# Effect of Energy Density on the Physical Properties of Resin-Based Restorative Materials when Polymerized with Quartz-Tungsten Halogen or LED-Light

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# **ABSTRACT**

Objectives: The purpose of this study was to evaluate flexural strengths, moduli, and maximum deflection of Clearfil AP-X (APX) and Ceram-X Mono (CXM) when cured with a quartz-tungsten halogen (QTH) or an LED-light (LED).

Methods: Specimens were made according to ISO 4049 and cured with QTH or LED for 10, 20 or 60 s. Flexural strength, modulus, and deflection were determined after 24 h water storage at 37°C and after thermocycling. Statistical significance was P<.05.

Results: Flexural strength did not depend on energy density or curing light and was significantly higher for APX than for CXM but decreased after thermocycling for both materials. Modulus and deflection depended on energy density. Modulus was significantly higher for APX than for CXM and increased for APX but decreased for CXM after thermocycling. Deflection decreased with increasing energy density and decreased after thermocycling. Though energy density did not influence flexural strength, it positively correlated with flexural modulus and negatively with maximum deflection.

Conclusions: Energy density did not influence flexural strength but modulus and deflection. Thermocycling affected all material properties. The LED was as effective as the QTH. (Eur J Dent 2010;4:183-191)

Key words: Resin-based filling material; Mechanical properties; LED; Curing device; Curing time.

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## INTRODUCTION

LED-curing lights are increasingly used to polymerize resin-based filling materials. These very modern curing devices offer several advantages, such as high power output and very low weight. Although the first-generation devices did not perform well,<sup>1,2</sup> the latest generation is reported to work optimally.<sup>3-5</sup> The lifetime of LEDs reaches 10,000 hours compared to approximately 50 hours for a quartz-tungsten halogen bulb. They also

cause less temperature increase during the polymerization of resin-based filling materials.<sup>6-8</sup>

There has been much research into the influence of LED-curing lights on the hardness, 3,9,10 shrinkage, 11 temperature rise, 3,6,8 cross-link density,4 and degree of conversion12-14 of resin-based filling materials. But only few studies were found considering flexural strength and flexural modulus. 9,15-17 These studies showed that the second generation of LED-curing lights performed similarly to the quartz-tungsten halogen devices (QTH). Although the energy densities of the LEDcuring lights were higher than those of the QTH, significant temperature increase was not measured in the pulp chamber and increased cell damage was not observed.<sup>3,6,8</sup> Only one of these studies investigated flexural strength according to EN ISO 4049, 18 and none measured flexural modulus. Although some publications compared the influence of energy densities of QTH and LED on hardness<sup>3,19-21</sup> and compressive strength,<sup>22</sup> none was found that compared the influence on flexural strength, flexural modulus, and deflection. The literature has described the effect of thermocycling on the physical properties of resin-based restorative materials when cured with QTH<sup>23,24</sup> but has not considered LED-lights and different energy densities. Therefore, there are only few possibilities for an accurate comparison of the results. Furthermore, no literature was found about the influence of LED-curing lights on flexural properties of ormocers.

An important feature of the resin matrix is that it should absorb energy and reduce stress concentrations by providing fracture toughness or ductility to maximize damage tolerance.<sup>25</sup> The matrices of resin-based restorative materials were also shown to be one essential reason for brittle fracture wear.<sup>26-28</sup> Maximum deflection measured during a three-point-bending test was used to obtain knowledge about the elasticity or toughness, respectively, of resin materials.<sup>29,30</sup>

Therefore, the goal of the present investigation was to determine flexural strength (according to EN ISO 4049), flexural modulus, and deflection of an ormocer (microhybrid composite with partial silicium-organically modified resin matrix) in comparison with a microhybrid resin-based filling material when polymerized with QTH or LED-light. The null hypothesis was that there is no difference

(a) between the investigated properties when irradiated with QTH or LED-light and (b) in the investigated properties between the ormocer and the microhybrid.

## **MATERIALS AND METHODS**

The ormocer Ceram-X Mono, shade M5 (Dentsply DeTrey GmbH, Constance, Germany), and the microhybrid Clearfil AP-X, shade A3 (Kuraray Europe GmbH, Frankfurt, Germany), were used as test materials (Table 1). Ceram-X Mono shade M5 is equivalent to Clearfil AP-X shade A3. The quartztungsten halogen light Hilux Ultra Plus (Benlioglu Dental Inc., Ankara, Turkey) with a 10 mm light guide and the LED-light curing device SmartLite PS with an 8 mm light tip (Dentsply DeTrey GmbH, Constance, Germany) were used to polymerize the materials in the constant polymerization mode. Each time after a series of ten specimens was cured, the output of each of the curing devices was controlled with a photometer (Curing Light Meter, Benlioglu Dental Inc.). Irradiances between 750 and 850 mW/cm<sup>2</sup> (mean 800  $\pm$  67 mW/cm<sup>2</sup>) were measured for the Hilux Ultra Plus and between 1100 and 1300 mW/cm2 (mean  $1200 \pm 98 \text{ mW/cm}^2$ ) for the SmartLite PS. No significant decrease of the output of either device was observed. The energy density of each curing device was calculated for the different exposure times (Table 2).

The preparation of the specimens was done according to EN ISO 4049.18 From each material, 120 specimens with a size of  $(25\pm2)$  mm x  $(2\pm0.1)$  mm x (2±0.1) mm were manufactured at 22.0 - 23.0°C (room temperature) and a relative humidity of 50%. Prior to polymerization, both sides of the specimens were covered with a 0.05 mm transparent polyester film. The initial curing location was in the center of the specimen. Two additional curing increments were used on either side of the initial curing location from the center of each specimen toward its end. The specimens were turned over, and the curing sequence was repeated on the bottom. The curing sequence resulted in a total of five curing increments on each side of each specimen (ten in total). The 120 specimens of each material were subdivided into three groups, each of 40 specimens. One half of the specimens of group 1 was cured with Hilux Ultra Plus, the other half with SmartLite PS for 10 s; the specimens of group 2 were cured for 20 s, group 3 for 60 s.

Flexural strength  $\sigma$  was calculated in MPa by:

 $\sigma = -$ 

Flexural modulus E was calculated by:

$$E = \frac{L^3}{4b \times h^3} \times \frac{F}{Y}$$

F = maximum strength in N

L = distance between the rests

b = width of the specimen

h = height of the specimen

F / Y = slope of linear part of the stress-strain curve

Maximum deflection was taken directly from the stress-strain-curve.

# Statistical analysis

Statistical analysis was conducted with SPSS software 12.0 (SPSS Software, Munich, Germany). Means and standard deviations were calculated. Normal distribution was proven by the Kolmogoroff-Smirnoff Test. Multiple comparisons were made for each of the tested properties with the univariate Anova followed by a Scheffe post hoc test and t-tests for unpaired samples. Correlations were calculated according to Pearson. Statistical significance for all tests was considered as P<.05.

# **RESULTS**

Means and standard deviations of flexural strength, flexural modulus, and maximum deflection are shown in Table 3. Significant differences of flexural strength, flexural modulus, and deflection were calculated between the microhybrid Clearfil AP-X and the ormocer Ceram-X Mono prior to and after thermocycling for all curing times and both of the curing devices (Tables 5, 6 and 7).

Clearfil AP-X showed significantly higher flexural strength than Ceram X Mono for all energy densities, curing devices, and aging conditions (Tables 3, 4 and 5). Except for Clearfil AP-X 20 s QTH-cured, neither prior to nor after thermocycling was a significant influence of energy density or curing device on flexural strength observed for the test materials (Tables 3, 4 and 5). Flexural strengths of all LED-light polymerized samples significantly decreased after thermocycling, which was not the case for all of the QTH-cured specimens (Tables 3 and 7). No correlation was found between energy density and flexural strength for any of the test materials (Tables 3 and 8).

Clearfil AP-X showed significantly higher flexural modulus than Ceram X Mono for all energy densities, curing devices, and aging conditions (Tables 3, 4 and 5). Flexural modulus increased for Clearfil AP-X with increasing curing time or energy density, respectively, and after thermocycling for both of the curing devices. The SmartLite-cured Clearfil AP-X specimens had higher modulus values than the Hilux Ultra Plus-cured specimens. In contrast to Clearfil AP-X, the flexural modulus of Ceram-X Mono remained constant or decreased after thermocycling (Tables 3, 4 and 5). Clearfil AP-X and Ceram-X Mono showed a significantly positive correlation between energy density and flexural modulus prior to as well as following thermocycling (Table 8).

Maximum deflection was significantly lower for Clearfil AP-X than for Ceram-X Mono when QTH-cured, but no differences were found for the LED-light-cured samples. No influence of curing time or curing device was detected either for non-thermocycled or for thermocyled Clearfil AP-X. Deflection decreased for Ceram-X Mono with increasing curing time and both curing devices (Tables 3 and 6). After thermocycling, the values of both test materials decreased (Tables 3 and 7). Ceram-X Mono showed a significant strong negative correlation between energy density and maximum deflection prior to and after thermocycling. but Clearfil AP-X did not. Further correlations were detected for both test materials for flexural strength and flexural modulus with maximum deflection (Table 8).

# **DISCUSSION**

This study investigated the influence of QTH or LED-light on flexural strength, flexural modulus, and deflection of two different types of resinbased filling materials according to EN ISO 4049.<sup>18</sup>

Flexural strength and flexural modulus are appropriate for evaluating the quality of the light-curing process<sup>9,15,24,31</sup> and maximum deflection furnished some knowledge about the materials' elasticity or toughness, respectively.<sup>29,30</sup> Literature reported that thermocycling also had an impact on the flexural properties.<sup>24</sup> Since the degree of conversion not only depends on the curing conditions but also on the chemical character of the resin matrix,<sup>32,33</sup> a microhybrid composite (Clearfil AP-X) and a microhybrid composite with partial silicium-organically modified resin matrix, so-called ormocer, (Ceram-X Mono) were chosen for this investigation. The spectral ranges of QTH and sev-

eral contemporary LED-lights (also SmartLite PS) were reported by the literature and documented in Table  $2.^{34,35}$ 

Several publications have shown that the combination of energy density and exposure time has significant influence on the degree of cure, flexural strength, and flexural modulus. 31,36,37 Peutzfeldt et al 19 found higher levels of degree of cure, flexural strength, and flexural modulus for TetricCeram with increasing energy densities. They concluded that the higher the energy density, the higher the degree of cure and mechanical properties. The present study could not confirm these findings for flexural strength but could for flexural modulus,

Table 1. Test materials.

Material	Formulation	Manufacturer
	Resin matrix: methacrylate modified	
Ceram-X-Mono <sup>1</sup>	polysiloxane, dimethacrylate resin	
#05110000198	Inorganic filler: Ba-Al-borosilicate glass,	
Shade: M5 = A3,	pyrogenic SiO <sub>2</sub>	
microhybride composite with	Filler load: 76 mass-%, 57 vol%	DeTrey Dentsply GmbH,
partial silicium-organically	Photoinitiator: camphorquinone	Constance, Germany
modified resin matrix	Synergist: ethyl-4-diemthylamino benzo-	
(Ormocer)	ate, UV stabilizer	
	Stabilizer: butylated hydroxy toluene	
	Resin matrix: Bis-GMA, Tegdma	
	Inorganic filler: Ba-glass, silica, pyro-	
Clearfil AP-X <sup>2</sup> #01122B,	genic SiO <sub>2</sub>	Kuraray Co. Inc.,
Shade: A3, microhybride	Filler load: 85.5 mass-%, 70 vol%	Kurashiki, Japan
	Photoinitiator: camphorquinone	
	Synergist: NI	

Bis-GMA = Bisphenol-A-dimethacrylate, Tegdma = Triethylenglycol dimetacrylate, HPMA = 3-Hydroxpropyl methacrylate, NI = No information

Table 2. Irradiances and energy densities of Hilux Ultra Plus and SmartLite PS.

	Curing time [s]	Irradiance [mW/cm²]	Energy density [mWs/cm²]	Spectral range [nm] *)	
	10	800	8000	400-520, broad,	
Hilux Ultra Plus QTH	20	800	16000	Flat distribution,	
	60	800	48000	maximum: 520	
	10	1200	12000		
SmartLite PS LED	20	1200	24000	450-470	
	60	1200	72000	peak: 460	

Camphorquinone spectral range: 350 - 550 nm, peak: 468 nm \*)

<sup>&</sup>lt;sup>1</sup> Formulation according to the literature<sup>3,4,39</sup>

<sup>&</sup>lt;sup>2</sup> Formulation according to the literature<sup>1,44</sup>

<sup>\*)</sup> according to literature<sup>22,43</sup>

**Table 3.** Flexural strength, flexural modulus, maximum deflection and (standard deviation) of Clearfil AP-X and Ceram-X Mono prior to (24 h) and after thermocycling (TC).

	Ceram-X Mono								
	Hilo				Lite PS ED)		Iltra Plus IAL)	SmartLite PS (LED)	
		24 h	5000 TC	24 h	5000 TC	24 h	5000 TC	24 h	5000 TC
	10s	114 (14)	103 (18)	124 (25)	109 (11)	86 (10)	57 (11)	87 (10)	57 (12)
Flexural strength [MPa]	20s	148 (33)	107 (17)	122 (22)	105 (18)	69 (15)	59 (9)	79 (9)	55 (8)
	60s	125 (22)	100 (18)	131 (19)	116 (21)	74 (21)	63 (11)	86 (13)	58 (15)
	10s	9550 (330)	12400 (800)	12430 (790)	12660 (330)	6110 (350)	5730 (410)	7190 (560)	7240 (860)
Flexural modulus [MPa]	20s	11180 (640)	12780 (320)	12100 (290)	15200 (990)	6650 (850)	5560 (360)	6920 (860)	7300 (240)
	60s	12140 (490)	12560 (340)	13000 (640)	15700 (850)	6810 (640)	6750 (290)	8070 (720)	7010 (450)
	10s	0.41 (0.07)	0.27 (0.03)	0.36 (0.03)	0.26 (0.03)	0.51 (0.05)	0.40 (0.03)	0.48 (0.04)	0.31 (0.03)
Maximum deflection [mm]	20s	0.39 (0.08)	0.27 (0.04)	0.42 (0.05)	0.29 (0.03)	0.44 (0.07)	0.40 (0.03)	0.42 (0.04)	0.30 (0.05)
	60s	0.37 (0.06)	0.32 (0.03)	0.36 (0.03)	0.36 (0.05)	0.41 (0.06)	0.30 (0.04)	0.38 (0.04)	0.28 (0.06)

**Table 4.** Significances (bold and italic) of flexural strength and flexural modulus between the materials, curing lights and curing times after 24 hours storage in water at 37°C (P<.05).

	Flexural strength after 24 hours in water at 37°C													
	Hilux Ultra Plus									SmartLite PS				
	Clearfil AP-X Ceram-X Mono								Cle	earfil AF	P-X	Cer	am-X M	lono
			10s	20s	60s	10s	20s	60s	10s	20s	60s	10s	20s	60s
		10s		0.253	0.999	0.402	0.005	0.028	1.000	1.000	0.960	0.499	0.001	0.405
	Clearfil AP-X	20s	0.002		0.841	0.000	0.000	0.000	0.794	0.681	0.980	0.000	0.000	0.000
Hilux Ultra		60s	0.000	0.497		0.031	0.000	0.000	1.000	1.000	1.000	0.049	0.003	0.031
Plus		10s	0.000	0.000	0.000		0.965	0.999	0.043	0.079	0.004	1.000	1.000	1.000
	Ceram-X	20s	0.000	0.000	0.000	0.975		1.000	0.000	0.000	0.000	0.934	1.000	0.964
	Mono	60s	0.000	0.000	0.000	0.875	1.000		0.001	0.002	0.000	0.996	1.000	0.999
		10s	0.000	0.081	1.000	0.000	0.000	0.000		1.000	1.000	0.066	0.004	0.044
	Clearfil AP-X	20s	0.000	0.600	1.000	0.000	0.000	0.000	0.999		1.000	0.117	0.010	0.080
		60s	0.000	0.000	0.667	0.000	0.000	0.000	0.971	0.481		0.007	0.000	0.004
Smart Lite PS		10s	0.000	0.000	0.000	0.203	0.974	0.999	0.000	0.000	0.000		1.000	1.000
F3	Ceram-X	20s	0.000	0.000	0.000	0.671	1.000	1.000	0.000	0.000	0.000	1.000		1.000
	Mono	60s	0.000	0.000	0.000	0.000	0.013	0.071	0.000	0.000	0.000	0.546	0.131	
					F	lexural	modulu	s after 2	24 hours	in wate	er at 37°	C		

which was detected to correlate strongly positively with energy density for both test materials prior to and after thermocycling (Table 8). However, due to the restricted number of experimental groups, the present study might have failed to reveal a signifi-

cant influence of the combination of energy density and curing time on flexural strength.

The correlation of energy density and flexural modulus found in the present investigation was positive and very strong prior to and after thermo-

**Table 5.** Significances (bold and italic) of flexural strength and flexural modulus between the materials, curing lights and curing times after 30 days storage in water at  $37^{\circ}$ C followed by 5000 thermocycles between + 5 and + 55°C (P<.05).

	Flexural strength after 24 hours in water at 37°C followed by 5000 thermocycles													
	Hilux Ultra Plus								SmartLite PS					
			Cle	earfil AF	P-X	Cer	am-X M	ono	Cl	earfil AF	P-X	Cer	am-X M	lono
			10s	20s	60s	10s	20s	60s	10s	20s	60s	10s	20s	60s
		10s		1.000	1.000	0.000	0.000	0.000	1.000	1.000	0.967	0.000	0.000	0.000
	Clearfil AP-X	20s	0.998		1.000	0.000	0.000	0.000	1.000	1.000	0.999	0.000	0.000	0.000
Hilux Ultra		60s	1.000	1.000		0.000	0.000	0.002	0.999	1.000	0.869	0.000	0.000	0.000
Plus	0 V	10s	0.000	0.000	0.000		1.000	1.000	0.000	0.000	0.000	1.000	1.000	1.000
	Ceram-X Mono	20s	0.000	0.000	0.000	1.000		1.000	0.000	0.000	0.000	1.000	1.000	1.000
	IVIOIIO	60s	0.000	0.000	0.000	0.156	0.037		0.000	0.000	0.000	1.000	1.000	1.000
		10s	1.000	1.000	1.000	0.000	0.000	0.000		1.000	1.000	0.000	0.000	0.000
	Clearfil AP-X	20s	0.000	0.000	0.000	0.000	0.000	0.000	0.000		0.993	0.000	0.000	0.000
Smart Lite		60s	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.983		0.000	0.000	0.000
PS		10s	0.000	0.000	0.000	0.001	0.000	0.979	0.000	0.000	0.000		1.000	1.000
	Ceram-X Mono	20s	0.000	0.000	0.000	0.000	0.000	0.941	0.000	0.000	0.000	1.000		1.000
	MOHO	60s	0.000	0.000	0.000	0.014	0.002	1.000	0.000	0.000	0.000	1.000	1.000	
			FI	exural	modulu	s after 2	4 hours	in wate	r at 37°	C follow	ed by 50	000 ther	mocycle	es

**Table 6.** Significances (bold and italic) of maximum deflection prior to and after thermocycling between the materials, curing lights and curing times (P<.05).

	Maximum deflection after 24 hours in water at 37°C															
	Hilux Ultra Plus										SmartLite PS					
	Clearfil AP-X Ceram-X Mono							Cle	earfil AP	P-X	Cer	am-X M	ono			
			10s	20s	60s	10s	20s	60s	10s	20s	60s	10s	20s	60s		
		10s		1.000	0.998	0.145	1.000	1.000	0.988	1.000	0.937	0.691	1.000	1.000		
	Clearfil AP-X	20s	0.889		1.000	0.039	0.983	1.000	1.000	1.000	1.000	0.338	0.999	1.000		
Hilux Ultra		60s	0.883	1.000		0.006	0.879	1.000	1.000	0.992	1.000	0.114	0.997	1.000		
Plus	0 V	10s	0.877	0.041	0.039		0.694	0.098	0.003	0.275	0.001	1.000	0.383	0.020		
	Ceram-X Mono	20s	0.044	0.000	0.000	0.943		0.999	0.762	1.000	0.600	0.987	1.000	0.965		
		60s	1.000	0.993	0.993	0.611	0.011		0.998	1.000	0.992	0.569	1.000	1.000		
		10s	1.000	0.869	0.863	0.930	0.080	1.000		0.970	1.000	0.060	0.931	1.000		
	Clearfil AP-X	20s	0.736	1.000	1.000	0.016	0.000	0.964	0.710		0.916	0.841	1.000	0.999		
Smart Lite		60s	1.000	0.999	0.999	0.469	0.005	1.000	1.000	0.988		0.024	0.842	1.000		
PS	0 4	10s	1.000	0.965	0.963	0.792	0.029	1.000	1.000	0.884	1.000		0.914	0.997		
	Ceram-X Mono	20s	1.000	0.998	0.938	0.497	0.006	1.000	1.000	0.985	1.000	0.939		1.000		
	IMOHO	60s	0.994	1.000	1.000	0.185	0.001	1.000	0.990	1.000	1.000	0.999	1.000			
					M	aximum	deflect	ion afte	r 24 hou	rs in wa	ter at 3'	7°C				
							followe	d by 500	00 therm	nocycles	5					

cycling for both of the test materials. The lower correlation value for Ceram-X Mono after TC (0.35, P<.0000) might indicate that the flexural modulus did not increase linearly with energy density, as was determined in the study by Peutzfeldt et al.31 Another explanation might be that the modulus decreased significantly after thermocycling. A certain explanation was not possible because of the limitations of this study. The results also show (Tables 2 and 3) that the highest energy density (SmartLite PS, 60 s curing time) resulted in the highest flexural modulus for both of the test materials. These results strongly supported the existing literature.

Energy density correlated strongly negatively with maximum deflection for Ceram-X Mono, indicating a higher degree of cure (Table 8), and was thus also in accordance with the results of Peutzfeldt et al.31 The fact that no correlation between energy density and deflection was found for Clearfil AP-X (Table 8) might be explained by the higher filler / matrix ratio that over-compensated the influence of energy density on deflection but not on flexural modulus, since this effect was much stronger.

The findings of the present study also showed that the formulation of the material itself influenced flexural strength, flexural modulus, and deflection (Tables 3 to 6). The difference in flexural strength was not only caused by the higher filler content<sup>38</sup> of Clearfil AP-X but also by the filler type (agglomerated pyrogenic SiO<sub>2</sub>) and the high con-

Table 7. Significances (bold and italic) between the 24 h storage in water at 37°C and the 30 days storage in water at 37°C followed by 5000 thermocycles between +5 and +55°C (P<.05).

			Flexural strength	Flexural modulus	Max. deflection
		10s	0.124	0.000	0.000
	Clearfil AP-X	20s	0.000	0.000	0.000
11.7 1111 111		60s	0.001	0.057	0.015
Hilux Ultra Plus		10s	0.000	0.150	0.000
	Ceram-X Mono	20s	0.181	0.000	0.099
		60s	0.125	0.830	0.000
		10s	0.039	0.396	0.000
	Clearfil AP-X	20s	0.024	0.000	0.000
C III DC		60s	0.038	0.000	0.980
Smart Lite PS		10s	0.000	0.855	0.000
	Ceram-X Mono	20s	0.002	0.156	0.000
		60s	0.000	0.000	0.000

Table 8. Correlations of light dose with flexural modulus and maximum deflection as wall as of maximum deflection with flexural strength and modulus after 24 h storage (24 h) and after thermocycling (TC) (P<.05).

	Correlation of energy density with										
Flexural modulus Maximum deflection											
	24 h	24 h	TC								
Clearfil AP-X	0.593 (P<.000)	0.579 (P<.000)	none	none							
Ceram-X Mono	0.528 (P<.000)	0.349 (P=0.007)	-0.591 (P<.000)	-0.435 (P<.000)							
		Correlation of maxi	num deflection with								
	Flexural	modulus	Flexural	strength							
	24 h	TC	24 h	тс							
Clearfil AP-X	-0.411 (P=0.005)	none	none	0.713 (P<.000)							
Ceram-X Mono	-0.397 (P=0.006)	-0.607 (P<.000)	0.447 (P=0.001)	0.556 (P<.000)							

tent of the more rigid Bis-GMA-containing organic matrix. Ceram-X Mono's organic matrix, containing methacrylate modified polysiloxanes and finely dispersed SiO<sub>2</sub> particles, was more elastic.<sup>4,39-41</sup> Clearfil AP-X was also found to have a significantly higher flexural modulus than Ceram-X Mono mainly due to its higher filler content (Table 1). The positive correlation between filler content, flexural strength, or flexural modulus was previously reported by Rodrigues Junior et al.<sup>38</sup> The strong negative correlation of flexural modulus with maximum deflection showed the loss of elasticity with increasing flexural modulus or increasing filler content, respectively.

EN ISO 4049<sup>18</sup> requires flexural strength ≥ 80 MPa, and the literature recommends flexural modulus ≥ 10000 MPa for resin-based filling materials used in occlusal areas.<sup>24</sup> Only Clearfil AP-X fulfilled these requirements prior to and after thermocycling independent of the light-curing device. The results showed that there were no significant differences between the flexural strength values of Ceram-X Mono and only for the 20 s irradiated samples of Clearfil AP-X when cured with QTH or LED-light. Both of the materials behave rather similarly after thermocycling independent of the curing light - sometimes flexural strength decreased and sometimes it did not. No correlation with the curing device was found. These findings supported the literature, 9,14,16,42 which concluded that LED-lights were as effective as QTH for polymerization of the materials used.

Furthermore, it was found that the flexural modulus of Clearfil AP-X remained constant or even increased after thermocycling, whereas the modulus of Ceram-X Mono remained constant or even decreased. Such behaviour of microhybrids and ormocers was also reported in the literature, and it was concluded that the significantly lower filler content of the ormocer could be one possible cause.<sup>24</sup> The test materials of this study also differed significantly in filler content, so that the same conclusion might be drawn. Finally, as already discussed in a preceding paragraph, LED-lights providing high energy densities resulted in significantly higher flexural moduli.

### **CONCLUSIONS**

Energy density did not influence flexural strength but did influence modulus and deflection.

The thermocycling process affected all tested properties of the materials. The LED was as effective as the QTH for polymerization of the materials used. Therefore, part (a) of the null hypothesis is accepted for flexural strength, rejected for flexural modulus, and partially rejected for maximum deflection, and part (b) was rejected.

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