



COMPARISON OF VARIOUS DESENSITIZING TOOTHPASTE – A CLINICAL STUDY

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ABSTRACT **BACKGROUND:** Dentine hypersensitivity is a commonly encountered problem that is triggered by an external stimuli. Various treatment modalities have been proposed, amongst them desensitizing paste is used extensively.

OBJECTIVE: Main objective of the study was to compare the efficacy of three commercially available desensitizing tooth paste-Sequel F^(R), Hiora k^(R), Vantej^(R).

MATERIALS AND METHODS: 30 subjects of dentin hypersensitivity were randomly selected and divided into 3 groups, desensitizing pastes were prescribed. Baseline values were recorded by tactile method and air blast test using visual analogue scale rating from 0-10. Efficacy was compared on baseline, 10 days and after 30 days.

STATISTICAL ANALYSIS: Wilcoxon signed rank test and Kruskal Wallis test was done to compare the efficacy of sensitivity reduction between the three groups and during different time interval (P value was set at 0.05).

RESULTS: Comparison of all the three pastes showed positive results within 30 days but Vantej^(R) showed statistically significant changes within 10 days indicating that it had rapid action.

CONCLUSION: All the tooth paste have convincing evidence in reducing dentinal hypersensitivity. Long term studies to facilitate better understanding can be advocated in the future.

KEYWORDS : Dentinal Hypersensitivity, Tooth Paste, Periodontal Therapy, Visual Analogue Scale, Hydrodynamic Theory.

INTRODUCTION:

Dentinal hypersensitivity is characterized by short sharp pain arising from exposed dentin in response to thermal, evaporative, tactile, osmotic or chemical stimuli. Canadian Advisory Board in 2003 suggested that Dentinal Hypersensitivity should be substituted for pathological condition and thus Dentin hypersensitivity can be identified as a distinct clinical entity¹. Dentinal hypersensitivity is a painful clinical condition that affects 8-57% of adult population which is majorly associated with dentin exposure to the oral environment^{2,3}. Several studies have reported non-cariou lesions and dentine hypersensitivity in adult population with prevalence rates ranging from 5-85%⁴ and 2-8 % to 74%⁵. These variations in percentages are due to differences in populations habits, dietaries and methods of investigations. The disease is prevalent with the age range of 20-50 years with higher prevalence for females^{1,6,7}. It is manifested in a manner that is physically and psychologically uncomfortable for the patient. Dentin exposure may result from various causes like excessive tooth brushing, exposure of non-bacterial acids in diet, periodontal therapy, and excessive occlusal force leading to tooth wear. The most widely accepted theory for dentinal hypersensitivity is hydrodynamic theory proposed by Brannstrom⁸. Clinical management of dentinal hypersensitivity is based on proper diagnosis, severity of condition, whether localized or generalized, eliminating the possible etiological causes, patient counselling about oral hygiene practices, diet and harmful habits.

There are two common methods to determine the intensity of dentine hypersensitivity one of them is through questionnaire and other is through clinical examination. The prevalence distribution of dentine hypersensitivity in the first method is estimated higher than the second method. The severity or degree of pain can be quantified using visual analogue scale or categorized scale (slight, moderate and severe). Another commonly used method is the "Schiff" scale⁹. It has been observed that some people with dentine hypersensitivity do not pursue treatment of the disease as they do not consider dentine hypersensitivity as a specific disease. However they seek professional

help when the intensity of pain is severe and affect their daily activity. Various strategies have been developed to treat dentin hypersensitivity. Desensitizing methods aim to control hydrodynamic mechanisms of pain which can be broadly divided into three categories:

- 1) Agents or products that reduce fluid flow within dentinal tubules themselves thereby blocking the tubules. eg. hydroxyapatite, calcium sodium phosphosilicate.^{10,11}
- 2) Those that interrupt the neural response to stimuli eg. potassium nitrate^{12,13,14,15,16}
- 3) Coagulation and protein precipitation of plasma in the dentinal fluid eg. Lasers¹

However, there is no gold standard for treatment of Dentinal hypersensitivity. Hence the objective of the study was to determine the efficiency of three different desensitizing paste Sequel F^(R) (5 %potassium nitrate and sodium monofluorophosphate 0.7% and triclosan 0.3%) Hiora -K^(R) (palakya 10 mg, Triphala 6 mg, Trikatu 4mg), Vantej^(R) (Novamin glass particle containing Calcium sodium phosphosilicate with 25% sodium, 25% Calcium, 6-8 % phosphate and remainder silica).

MATERIALS AND METHOD –

The present study was randomised control single blind conducted at outpatient department of A.J Institute of dental Sciences. A total of 30 patients (18 male and 12 female) who visited the department participated in the study. Informed consent was obtained from each patient. They were randomly divided into 3 groups –Group A - Sequel F^(R), Group B - Hiora^(R) and Group C - Vantej^(R).

INCLUSION CRITERIA-

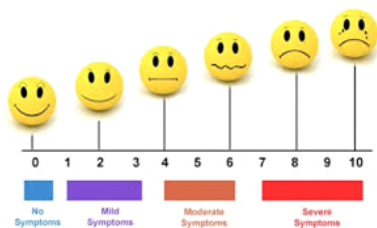
- Patients with generally good health with minimum of two hypersensitive teeth with cervical abrasion
- Gingival recession less than 1mm loss of dentine in depth which did not require any restoration or treatment,
- Compliant patients.

EXCLUSION CRITERIA:

- Advanced periodontal disease, gross carious lesions, mobility.
- Gross oral pathology
- Medical condition, pregnant or lactating women and
- Patient using other desensitizing toothpaste were excluded from the study.

Baseline sensitivity values were recorded before starting the treatment by tactile method and air blast test using visual analogue scale which was repeated after 10 and 30 days. Tactile sensitivity test was assessed using probe with slight manual pressure in mesio-distal direction on hypersensitive areas of tooth. Sensitivity was also assessed by air blast method where the evaporative stimuli was applied in the form of cold air from a dental unit syringe (at 20 ± 3 °C at 40 to 65 psi) for 1sec from a distance of 1 cm from the affected area. The degree of hypersensitivity was recorded according to VAS. The Scores were given on 10 cm sensitivity VAS which has ratings from:

- 0-1: no pain
- 2-3: mild pain
- 4-6: moderate pain
- 7-10: severe pain.



Subjects >5 on VAS were included in the study. The effectiveness of two times application of desensitizing paste for period of 30 days among the subjects were analysed.

RESULTS:

Wilcoxon signed rank test and Kruskal Wallis test was done to compare the efficacy of sensitivity reduction between the three groups and during different time interval. (P value was set at 0.05).

There was no report of any side effects on usage of either of the dentifrices.

In the present study, highly significant reduction of both objective and subjective symptoms was observed from baseline to 30 days (Table A and Table B)

Table A

Parameter: Objective assessment

Group	N	Median	IQR	Minimum	Maximum	Friedman test Value	p value
Group A	DAY 0	10	3.00 (2-3)	1	3	16.545	.000 HS
	10 DAYS	10	1.50 (1-2.25)	0	3		
	30 DAY	10	1.00 (0.75-1)	0	2		
Group B	DAY 0	10	2.00 (2-2)	2	3	14.000	.001 HS
	10 DAYS	10	2.00 (1-2)	1	2		
	30 DAY	10	1.00 (1-1)	1	2		
Group C	DAY 0	10	2.00 (1.75-2.25)	1	3	12.000	.002 HS
	10 DAYS	10	1.00 (0-2)	0	2		
	30 DAY	10	1.00 (0-2)	0	2		

Table B

Parameter: Subjective assessment

Group	N	Median	IQR	Minimum	Maximum	Friedman test Value	p value
Group A	DAY 0	10	2.50 (2-3)	1	3	15.548	.000 HS
	10 DAYS	10	2.00 (1-2)	1	2		
	30 DAY	10	1.00 (1-1)	0	2		
Group B	DAY 0	10	2.00 (2-2)	1	3	12.000	.002 HS
	10 DAYS	10	1.50 (1-2)	1	2		
	30 DAY	10	1.00 (1-1)	1	2		
Group C	DAY 0	10	2.00 (2-3)	1	3	13.130	.001 HS
	10 DAYS	10	1.00 (0.75-2)	0	2		
	30 DAY	10	1.00 (0-2)	0	2		

PAIR WISE COMPARISON OF OBJECTIVE ASSESSMENT WAS DONE BY WILCOXON SIGNED RANK TEST (TABLE C)

- During 0-10 days significant reduction was noted in all the 3 groups.
- In 10-30 days significant reduction was noted only with group A and group B and no statistically significant changes with group C.

- During 0-30 days statistically highly significant reduction was noted with group A and group B and group C showed significant reduction

Table C

Pairwise comparison by Wilcoxon signed rank test

Group	10 DAYS - DAY 0		30 DAY - DAY 0		30 DAY - 10 DAYS	
	p value		p value		p value	
Group A	.011	sig	.004	HS	.020	sig
Group B	.025	sig	.004	HS	.025	sig
Group C	.024	sig	.024	sig	1.000	

PAIR WISE COMPARISON OF SUBJECTIVE ASSESSMENT WAS DONE BY WILCOXON SIGNED RANK TEST (TABLE D)

- In 0 -10 days highly significant changes were noted with group A and statistically significant changes were noted with group C and no statistically significant changes were noted with group B.
- In 10-30 days' time period significant reduction was noted with group A and group B and no significant changes with group C.
- 0-30 days highly significant changes were noted in group A and group B and significant changes in group C.

Table D

Pairwise comparison by Wilcoxon signed rank test

Group	10 DAYS - DAY 0		30 DAY - DAY 0		30 DAY - 10 DAYS	
	p value		p value		p value	
Group A	.008	HS	.006	HS	.020	sig
Group B	.059		.007	HS	.046	sig
Group C	.014	sig	.016	sig	.157	

However according to the results of this study no statistically significant difference was seen within the 3 groups at 0 day, 10 day and 30 days (Table E and Table F-Kruskall Wallis test)

GROUPWISE COMPARISON BY KRUSHALL WALLIS TEST Table E

Parameter: Subjective assessment

Group	N	Median	IQR	Minimum	Maximum	Kruskall wallis test	p value
Group B	10	2.00 (2-2)	1	3			
Group C	10	2.00 (2-3)	1	3			
10 DAYS	Group A	10	2.00 (1-2)	1	2	2.578	.276 NS
	Group B	10	1.50 (1-2)	1	2		
	Group C	10	1.00 (0.75-2)	0	2		
30 DAY	Group A	10	1.00 (1-1)	0	2	.191	.909 NS
	Group B	10	1.00 (1-1)	1	2		
	Group C	10	1.00 (0-2)	0	2		

Table F

Parameter: Objective assessment

Group	N	Median	IQR	Minimum	Maximum	Kruskall wallis test	p value
Group B	10	2.00 (2-2)	2	3			
Group C	10	2.00 (1.75-2.25)	1	3			
10 DAYS	Group A	10	1.50 (1-2.25)	0	3	1.956	.376 NS
	Group B	10	2.00 (1-2)	1	2		
	Group C	10	1.00 (0-2)	0	2		
30 DAY	Group A	10	1.00 (0.75-1)	0	2	.720	.698 NS
	Group B	10	1.00 (1-1)	1	2		
	Group C	10	1.00 (0-2)	0	2		

DISCUSSION:

Dentin hypersensitivity is a common clinical condition in permanent teeth characterized by an exaggerated response to non-noxious stimuli. It is mainly caused by loss of enamel or cementum leading to the exposure of dentine to the oral environment¹⁸. There have been various materials and methods proposed to reduce and treat dental hypersensitivity. These include the uses of dentifrices , composites, lasers ,bioglass and so on.

The desensitizing tooth paste is one of the most commonly and widely used method for dental hypersensitivity because of its low cost ,ease of use and application. These materials usually exert their effect through sealing the dentinal tubules thereby reducing the transmission of nerve impulse.

Thus the goal of the study was to determine the efficacy of three desensitizing tooth paste - Sequel F[®], Hiora K[®], Vantej[®].

The potassium nitrate present in Sequel F[®] inhibits repolarisation of sensory nerve endings thereby reducing pain¹⁹. According to Wilchgers and Ermert and Kim potassium nitrate has an effective desensitizing action. It is believed that the increase in the concentration of extracellular potassium around the nerve fibres causes their depolarization, avoids repolarisation and blocks the axonic action and passage of nerve stimulus, thus inactivating the action potential²⁰. Markowitz K and Kim also stated that a prolonged period of depolarization results in inactivation of action potential thus divalent cation solution stabilize the nerve membrane without changing the membrane potential²¹. Shreya et al stated that subjects treated with 5% potassium nitrate in the present study showed significant improvement in hypersensitivity scores thus claimed the efficacy of the agent in reducing hypersensitivity²². In this present study, thereby significant reduction of symptoms with Sequel F[®] from baseline to 30 days. Hiora K[®] contains herbs which act on the tooth and have protective and antimicrobial action. Present study showed significant efficacy of Hiora K[®] toothpaste in sensitivity reduction as there was remarkable symptomatic relief noted in all the patients. A study conducted by Sukumaran et al stated that Hiora-K[®] was safe and it effectively reduced dental hypersensitivity²³. Novamin is a bioactive glass/ceramic with calcium sodium phosphosilicate as an active ingredient which works on the principle of tubule occlusion by infiltration of precipitation products which when exposed to aqueous media provides calcium and phosphate ions that form HCA with time. Combination of HCA layer results in physical occlusion of dentinal tubules which relieves hypersensitivity²⁴. A review of literature by Gendreau et al based on randomized controlled clinical trials, support the use of novamin tooth paste formulation in providing relief of pain from dental hypersensitivity²⁵.

A comparative study by Parkinson and Wilson concluded that calcium sodium phosphosilicate imparts significant level of dentinal occlusion thus with durable occlusive deposits following 4 days of twice daily brushing in vitro²⁶. A study conducted by Aditya et al stated that novamin crystals have been highly effective in treatment of dentine hypersensitivity pain²⁷. Narong dij et al stated that calcium sodium phosphosilicate tooth paste showed a greater benefits at an early stage compare to potassium nitrate²⁸. Results with Vantej[®] in the present study also revealed that statistically significant changes were noted from 0-10 days indicating that it had a rapid action. In our study, the comparison of all the three toothpastes showed positive results within 30 days. However the placebo effect should be taken into consideration while comparing between the desensitizing agents because of the individual variability in pain response in dental hypersensitivity studies we need to include large sample size to get statistically significant results²⁹. The formation of reparative dentin may also attribute to the occlusion of dentinal tubules and reduction of sensitivity with time. All these factors must be considered while analysing the results.

CONCLUSION:

All the toothpastes have convincing and supporting evidence for efficacy in reducing dental hypersensitivity. Long term studies to facilitate better understanding of performance of these desensitizing toothpaste can be advocated in the future. Inclusion of a placebo group to determine whether the results were due to placebo effect might be beneficial.

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