate no statistically significant differences between depths of cavity at same post-irradiation time in the same exposure method; the same upper-case superscript letters indicate no statistically significant differences between post-irradiation times at same depth of cavity in the same exposure method (p>0.05)

KHN=Knoop hardness number

Exposure method	Post- irradiation time	Depth of cavity (mm)					
		0,5	2	4	6	8	10
Exposure time of 20 s (20 s)	0.5 hours	43.48±0.56 ^{αaC}	39.89±0.92 ^{αbC}	37.56±0.40 ^{αcC}	35.89±0.73 ^{αcdC}	34.88±1.27 ^{αdC}	33.08±1.71 ^{αeC}
	24 hours	45.76±0.74 ^{αaB}	41.13±0.68 ^{αbB}	39.39±0.32 ^{αcB}	39.19±0.31αCAB	38.67±0.21αCAB	38.59±0.12 ^{αcAB}
	7 days	46.79±0.59 ^{αaA}	42.94±0.59 ^{αbA}	41.62±0.28αbA	40.03±1.14 ^{αcA}	39.62±1.28 ^{αcA}	39.04±1.07 ^{αcA}
Bonding + exposure time of 20 s (B+20 s)	0.5 hours	$41.71 \pm 0.54^{\beta aC}$	38.63±0.60 ^{αbC}	37.67±0.58αbcC	37.07±0.58αcC	$35.49{\pm}0.98^{\alpha dB}$	$35.01 \pm 0.90^{\alpha dB}$
	24 hours	46.11±0.59 ^{αaB}	43.18±0.75 ^{βbB}	41.84±0.62 ^{βcB}	41.70±0.49 ^{βcB}	41.34±0.82 ^{βcA}	41.17±0.84 ^{βcA}
	7 days	47.50±0.25 ^{αaA}	$45.51 {\pm} 0.95^{\beta bA}$	$43.34{\pm}0.83^{\beta\text{cA}}$	$43.35{\pm}0.88^{\beta\text{cA}}$	$42.54{\pm}0.70^{\beta cA}$	$42.24{\pm}0.90^{\beta cA}$
Exposure time of 40 s (40 s)	0.5 hours	48.83±0.36 ^{γaB}	44.42±0.33 ^{βbC}	42.83±0.50 ^{βcC}	40.83±0.87 ^{βdC}	38.60±0.58 ^{βeC}	$37.57 \pm 0.29^{\beta fC}$
	24 hours	51.89±0.61 ^{βaA}	49.04±0.66YbAB	47.57±0.57 ^{ycAB}	46.73±0.46 ^{ycAB}	44.76±0.59 ^{ydAB}	43.98±0.58 ^{ydAB}
	7 days	52.30±0.53 ^{βaA}	49.47±0.52 ^{ybA}	47.92±0.76 ^{ycA}	47.03±0.51 ^{ycA}	45.36±0.68 ^{YdA}	44.76±0.77 ^{vdA}
Exposure time of 120 s (120 s)	0.5 hours	48.57±0.51γaC	48.74±0.40 ^{yaC}	$48.84{\pm}0.34^{\text{yaC}}$	48.95±0.26 ^{yaC}	48.98±0.15 ^{yaC}	48.90±0.26 ^{yaC}
	24 hours	52.40±0.25 ^{βaB}	52.35±0.27 ^{δaB}	52.20±0.24 ^{δaB}	52.40±0.28 ^{δaB}	$52.67 \pm 0.26^{\delta aB}$	$52.26 \pm 0.50^{\delta aB}$
	7 days	$53.06 \pm 0.19^{\beta aA}$	53.03±0.21 ^{δaA}	$52.98 \pm 0.09^{\delta aA}$	$53.00{\pm}0.28^{\delta aA}$	$53.27 \pm 0.33^{\delta aA}$	52.92±0.26 ^{δaA}

Table 2- Mean KHN±SD for KHN for Unifil Core EM (UCE) at 6 depths of cavity after 3 post-irradiation times by 4 light-exposure methods

The same lower-case superscript Greek characters indicate no statistically significant differences between exposure methods at same post-irradiation time in the same depth of cavity; the same lower-case superscript letters indicate no statistically significant differences between depths of cavity at same post-irradiation time in the same exposure method; the same upper-case superscript letters indicate no statistically significant differences between post-irradiation times at same depth of cavity in the same exposure method (p>0.05)

KHN=Knoop hardness number

Exposure method	Post- irradiation time	Depth of cavity (mm)					
		0,5	2	4	6	8	10
Exposure time of 20 s (20 s)	0.5 hours	36.26±1.31αaC	34.18±0.97 ^{αbC}	32.99±0.64 ^{αbC}	31.69±0.64 ^{αcC}	29.10±0.49 ^{αdC}	27.31±0.39 ^{αeC}
	24 hours	41.22±0.81 ^{c/aB}	38.42±1.14 ^{αbB}	38.35±1.07 ^{αbB}	38.42±1.19 ^{αbB}	$37.42 \pm 1.33^{\alpha bB}$	36.87±1.50 ^{αbB}
	7 days	42.86±0.80 ^α	40.52±1.04 ^{αbA}	39.79±1.20αbcA	39.56±1.15αbcA	38.37±1.01αcA	38.08±1.23 ^{αcA}
Bonding + exposure time of 20 s (B+20 s)	0.5 hours	$32.23{\pm}0.97^{\beta aC}$	30.78±0.79 ^{βbC}	30.58±0.65 ^{βbC}	30.54±0.47 ^{βbC}	30.10±0.93 ^{αbC}	29.49±0.79 ^{βcC}
	24 hours	40.12±0.76αaB	37.97±1.08 ^{αbB}	37.56±0.25 ^{αbcB}	37.50±0.11 ^{αbcB}	37.39±0.21 odbcB	36.79±0.42 ^{αcB}
	7 days	42.06±0.40 ^{ccaA}	40.21±0.28 ^{αbA}	39.08±0.24 ^{αcA}	39.19±0.26αCA	38.82±0.26 ^{αcA}	$38.09 \pm 0.43^{\alpha dA}$
Exposure time of 40 s (40 s)	0.5 hours	37.19±0.24 ^{γaC}	36.50±0.37 ^{γaC}	35.28±1.15 ^{γbB}	34.14±0.61 ^{ycC}	$32.22 \pm 0.17^{\beta dB}$	30.36±0.13 ^{yeC}
	24 hours	$42.31 \pm 0.38^{\beta aB}$	40.93±0.53 ^{βbB}	40.90±0.61 ^{βbA}	$40.62 \pm 0.49^{\beta bcB}$	39.80±1.10 ^{βbcA}	39.51±0.87 ^{βcB}
	7 days	$43.07 \pm 0.15^{\beta aA}$	42.12±0.45 ^{βbA}	41.87±0.45 ^{βbA}	41.97±0.47 ^{βbA}	40.76±0.61 ^{βcA}	40.77±0.69 ^{βcA}
Exposure time of 120 s (120 s)	0.5 hours	37.35±0.32 ^{γaC}	$37.60 {\pm} 0.35^{\delta aC}$	$37.46{\pm}0.25^{\delta aC}$	$37.59 {\pm} 0.18^{\delta aB}$	37.36±0.21 ^{yaC}	$37.32 {\pm} 0.25^{\delta aB}$
	24 hours	$42.58 \pm 0.30^{\beta aB}$	42.67±0.62 ^{yaAB}	42.60±0.46 ^{yaAB}	42.59±0.46 ^{yaA}	42.62±0.55 ^{yaAB}	42.7±0.44 ^{yaA}
	7 days	$42.78 \pm 0.40^{\beta aA}$	42.98±0.77 ^{yaA}	42.97±0.86 ^{yaA}	42.81±0.45 ^{yaA}	43.00±0.85 ^{yaA}	43.06±0.60 ^{yaA}

For DCP, with the B+20 s method, no significant differences in the KHN values were observed at 0.5 hours post-irradiation at all depths of cavity, except 0.5 mm (p>0.05). However, at 24 hours and 7 days post-irradiation, the B+20 s method resulted in significantly higher KHN values than the 20 s method at all depths of cavity, except 0.5 mm (p<0.05). On the other hand, for UCE, at 0.5 hours post-irradiation, the B+20 s method showed significantly lower KHN values than the 20 s method up to the depth of 6.0 mm; however, the KHN values were significantly higher at the depths of 8.0 mm and 10.0 mm (p<0.05). At 24 hours and 7 days post-irradiation, no significant differences in KHN values were observed between the B+20 s and 20 s method at all depths of cavity (p>0.05).

For all light-exposure methods, except the $120\,\mathrm{s}$ method, the UCE exhibited significantly higher KHN values at all depths of cavity, except at $4.0\,\mathrm{mm}$ in the $40\,\mathrm{s}$ method, at 7 days post-irradiation than at $0.5\,\mathrm{hours}$ and $24\,\mathrm{hours}$ post-irradiation (p<0.05). It was found that the KHN values of DCP were significantly higher than those of UCE at all depths of cavity and post-irradiation times, regardless of the exposure method (p<0.05).

DISCUSSION

The results of this study indicate that the KHNs

of dual-cured core build-up resin composites depend on the light-exposure method, including irradiation duration, use of bonding agent, depth of cavity, post-irradiation time, and material brand. Therefore, the research hypotheses formulated for this study must be rejected.

KHN has been shown to be a good indicator of the degree of conversion/polymerization based on its good correlation with infrared spectroscopy^{9,10,31}. However, the prediction of an absolute value of degree of conversion by means of an absolute hardness value is not achievable, since other factors such as type and size of filler, filler load, monomer composition, quantity of initiators, and the ratio of chemical- and light-cured components strongly influence the final quantity of reacted monomers^{2,5,14}. Microhardness data from the same resin cement only should be compared according to the depth of the root canal or time elapsed since luting³. KHNs could be used to reflect the degree of conversion at different depths of a resin composite³². Therefore, KHNs were measured in the present study to reflect monomer conversion at different depths of cavity and post-irradiation times in the dual-cured core build-up resin composites.

The KHNs of both core build-up resin composites irradiated for 20 or 40 seconds were the highest at the depth of 0.5 mm, and gradually decreased with increasing depth. This phenomenon could

be simply attributed to the direction of photoinitiation. Light irradiation was focused on the top surface of the cavity. Therefore, polymerization of the resin composites, by means of photoactivated free radicals, may occur immediately at the shallow depths of cavity. This finding, that is, KHN of light- and dual-cured resin composites are affected by depth of cavity, has been reported previously^{4,6-8}. The chemical-curing mechanism of dual-cured resin composites is usually based on a redox reaction of benzoyl peroxide with aromatic tertiary amines, which generates free radicals that break the aliphatic carbon double bonds to initiate the polymerization process. It is supposed that immediate photo-activation after light irradiation, despite causing a rapid increase in the viscosity of the polymer matrix, does not hinder migration of the activated free radicals responsible for further chemically induced polymerization. Although the photo-activated free radicals at shallow depths of cavity could induce chain propagation of the resin polymer in the downward direction, the exact polymerization mechanism of dual-cured core build-up resin composite at greater depths of cavity remains unknown. It is difficult to distinguish clearly between the depths of cavity at which polymerization of the resin composite occurs through photo-initiation and those at which polymerization occurs by means of chemical initiation alone.

Evaluation of the duration of photo-activation revealed that a longer exposure time of 40 seconds resulted in significantly higher KHN values than did a shorter exposure time of 20 seconds at all depths of cavity and post-irradiation times for both resin composites; however, this trend was not observed in UCE at the depth of 0.5 mm. Thus, our study showed that prolonged irradiation durations resulted in increased hardness values. Light-curing units with lower power density needed longer light-exposure times to produce a similar microhardness value of resin composite as that of light sources with high power density¹¹. Therefore, with longer light-exposure durations, which result in higher energy densities at a given irradiance, more photo-sensitizer molecules are activated, which in turn increase the free radical concentration and consequently the conversion of double bonds. However, at the depth of 0.5 mm, no significant differences were observed in the KHN values between the light-irradiation durations of 40 seconds and 120 seconds (40 seconds each on the top and either side of the cavity) for both resin composites. This finding might be attributed to the fact that the polymer network developed during light exposure for 40 seconds does not allow any additional mobility of the polymer chains in order to increase monomer conversion, indicating that the light-irradiation duration of 40 seconds alone is sufficient.

Dentin hardness ranges from KHN values of 50 to 70, depending on the distance from the amelodentinal junction²¹. The mean microhardness (KHN) value of 52.9 for DCP with 120 s method at the depth of 10.0 mm at 7 days post-irradiation therefore predicts similar mechanical properties to that of dentin. The equal degree of polymerization within the core material may support a uniform distribution of stress along the tooth-material interface under load. Longer light-exposure durations result in superior microhardness, but, at the same time, contribute to an increase of shrinkage and contraction stress of the dualcured core build-up resin composite³⁸. In cases of significant coronal destruction, it is necessary to replace the lost tooth structure with a core buildup material to attain full-coverage restoration. The cast post and core, prefabricated post and core materials, and coronal-radicular build-ups are the available options for this purpose. The fracture resistance and survival probability of post and core restorations depend on several factors such as the post material, luting agent, amount and condition of residual tooth structure, core material, preparation of the tooth for restorative procedures, and, finally, the fixed restoration^{16,22,25,34}. Therefore, when using dual-cured core build-up resin composites, it is preferable to prepare the composites by using the effective exposure method.

Before core build-up resin composites are used, a bonding agent must be applied on the cavity wall. The effectiveness of the bonding agent on the KHN as an indirect method of monomer conversion differed between the 2 resin composites. For DCP, at 24 hours and 7 days post-irradiation, application of the bonding agent resulted in significantly higher KHNs than the 20 s method at all depths of cavity; however, this was not observed with UCE. The initiator and/or accelerator present in the bonding agent would promote monomer conversion of the dual-cured core build-up resin composites. On the other hand, the acidic monomer in the bonding agent inhibits the amine co-initiator in the dualcured materials¹⁶, which in turn adversely affects polymerization of dual-cured core build-up resin composites. To prevent this inhibitory effect, aromatic sulfinic acid sodium salts are sometimes added to the bonding agent^{15,37}. The differences in compositions between the 2 core build-up resin composites might be responsible for the difference in their KHN behaviors.

In the present study, both resin composites at all depths of cavity had a post-curing effect 7 days after irradiation, showing statistically higher KHN values than those at 0.5 hours post-irradiation for all light-exposure methods. These results are

consistent with those of a previous study³, but not with those of another study⁴¹, which did not find changes in microhardness values 24 hours after irradiation. However, the polymerization reaction of the dual-cured materials might be specific¹⁰, and these previous studies used resin cements, whereas dual-cured core build-up resin composites were used in the present study. In fact, in the former study, neither did the authors evaluate dual-cured core build-up resin composites nor did they evaluate hardness behavior of dual-cured core build-up resin composite in simulated depths of cavity.

Dual-cured resin composites have been introduced for use as both luting and core buildup materials. Superior physical properties are important for a successful restoration. In this study, it was found that the KHNs of DCP were superior to those of UCE, regardless of exposure methods. Various factors can influence the microhardness of a resin composite, such as filler load, type, or size, or resin matrix type^{17,18}. Increasing the monomer viscosity decreased the hardness²⁷. In this study, the association between KHN values of the resin composite and composition of filler and matrix resin was not evaluated. On the other hand, the differences between the KHNs at the depths of 0.5 mm and 10.0 mm in UCE were lower than those in DCP, regardless of the light-exposure method. Dual-cured materials differ markedly in terms of the relative content of light- and selfactivated catalysts¹³. Differences in the degree of conversion among materials when subjected to various curing protocols may consequently be attributed to variations in catalyst systems. In the present study, since similar KHNs were observed irrespective of the light-exposure method at greater depths of cavity in UCE rather than DCP, it might be inferred that the former exhibits high levels of chemically curing activators compared with the latter, compensating for attenuation of light energy at greater depths of cavity. Indeed, the polymerization behavior of dual-cured resin composites is strongly related to the material and can vary as a function of composition³⁵. The speed of the polymerization reaction is strongly influenced by inhibitor concentration in the unfilled light-cured methacrylate-based systems³⁰. Therefore, the curing mechanism of a specific composite material may not be applicable to other materials.

CONCLUSIONS

Within the limitations of the present study, the following conclusions can be drawn:

1. The microhardness of the dual-cured core build-up resin composites vary depending on the light-exposure method, including irradiation duration, use of bonding agent, depth of cavity, post-irradiation time, and material brand.

- 2. For both resin composites, irradiation for 120 seconds does not result in significant differences in KHNs among all 6 depths of cavity at all post-irradiation times.
- 3. Based on the physical property of microhardness behavior, dual-cured core build-up resin composites are preferably prepared by the effective exposure method.

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