





EFFECTIVENESS OF VARIOUS DENTAL BLEACHING TECHNIQUES- AN IN VITRO STUDY

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Introduction

Stained or discolored teeth make it troublesome for a person to look and feel good. Bleaching is a conservative and efficacious practice in dentistry to whiten discolored and stained teeth. At present, home bleaching (10% Carbamide peroxide) and in-office bleaching $(25\% H_2O_2)$ can be served with a number of available whitening methods.

Various light sources have been used for activation of bleaching agents in power bleaching. However, the effect of light energy source in activation of bleaching gel remains unclear.

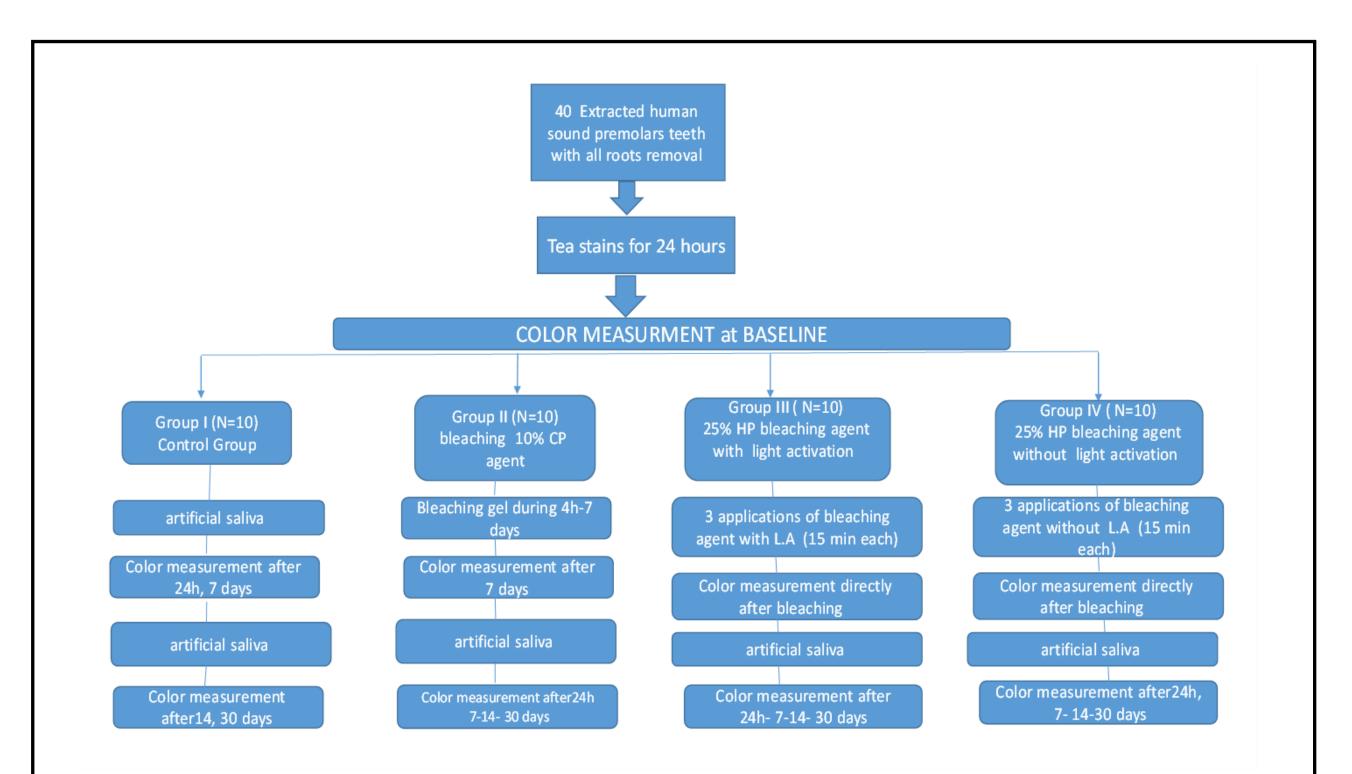
Aim of the study

Hence, this study aimed to evaluate the efficacy of home bleaching agent (10% Carbamide Peroxide) and in-office bleaching agent (25% hydrogen) peroxide) with and without light activation.

Materials and methods

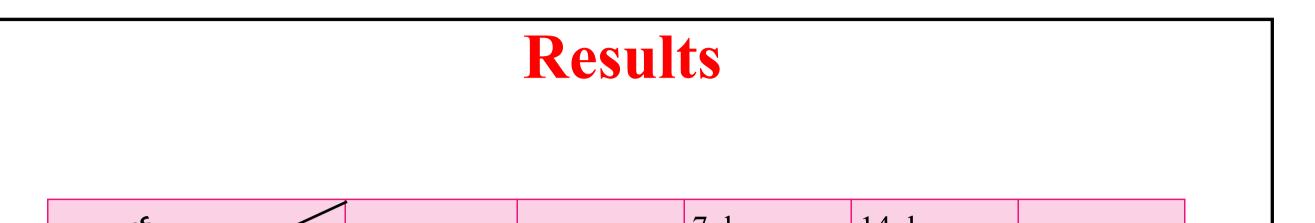
Ethical Approval

Ethical approval was obtained from the research center of Riyadh Elm University. Study was registered with number FPGRP/43430006/128. Study design



Statistical analysis

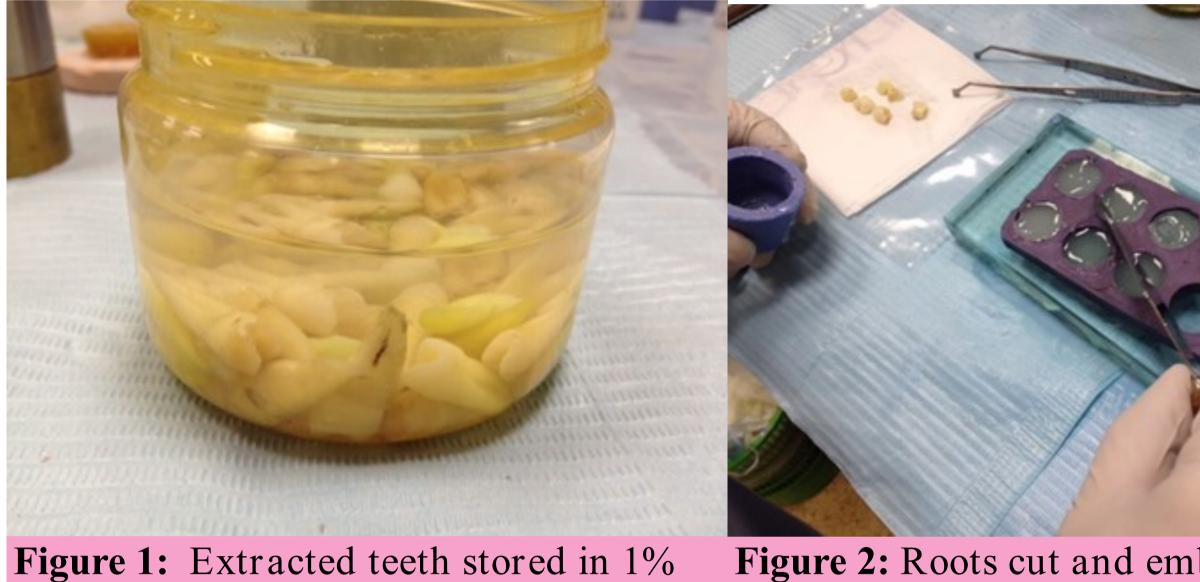
Color change related to time was measured using repeatedmeasures ANOVA. One-way ANOVA with post hoc Tukey's multiple comparison tests.



It was a lab based study that compared the effectiveness of (10%) Carbamide Peroxide bleaching agent and (25%) hydrogen peroxide bleaching agent with light activation by Zoom In Office Light Activator (LED-Philips Zoom, Discus Dental, Inc. USA, 400-505 nm (violet color)) and without light activation.

Specimen teeth collection

Sixty (60) human sound premolars teeth extracted for orthodontic purposes in Rivadh Elm University Hospitals were collected and (40) of them were used in the experiment.



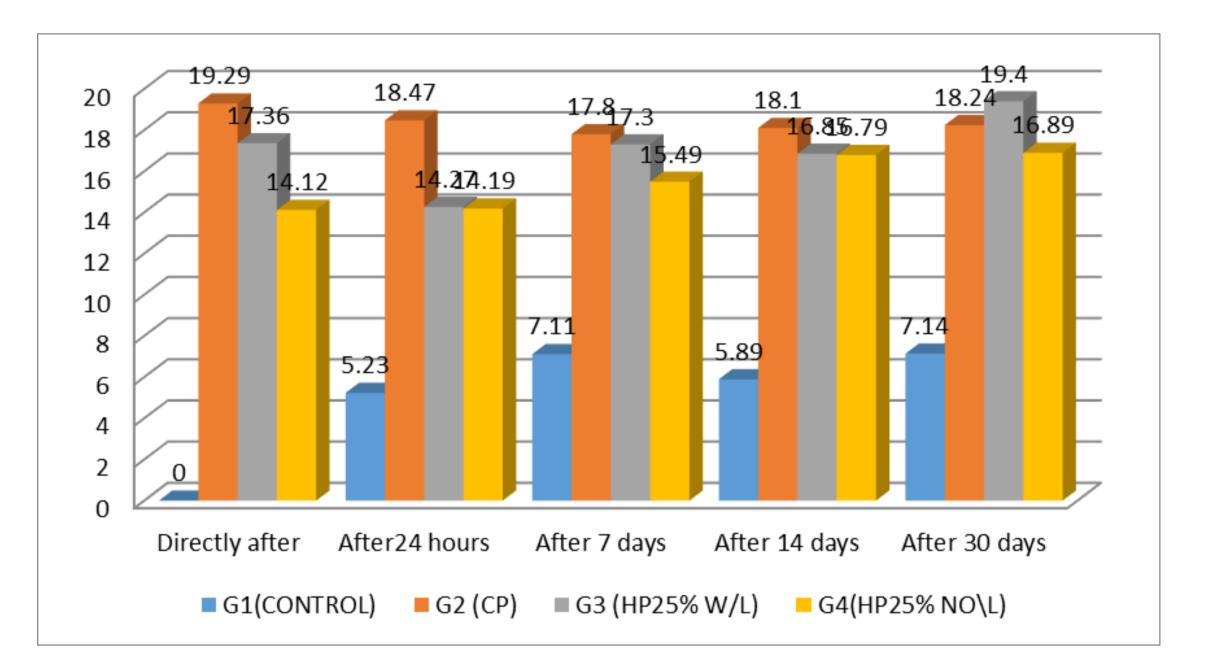
Thymol solution

Figure 2: Roots cut and embedded in orthoresin



GROUPS JODS	Direct After bleaching	24 hours after bleaching	7 days	14 days	30 days	
GROUP PERIODS			After bleaching	After bleaching	after bleaching	
CONTROL. No bleaching	_	5.23 (2.85)	7.11 (3.88)	5.89 (3.35)	7.14 (3.24)	
10% Carbamide Peroxide.	19.29 (6.67)	18.47 (7.70)	17.80 (6.17)	18.10 (6.04)	18.24 (5.55)	
25% Hydrogen peroxide with light activation	17.36 (4.02)	14.27 (3.88)	17.30 (2.69)	16.85 (3.01)	19.41 (4.11)	
25% Hydrogen peroxide without light activation	14.12 (3.86	14.19 (4.63)	15.49 (4.87)	16.79 (6.09)	16.89 (6.09)	

Mean of ΔE values and Standard Deviations (SD) for all groups at each evaluation period



Mean of ΔE values at different time intervals

Table 6: Mean of ΔE values, Standard Deviations (SD) and <i>P</i> . values of a evaluation periods							
Carolos activos	Direct	24 hours	7 days	14 days	30 days		
	After	after	after	after	after		
	bleaching	bleaching	bleaching	bleaching	bleaching		

Figure 3: Immersion of sample crown in orthoresin block

Figure 4: Sample after complete setting of orthoresin ready for bleaching

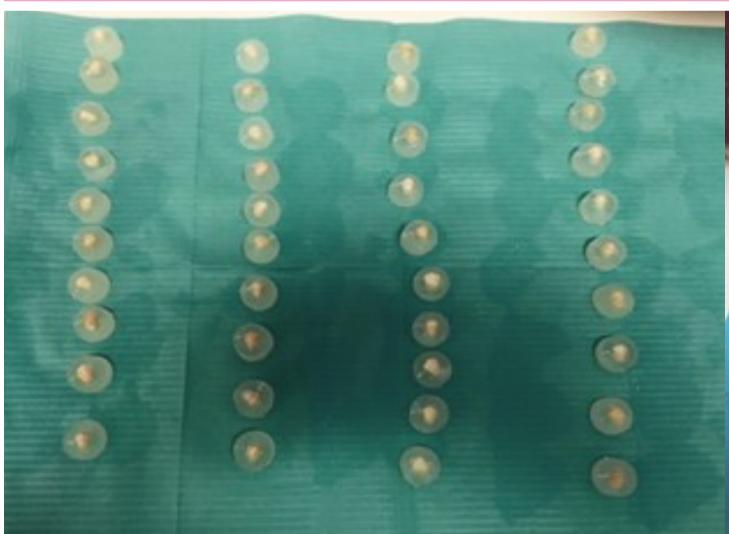


Figure 5: Samples randomly divided into four groups



Figure 6: Coding and grouping of samples in separate containers

Stain development

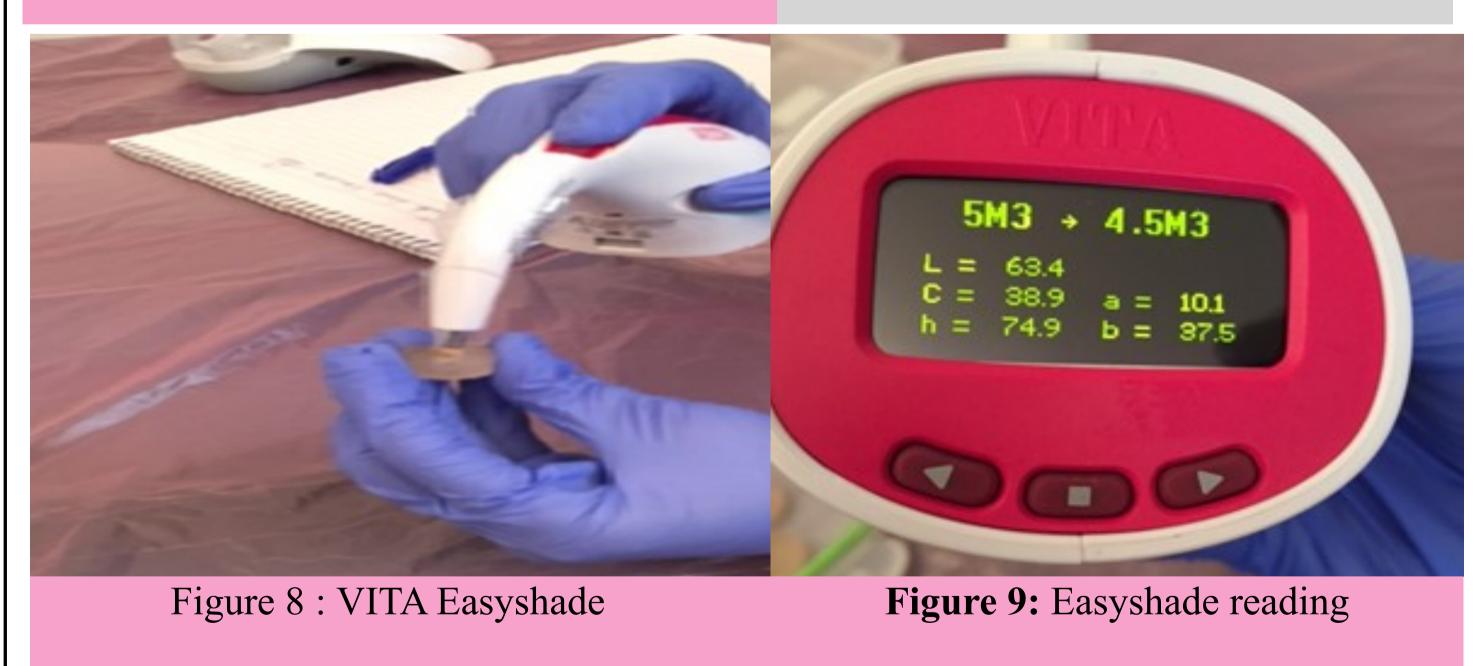




Color measurements were based on the Commission Internationale de l'Eclairage L*a*b* mode. In this mode, the L* represents the light value (brightness), the a* represents either green (- a*) or red (+ (a*), and the b* represents either blue $(-b^*)$ or yellow $(+b^*)$. The difference between the color coordinates was calculated as

 $\Delta E = \{ (\Delta L^*) 2 + (\Delta a^*) 2 + (\Delta b^*) 2 \} \frac{1}{2}$

Figure 7: Direct staining of crowns by using a standardized tea solution



*G1	-	5.23(2.85)	7.11(3.88)	5.89(3.35)	7.14(3.24)
**G2	19.29(6.67)	18.47(7.70)	17.80(6.17)	18.10(6.04)	18.24(5.55)
Р.	-	.000	.000	.000	.000
*G1	-	5.23(2.85)	7.11(3.88)	5.89(3.35)	7.14(3.24)
***G3	17.36(4.02)	14.27(3.88)	17.30(2.69)	16.85(3.01)	19.41(4.11)
Р.	-	.002	.000	.000	. 000
*G1	-	5.23(2.85)	7.11(3.88)	5.89(3.35)	7.14(3.24)
****G4	14.12(3.86)	14.19(4.63)	15.49(4.87)	16.79(6.09)	16.89(6.09)
Р.	-	.002	.001	.000	.000
**G2	19.29(6.67)	18.47(7.70)	17.80(6.17)	18.10(6.04)	18.24(5.55)
***G3	17.36(4.02)	14.27(3.88)	17.30(2.69)	16.85(3.01)	19.41(4.11)
Р.	1.000	.442	1.000	1.000	1.000
**G2	19.29(6.67)	18.47(7.70)	17.80(6.17)	18.10(6.04)	18.24(5.55)
****G4	14.12(3.86)	14.19(4.63)	15.49(4.87)	16.79(6.09)	16.89(6.09)
Р.	.070	. 412	1.000	1.000	1.000
***G3	17.36(4.02)	14.27(3.88)	17.30(2.69)	16.85(3.01)	19.41(4.11)
****G4	14.12(3.86)	14.19(4.63)	15.94(4.87)	16.79(6.09)	16.89(6.09)
Р.	.629	1.000	1.000	1.000	1.000

*G1 (none bleaching group), **G2 (bleaching by 10% Carbamide Peroxide) ***G3 (bleaching by 25% HP with light activation (LED \ZOOM) ****G4 (bleaching by 25% HP without light activation).

Discussion and Conclusions

In this study, the degree of bleaching obtained via the in-office technique (25% HP, three applications for 15 minutes each) was similar to that obtained by the home bleaching technique (10%) CP/4 hours/7 days) regardless of the use of a light source.

This result is in line with the findings of Sulieman et al (2005), and contradictory to the Zekonis et al (2003) and Auschill et al (2005).

 ΔE values did not differ significantly between in-office bleaching technique using light activation and use of bleaching gel without light activation. This finding is similar to that reported by by Kugel et al (2006), in which in-office tooth whitening with light activation did not exhibit benefit over the chemically activated tooth whitening system after a 2-weeks. Similar finding was reported by Marson et al (2008) hydrogen peroxide did not show improvement with the use of any light sources tested (halogen light, LED, LED/Laser). High concentration HP (25%) is used during in office bleaching, light may not contribute much to the bleaching results. Our results contradict with findings of Browning and Swift (2011), Dominguez et al (2011) that demonstrated the efficacy of the association of light sources with in office bleaching systems. Within the limitations of this study, it can be concluded that using (25%) hydrogen peroxide bleaching agent (irrespective of using light activation) gave same effects as the prolonged application of (10%) Carbamide Peroxide bleaching agent. However, optimal bleaching can be achieved with any techniques used in this experiment.

Sample groups



Figure 10: (G1)Samples of control group in artificial saliva

to 10% CP bleaching agent.

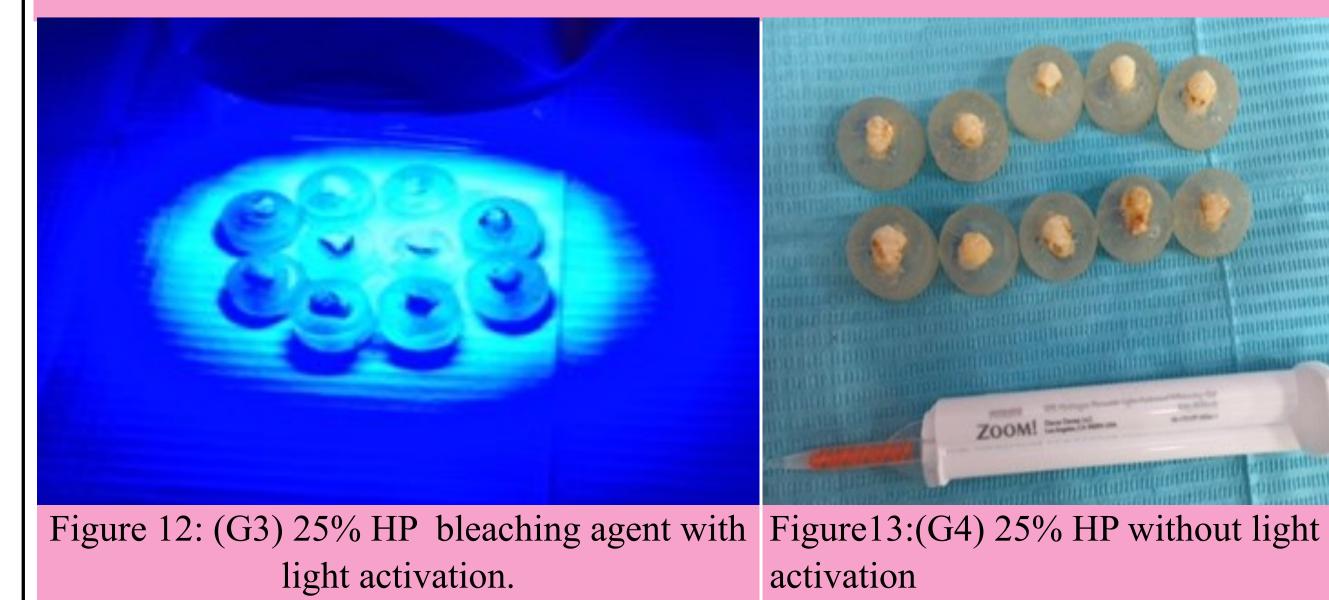


Figure 11: (G2) Group samples' subjected

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Recommendations

Light activation is not required during bleaching Similar studies with different bleaching materials are required Further clinical trials needed to measure effect of light on the pulp temperature.