Water sorption of heat-cured acrylic resin

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ABSTRACT

Aims: To evaluate the effect of the different curing methods conventional water bath heat—curing and microwave energy curing method and different immersion periods on the water sorption and solubility of the different types of heat—cured acrylic resins. Materials and methods: Water sorption and solubility were measured by means of mass change in the materials after water saturation and dehydration. Two different commercial heat—cured acrylic resins Major Base 2, and Quayle Dental resins were used in this study. Specimens were divided into two groups depending on the curing method. Thirty two specimens were prepared, eight specimens for each material in each group. Specimens were immersed in distilled water, and then subsequently removed from their containers at 1 week and 1 month of immersion for evaluation. Mean values wear compared statistically with one way analysis of variance followed by Duncan's multiple range test to determined the significant different among the groups at (p< 0.05) level of significance. Results: showed that the curing method and immersion period have a significant effect on the water sorption and solubility ratios of the resins. Curing by microwave energy method and increasing immersion period caused increasing in the ratios. The type of heat—cured acrylic resin has an effect but the difference was not significant. Conclusion: the curing method, immersion period and types of the acrylic resin have an effect on the absorption and solubility ratios.

Key words: Water sorption, acrylic resin, conventional heat-cured, microwave .

Al- Nori AKh, Ali AA, Rejab LT. Water sorption of heat–cured acrylic resin. *Al–Rafidain Dent J.* 2007; 7(2): 186–194.

Received: 14/6/2006 Sent to Referees: 2/7/2006 Accepted for Publication: 19/8/2006

INTRODUCTION

The most popular denture base material is heat-cured acrylic resin poly methyl methacrylate (PMMA). Acrylic polymers have been used as facings for fixed bridges to improve the esthetic value of the restoration. 11-41 Poly methyl methacrylate possesses dimensional change during processing and in service. (5) Poly methyl methacrylate absorb water slowly over period of time. This imbibition is due primarily to the polar properties of the resin molecules. (6,7) Sorption of material represents the amount of water adsorbed on the surface and absorbed into the body of the material during fabriccation or while the restoration is in service. Since both adsorption and absorption are involved the term sorption is usually used to include the total phenomena. Usually a serious warpage and dimensional change in the material are associated with a high percentage of water sorption. (8) Water sorption has detrimental effect on the colour stability and the wear resistance. (9) Solubility represents the mass of the soluble materials from polymers. The only soluble materials present in denture base resins are initiators, plasticizers, and free monomer. Any observed loss of weight of the resin is the measure of the specimen's solubility. (6) Water sorption and solubility were measured by means of mass change in the materials after water saturation and dehydration. (10-12) Curing processes have been modified in order to improve the physical and mechanical properties of those materials, and also to aff-

186

ord the technical work of the professionals. Different polymerization methods have been used: heat, light and microwave energy. (13,14) This last method has the advantage of reducing time of curing. (15) The major advantages of microwave heating over conventional heating are: (1) the inside and outside of substance are almost equally heated, and (2) temperature rise rapidly. (7)

The aim of this study was to evaluate the effect of the different curing methods, namely conventional water bath heat—curing and microwave energy polymerization method and also different immersion periods on the water sorption and solubility of the different types of heat—cured acrylic resins.

MATERIALS AND METHODS

The two different commercial heatcured acrylic resins were used in this study. These were Major Base 2 (M) (Major prodotti company, Italy) and Ouayle Dental (England) (QD). The resins wear mixed at powder/liquid ratio of (2.5/1) by weight. (16, 17) The resins specimens for the water sorption and solubility were prepared as disks with a dimension (50 \pm 1 mm in diameter and 0.5 ± 0.1 mm thikness). (18) Specimens were divided into two groups depending on the curing method, "Group I" water bath method, "Group II" microwave energy curing method, and the two different resins were used in each group. Eight specimens for each material in each group were prepared, so the final total number was thirty

two. For conventional water bath technique specimens cured with the conventional denture-flasking procedure. The specimens processed at 74 °C for 1.5hours, then the temperature of the water bath increase to boiling (100 °C) for additional one hour. (8) While for the curing cycle for microwave technique, a new Iraqi Fiber Reinforced Plastic (FRP) flask was placed in the microwave oven (Samsung, Model RE-570 D, 0.6 cuft, Korea) for 30 minutes at low setting (80 watts): 15 minutes per side, followed by 1.5 minutes at the high setting (500 watts). (19) Initially, specimens were dried over silica gel in a desiccator at 37 °C and weighed to an accuracy of 0.0001 g using an electronic balance (Mettler PM 460, Germany). This was considered to be the initial weight of the specimen (W1). Specimens then were immersed in distilled water, each specimen being in separate containers. The specimens subsequently were removed from their containers at 1 week and 1 month. Excess water was removed by blotting with filter paper and the weight of the specimen was recorded (W2) at each occasion. This represent the weight of the specimen after absorption of the distilled water. The procedure was repeated after interval of 1 week and 1 month. The amount of soluble materials lost was measured by drying the specimens in the desiccator after each absorption cycle and recorded as (W3). The percentage of absorption and solubility were determined as follows (18):

(1) Absorption % =
$$\frac{(W2 - W3) g}{(W1) g} \times 100$$

= $\frac{(\text{Weight after absorption - final weight after desication}) g}{(\text{Initial weight}) g} \times 100$
(2) Solubility % = $\frac{(W1 - W3) g}{(W1) g} \times 100$
= $\frac{(\text{Initial weight - final weight after desication}) g}{(\text{Initial weight}) g} \times 100$

Mean values and standard deviation wear calculated Mean values wear compared statistically with one way analysis of variance (ANOVA) followed by Duncan's multiple range test to determined the significant different among the groups at p< 0.05 level of significance.

RESULTS AND DISCUSSION

The results of the mean value of the water sorption and solubility in this study are shown in Tables (1 and 2). The results of ANOVA at p< 0.05 level of significance are shown in Tables (3–10) and the results of Duncan's multiple range test were represented in Figures (1–3). The results in Figure (1) represent the ratios of the water sorption and solubility of the QD and M resins in the tested groups Group I (water bath method) and Group II (microwave energy curing method) after 7 days of immersion. The ratios of the QD resin were higher than

that of the M resin but the difference was not significant as shown in Tables (3 and 4). Denture plastics of the same type may vary in water sorption because of the presence of additives. (11) This result come in agreement with the finding of Peter and Ernst and others (20, 21) when examined the physical properties of the different brands of heat—cured denture base resins that had different additives, the resins exhibit significant differences in residual MMA monomer, water up take and ratios of water sorption and solubility.

Table (1): Means value of the water sorption rate of the tested materials cured by two different curing methods for different immersion periods.

Tested groups			Water sorption r	ratio (weight %)			
		Quayle Do	ental resin	Major resin			
	No	7 days immersion	1 month immersion	7 days immersion	1 month immersion		
	•	Mean <u>+</u> SD	Mean <u>+</u> SD	Mean <u>+</u> SD	Mean <u>+</u> SD		
Group I	8	1.0168 <u>+</u> 0.2075	1.8221 <u>+</u> 0.8892	0.9864 <u>+</u> 0.1374	1.4853 <u>+</u> 0.1752		
Group II	8	1.3168 <u>+</u> 0.2085	2.9221 <u>+</u> 0.8792	1.2436 <u>+</u> 0.1813	2.3424 <u>+</u> 0.3979		

SD: Standard deviation; No: Samples number; Group I: Water bath curing method; Group II: Microwave energy curing method.

Table(2): Means value of the water solubility rate of the tested materials cured by two different curing methods for different immersion periods.

Tested groups			Water solubility 1	ratio (weight %)			
		Quayle D	ental resin	Major resin			
	No	7 days immersion	1 month immersion	7 days immersion	1 month immersion		
		Mean <u>+</u> SD	Mean <u>+</u> SD	Mean <u>+</u> SD	Mean <u>+</u> SD		
Group I	8	0.7192 <u>+</u> 0.6767	0.9821 <u>+</u> 1.0689	0.2743 <u>+</u> 0.1097	0.3484 <u>+</u> 0.1351		
Group II	8	1.5192 <u>+</u> 0.6778	1.5964 <u>+</u> 0.68	0.9886 <u>+</u> 0.161	1.5712 <u>+</u> 0.7251		

SD: Standard deviation; No: Samples number; Group I: Water bath curing method; Group II: Microwave energy curing method.

Table (3): ANOVA for the means of the water sorption ratio of the tested materials cured by different curing methods after 7days of immersion.

			Water	sorption r	atio (wo	eight %)	
Source of variance	df	Wate	er bath (Gr	oup I)	Microwave (Group II)		
		MS	F-value	<i>P</i> –value	MS	F-value	<i>P</i> –value
*Tested materials	1	0.0032	0.10	** 0.753	0.0188	0.49	** 0.496
Error	12	0.0312			0.0382		
Total	13						

df: Degree of freedom; MS: Mean square.

Table (4): ANOVA for the means of the water solubility ratio of the tested materials cured by different curing methods after 7 days of immersion.

			Water	solubility r	atio (v	weight %)	
Source of variance	df	Water bath(Group I)			Microwave(Group II)		
		MS	F-value	<i>P</i> -value	MS	F-value	<i>P</i> -value
*Tested materials	1	0.693	2.95	** 0.112	0.986	4.06	** 0.067
Error	12	0.235			0.243		
Total	13						

df: Degree of freedom; MS: Mean square.

Table (5): ANOVA for the means of the water sorption ratio of the different materials in the tested groups after 7 days of immersion.

Source of variance			Water	sorption ratio (weight %)				
	df	Qua	yle Dental	l resin	Major resin			
		MS	F-value	<i>P</i> -value	MS	F-value	<i>P</i> -value	
*Tested groups	1	0.3150	7.24	** 0.020	0.2314	8.95	** 0.011	
Error	12	0.0435			0.0259			
Total	13							

df: Degree of freedom; MS: Mean square.

Table (6): ANOVA for the means of the water solubility ratio of the different materials in the tested groups after 7 days of immersion.

			Water	solubility	ratio (weight %)	1	
Source of variance	df	Qu	ayle Denta	l resin	Major resin			
		MS	F-value	<i>P</i> –value	MS	F-value	<i>P</i> -value	
*Tested groups	1	2.240	4.88	** 0.047	1.7857	94.13	*** 0.000	
Error	12	0.459			0.0190			
Total	13							

df: Degree of freedom; MS: Mean square

^{*}Tested materials: Quayle Dental and Major heat cured acrylic resin .

^{**} Means are not have significant different at P≤0.05.

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^{**} Means are not have significant different at P≤0.05.

^{*} Group I (resins cured by water bath) and Group II (resins cured by microwave)

^{**} Means are significant different at $P \le 0.05$. *** Means are highly significant different at $P \le 0.01$.

Table (7): ANOVA for the means of the water sorption ratio of the different materials for different immersion periods cured by water bath method(Group I).

Source of variance	df	Qu	ayle Denta	l resin	Major resin			
		MS	F-value	<i>P</i> –value	MS	F-value	<i>P</i> –value	
*Immersion periods	1	2.270	5.44	** 0.038	0.8710	35.13	*** 0.000	
Error	12	0.417			0.0248			
Total	13							

df: Degree of freedom; MS: Mean square.

Table (8) ANOVA for the means of the water sorption ratio of the different materials for different immersion periods cured by microwave energy method (Group II).

Source of variance		Water sorption ratio (weight %)							
		Quayle Dental resin			Major resin				
		MS	F-value	<i>P</i> -value	MS	F-value	<i>P</i> -value		
*Immersion periods	1	9.019	21.63	** 0.000	4.2263	44.21	** 0.000		
Error	12	0.417			0.0956				
Total	13								

df: Degree of freedom; MS: Mean square.

Table (9): ANOVA for the means of the water solubility ratio of the different materials for different immersion periods cured by water bath method (Group I).

		Water solubility ratio (weight %)								
Source of variance	df	Qu	ayle Denta	l resin	Major resin					
		MS	F-value	<i>P</i> -value	MS	F-value	<i>P</i> –value			
*Immersion periods	1	0.242	0.30	** 0.593	0.0192	1.27	*** 0.282			
Error	12	0.800			0.0151					
Total	13									

df: Degree of freedom; MS: Mean square.

^{*}Immersion periods: 7 days and 1 month immersion period.

^{**} Means are significant different at P≤0.05

^{***} Means are highly significant different at P≤0.01

^{***} Means are significant different at P≤0.05.

^{*}Immersion periods: 7 days and 1 month immersion period.

^{**} Means are highly significant different at P≤0.01.

^{*}Immersion periods: 7 days and 1 month immersion period.

^{**} Means are not significant different at P≤0.05.

^{***} Means are significant different at P≤0.05.

Table (10) ANOVA for the means of the water solubility ratio of the different materials for different immersion periods cured by microwave energy method (Group II)

		CHCIE	zy miemou (Oroup II).					
		Water solubility ratio (weight %)							
Source of variance	df	Quayle Dental resin			Major resin				
		MS	F-value	<i>P</i> -value	MS	F-value	<i>P</i> -value		
*Immersion periods	1	0.021	0.05	** 0.835	1.188	4.31	** 0.060		
Error	12	0.461			0.276				
Total	13								

df: Degree of freedom; MS: Mean square.

^{**} Means are not significant different at P≤0.05.

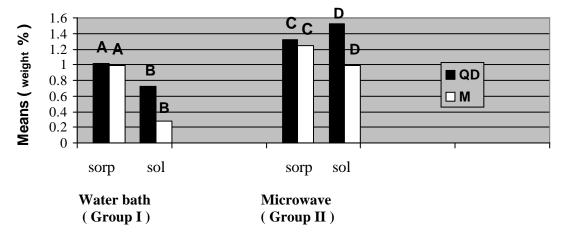


Figure (1): Duncan multiple rang test for the water sorption and solubility ratios of the different materials(QD and M) in the tested groups after 7 days of immersion. Means with the same letter have no significant difference at p < 0.05; Sorp: Water sorption rate.Sol: Water solubility rate; QD: Quayle Dental heat-cured acrylic resin; M: Major heat-cured acrylic resin.

The result in Figure (2) represent the ratio of the water sorption and solubility of the tested resins after 7 days of immersion in the tested groups. The ratios for the QD and M resins in Group II (microwave energy method) were higher than that of the Group I (water bath method), the differentce was significant as shown in Tables (5and 6). This finding is disagreement with the finding of Reitz et al., and others (22-24) in that the two polymerization methods microwave irradiation and heat activation did not cause differences in the properties of the resins. The explanation of increasing the ratios in Group II is that the microwave oven for curing resin was much faster than a conventional water bath, and degree of curing also increased a little. Microwaves act only on the monomer, which decreases in the same proportion as polymerization increases. Microwave cause the methyl methacrylate monomer molecules within the acrylic resin to orient themselves in electromagnetic field at frequency 2450 MHz, and numerous polarized molecules are flipped over rapidly and generate heat due to molecular friction. Initiating radicals are then able to react with monomers to start polymerization. Microwave processing temperatures beyond 100.3 °C caused vaporization of the monomer and produced porosity and the voids in the structure of the cured resin will increase water sorption. (7, 24, 25)

^{*}Immersion periods: 7 days and 1 month immersion period.

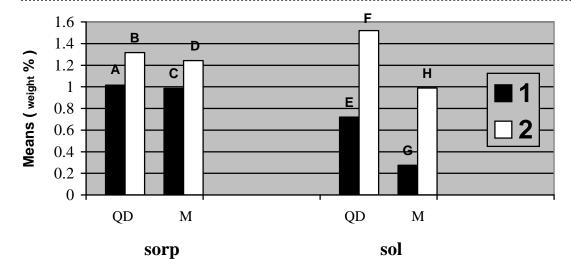


Figure (2): Duncan multiple rang test for the effect of the curing methods on the water sorption and solubility ratios of each material (QD and M) in the tested groups after 7days of immersion.

Means with the different letter have a significant difference at p < 0.05; (1): Water bath curing method (Group I); (2): Microwave energy curing method (Group II); Sorp: Water sorption rate.Sol: Water solubility rate; QD: Quayle Dental heat-cured acrylic resin; M: Major heat-cured acrylic resin.

The results in Figure (3) showed the effect of the immersion period on the ratios of the water sorption and solubility of the different resins (QD and M) in the tested groups. The ratios after one month were higher than after 7 days. The difference was significant except that the difference of the water solubility ratio of the QD resin in Group I and of the two resins in Group II was not as shown in Tables (7-10). The significant differences indicated that the water uptake and leaching of the soluble mass of the resin are continues process in a period reached to 1 month. The water sorption is gradual process taking many months to complete. (9) This result come in agreement with the finding of Stafford et al (21) in that the resins expanded during the storage in water for one month. The result of the effect of the immersion period on the solubility disagreement with the finding of Varpu and Pekka ⁽⁶⁾ in that the largest amount of residual monomer usually leached from denture base PMMA within the first few days of water storage.

CONCLUSION

The results of this study appeared that the curing method and immersion period have an effect on the water sorption and solubility ratios of the different types of heatcured acrylic resin, QD and M resins. The results appeared after 7 days of immersion the ratios of the water sorption and solubility of the QD were higher than that of M when cured by both conventional water bath and microwave processing. Also the ratios of the both resin were significantly higher when cured by microwave processsing method than when cured by conventional water bath method. The result also appeared that the ratios of the two resins after one month were higher than after 7 days when cured by both curing methods and the difference was significant except that the difference of the water solubility ratio of QD in Group I and that of the two the resins in Group II was not.

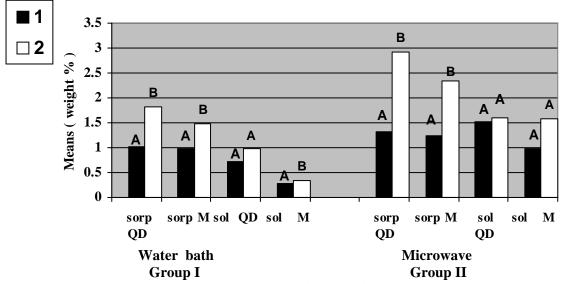


Figure (3): Duncan multiple rang test for the effect of immersion period on the water sorption and solubility ratios of (QD and M) materials in the tested groups. Means of the ratios with different letters for each resin have a significant difference at p < 0.05; (1): 7 days of immersion; (2): 1 month of immersion; Sorp: Water sorption rate; Sol: Water solubility rate. QD: Quayle Dental heat-cured acrylic resin; M: Major heat-cured acrylic resin.

REFERENCES

- 1. Skinner EW. Science of Dental Materials. 8th ed. W.B. Saunder Co. Philadephia, USA. 1982; p: 244.
- 2. McCabe JF. Anderson's Applied Dental Materials. 6th ed. Blackwell Scientific Publications, London,UK. 1985; P: 84.
- 3. Craig RG, O'Brien WJ, Powers JM. Dental Materials. 6th ed. Mosby Co. Missouri, USA. 1996; p: 258.
- 4. Chen S, Liang W, Yen P. Reinforcement of acrylic denture base by incorporation of various fibers. *J Biomed Mater Rest.* 2001; 58: 203–208.
- Widad A, Reta M, Abd Alwahid K. Filler reinforced acrylic denture base material. Part 2– Effect of water sorption on dimensional changes and transverse strength. *J Coll Dentistry*. 2005; 17(1): 6–10.
- 6. Varpu M, Vallittu PK. Water sorption and solubility of glass fiber–reinforced denture polymethyl methacrylate resin. *J Prosthet Dent.* 1996; 76(5): 531–534.
- 7. Lai CP, Tsai MH. Morphology and properties of denture acrlic resins cured by microwave energy and conventional water bath. *Dent Mat.* 2004; 20: 133–141.
- 8. Craig RG. Restorative Dental Materials.

- 11th ed. Mosby Co. Missouri, USA.2002; p: 656.
- Van Nort R. Introduction to Dental Materials. 2^d ed. Mosby Co. Missouri, USA. 2002; P: 96–112.
- 10. American Dental Association Specification. Guide to dental materials and device.7th ed. 1975; p: 205.
- 11. Craig RG. Restorative Dental Materials. 10th ed. Mosby Co. Missouri, USA. 1997; p: 513.
- 12. Varpu M, Katja K. Water sorption solubility, and effect of post–curing of glass fiber reinforced polymers. *Biomaterials*. 1999; 20: 1187–1194
- 13. Al-Haydary ASA. The use of microwave energy in polymerization of radio-opaque denture base polymers. MSc Thesis. Collage of Dentistry. University of Baghdad. 1992.
- 14. Garcia RC, Junior JA, Rached RN. Effect of denture cleanser on the surface roughness and hardness of a microwave cured acrylic resin and dental alloys. *J Prosthet De*nt. 2004; 13: 173–178.
- 15. Azzari MJ, Cortizo MS. Effect of the curing conditions on the properties of an acrylic denture base resin microwave–polym-

- erised. J Dent. 2003; 31: 463-468.
- Mitchell L, Mitchell DA. Oxford Handbook of Clinical Dentistry. 3rd ed. Oxford University Press. 1999; Pp:700–701.
- 17. Jagger D, Harrison A, Milward P. The effect of the addition of poly (methyl methacrylate) fibers on some properties of high strength heat—cured acrylic resin denture base material. *J Oral Rehabil*. 2003; 30: 231–235.
- 18. Kazanji MN, Watkinson AC. Soft lining materials: their absorption of, and solubility in, artificial saliva. *Brit Dent J.* 1988;165: 91–94.
- 19. Radhwan HH. Comparison of some physiccal properties of acrylic denture base material cured by water bath and microwave techniques. *Al–Rafidain Dent J.* 2003; 3(2): 143–147.
- 20. Peter P, Ernst UR. Residual methyl methacrylate monomer, water sorption, and water solubility of hypoallergenic denture base

- materials. *J Prosthet Dent.* 2004; 92(1): 72–78.
- 21. Stafford GD, Bates JF, Huggett R. A review of some denture base polymers. *J Dent*. 1980; 8(4):292–306.
- 22. Reitz PV, Sanders JL, Levin B. The curing of denture acrylic resins by microwave energy: physical properties. *Quint Int* 1985; 16: 547–551
- 23. Derrick LW, Daniel BB, Steven AA. Effect of polyethylene fiber reinforcement on the strength of denture base resins polymerized by microwave energy. *J Prosthet Dent.* 1994; 72:635–638.
- 24. Capagnoni MA, Barbosa DB. The effect of polymerization cycles on porosity of microwave–processed denture base resin. *J Prosthet Dent.* 2004; 91(3): 281–285.
- 25. John C and Eugene P. Porosity reduction and its associated effect on the diametral tensile strength of activated acrylic resins. *J Prosthet Dent.* 1985; 53(3): 374–379.