

Original Research Article

**Erosive effects of Pediatric liquid medicinal syrups with remineralising agents on primary enamel: An in vitro comparative study.**

**Abstract :**

**Background:** The use of liquid medicinal syrups in childhood to the reasons that children suffer from illness quite often. The acidic component in the formulations may cause erosive to the dental tissues. **Aims:** The aim of study was to test the erosivity of common Paedtric medications and to assess the change of the erosive potential of the medications in the presence of commercial Remineralilising agents. **Materilas and methods:** A total of 27 extracted/exfoliated non carious Deciduous molars were used. 8 medical syrups were used.

**Statistical Analysis used:** NOVA and post hoc analysis were applied. An Tukey alpha value of 0.05 was employed.

**Conclusion:** The knowledge of the erosive potential of commonly used syrups is mandatory as erosion in children teeth.

**Key words;** Deciduous molars, Erosion, Liquid medicinal syrups.

## Introduction

Dental erosion is defined as a form of irreversible non carious tooth loss, which results in chemical dissolution of mineral from the tooth when the PH of the oral environment falls below 5.5.<sup>[1]</sup>

It has been increasingly recognized as a problem of all ages of particular concern is the wearing of tooth enamel that is caused by dietary erosion, which is usually observed in individuals who consume fruit juices and carbonated soft drinks and in those on long term acidic medications.<sup>[2]</sup>

Dental erosion is increasingly recognised as a condition affecting the younger age groups, and in previous studies on the primary dentition a high prevalence of up to 82% has been reported.

Dental erosion has been found to be a common cause of tooth wear and it may be the main contributing factor in severe tooth wear, more than attrition and abrasion.<sup>[3]</sup> Liquid oral medicines are usually prescribed for children to aid compliance.<sup>[4]</sup> Acidic preparations are often necessary for drug dispersion, chemical stability maintenance, to ensure physiological compatibility and to improve flavor.<sup>[4,5]</sup>

In addition to the acidic components, other factors such as prolonged and frequent ingestion (i.e., two or more times daily), bedtime and between meals consumption, high viscosity and the collateral effect of reduced salivary flow, may contribute to increase the risk for medication-induced dental erosion.<sup>[6]</sup>

## **Materials & methods;**

The study was approved by the Institutional Ethical Committee (IERB,MDC 8/2020) A total of 27 extracted/non carious deciduous molars were used .Before use, the teeth were hand scaled and cleaned. The teeth were checked for structural abnormality that could possibly interfere with results, and on this basis, the teeth selection was done.

The roots were removed at the Cemento enamel junction. The crowns were fixed in a wax block parallel to the surface. The specimen before immersion cycle was stored at 37 ° C in Artificial saliva for 24 h.

The 8 commonly available liquid medicaments (PLM) were selected for the study and their endogenous PH was measured using a PH electrode meter (Eutech PH 700).

### **PH and TA measurements.**

The PH of each Pediatrics liquid medicaments was measured using a PH electrode meter. (Microelectrode,Inc,new Hampshire, USA),Which was calibrated at the start of every session using buffer standards of PH 4 and PH 7 (Islab,wertheim,Germanay).Every drink was tested with standard volume (20 ml) from a freshly opened bottle. Every experiment was performed in triplicate at room temperature (22 c ) and the mean and standard deviation obtained .phosphate – buffered saline (PBS) was employed as the control.

The TA of each drink was assayed using a standard protocol of acid-base reaction <sup>[7]</sup>.In brief, 0.1 M Sodium hydroxide (NaoH) was progressively added to 20 ml of each drink until neutralization (PH7) was attained using the same electrode .The volume of 0.1 M NaoH required was recorded.

## **PH and TA acidity measurements of representative drinks after addition of Remineralising agents.**

Representative items [Table 2] were randomly selected from the each category in [Table 1] and were tested to examine the effects after the addition of various Remineralising agents.

These remineralising agents were tooth mousses.

1. TM; GC Corporation, Tokyo, JAPAN.
2. Tooth mousses plus. (TMP ,GC Corporation, Tokyo).
3. Clinpro (3 M, Minnestota, USA).
4. 1.23% NAF and Artificial saliva. (AS).

To stimulate intraoral conditions where there is dilution of the applied paste by saliva, TM, TMP and Clinpro TM were applied as slurries. [1 g of paste to 10 ml distilled water. Neutral NAF solution was applied as a 1.23% wt/vol solution .TM, TMP, Clinpro TM ,NAF and AS were individually added to each representative drink at the following concentration.

The PH and TA were recorded after 5 month to as certain the effects were due to dilution; test with control (PBS) solution was performed.

All experiments were performed in triplicate.

### **Statistical analysis;**

The students t test (two-tailed), one way ANOVA were performed on instant 3 (Graph pad software, California, USA). An Tukey alpha value of 0.05 was employed.

## RESULTS

[TABLE 1] The PH of each Pediatric liquid medicament was measured using a PH electrode meter. The PH ranged between 3.09 to 5.7. The minimum PH value minimum for S1 syrup (3.09) & maximum for S8 (5.7).

[Table 2][Graph 1]After adding the Reminerasing agents the PH value is changes to  $> 5.5$ .

[Table 3][Graph 2]The TA of each Pediatric liquid medicament was minimum for S5 and maximum for S4 .After adding Remineralising agents the TA value changes.

### Surface changes of primary enamel

Primaryenamel surface treated with various PLMs, and artificial saliva was observed under the SEM at 1 min, 10 min, and 8 h.

In most of the medicaments, etched prism pattern and crater formation were observed on the primary enamel surface immersed in medicaments at 1 min, 10 min, and 8 h time intervals and sporadic rod ends were seen at the end of 1 min time interval in **Tonoferol**only[Fig 1]. In **artificial saliva**, no surface changes were observed on primary enamel surface in SEM [Fig 2].

**In antibiotic group** (Ofloxacin, Cefixime, and Albendazole), In Ofloxacin etchedprism pattern was seen after 1 min interval [Fig 3] and crater formation was seen after 10 min and 8h interval, respectively, [Fig 4]whereas, in Albendazole, nothingsignificant was found after1 min interval, but etched prism pattern was seen after 10 min and crater formation was seen after 8 h interval. [Fig 5].

**In analgesics group** (Paracetamol and Ibuprofen), etched prism pattern was observed after 1 min interval and crater formation was seen after 10 min and 8 h interval in both the medicaments, [Fig 6] respectively.

In Nimulide group sporadic rod ends were seen only in this group after 1 min interval. Etched prism pattern was seen after 10 min and crater formation was seen after 8 h interval. [Fig 7]

In **Anti histamine group** (Diphenyl hydramines) no significant changes on primary enamel surface were found after 1 min and 10 min interval, where as crater formation was seen after 8 h interval in both the medicaments [Fig 8].

## Discussion

Dental erosion is increasingly recognized as common conditions in Pediatric dentistry with complications of dental sensitivity, altered aesthetics and loss of occlusal vertical dimension, eating difficulties, pulp exposure and abscess. <sup>[8,9]</sup>

The high levels of consumption of acidic drinks are likely to be associated with the alarming prevalence of erosion among the pre-school children reported in the studies of Millward etal <sup>[10]</sup> and Al-malik etal <sup>[11]</sup>

Interestingly, all Pediatric medications examined in the present study exhibited PH in a similarly range between 3.09 to 5.7 .This PH range of the Pediatric medications is in accordance with the handful of previously papers published on prescription medications. <sup>[12,13]</sup>

Therefore, Paediatric medications have PH values below the critical PH for Demineralization occur.

The findings of present study are similar to those done by Babu KLG etal <sup>[14]</sup> which examined the PH of 8 medications ranging from 6.05 to 6.77 and its effects on primary teeth.

In a study conducted by Greenwood ME etal <sup>[15]</sup> the liquid syrup Dimetapp (Brompheramine and Phenylephrine) had an acidic PH of 2.86 .The PH of the artificial saliva in the present study was found to be 7.1.

The other difference between deciduous and permanent tissues may also be of importance. For example deciduous teeth demonstratates a higher degree of enamel porosity and lower degree of mineralization than permanent teeth. In addition, deciduous enamel has repeatedly been noted to have a higher content of Co2 and carbonated as well lower content of phosphorus than the permanent tissue. <sup>[16]</sup>

Deciduous teeth are more susceptible to caries like acid attack than permanent teeth invitro according to Feather stone and Melberg. <sup>[17]</sup>

Even though the PH of PLM was not near the critical PH of the oral cavity, erosion of PES was evident when subjected to study under SEM. This is in agreement with the studies of Babu KLG et al<sup>[14]</sup> and Green wood ME et al<sup>[15]</sup> who used SEM to evaluate the Erosion potential of liquid syrup on rats enamel and human extracted primary teeth respectively

In the present study Etched prism pattern were seen in enamel surface treated with Azithromycin syrup for 10 min and Crocin syrup for 1 minute and Multivitamin syrup for 10 minutes. These findings were in agreement with findings of Babu KLG et al<sup>[14]</sup> who reported similar typical prism pattern on enamel surface treated with Amoxicillin for 1 minutes, Theophylline for 1 minutes and Multivitamin for 10 minutes.

Enamel surface treated with crown suspension for 1 minutes Ibugesic plus suspension for 1 minute, Mox Red mix suspension for 1 minute, Although suspension for 10 minutes and 8 hours Valparin 200 syrup for 10 minutes and 8 hours Visyneral syrup for 1 minute and 8 hours and delicious syrup for 1 minute and 8 hours all showed the etched prism pattern but this was not of the classical appearance. This atypical appearance of enamel surface can be supported by the study done by Grando LJ et al<sup>[18]</sup> who stated that complexity can be seen in the different patterns of loss of enamel structure in eroded Deciduous teeth.

In the present study there seemed to be no direct relationship between the PH of the selected PLMS and the Erosive changes seen on the PES. This findings is in accordance with that of Babu KLG et al. <sup>[14]</sup>

Morch Tet al<sup>[19]</sup> and Onose H et al<sup>[20]</sup> reported that the Chelating agents present in PLMS like sodium salt of various aminoacids and lactate at or near neutral PH, increase the uptake of radioactive phosphorus by loss of ca from enamel. This factor may be responsible for the erosive pattern seen in the surface of PES in the present study.

In the present study, demonstrated that the addition of Remineralizing agents like TM, TMP, neutral NAF, Clinpro and AS resulted in significant changes in PH and TA compared to the control group.

This present study was similar to study done by Manton et al<sup>[21]</sup> The addition of 0.2% CPP-ACP to acidic drinks showed that a slight increase in PH (0.5 units) was associated with a significant reduction in erosive potential.

This present a study result is in agreement with study by Dawes et al<sup>[22]</sup> which suggests an inverse proportion relationship between PH and Ca and phosphate concentration.

CPP-ACP consists of biologically active peptide casein phosphor peptides that holds and stabilizes Ca and phosphate that holds and stabilizes Ca and Phosphate ions in a bio available form, and prevents precipitation. These ions become available at the tooth surface when the PH drops, so that Demineralization is inhibited<sup>[23]</sup>.

Maguire et al<sup>[5]</sup>, Arora et al<sup>[24]</sup> Therefore, PH, TA are not the only parameters in determining the erosive potential .Furthermore, different conc of active ingredients in medicines may indirectly affect the PH,TA due to the different levels of acids added to mask the taste of the active ingredients

## **Conclusion**

In conclusion, the present study demonstrates that pediatric over-the counter medications have erosive potential. It also demonstrates that commercially available Remineralising products, Tooth mouses, Tooth mouse plus, 1.23% Neutral sodium fluoride, and Artificial saliva have the potential to reduce the erosivity of pediatric over the counter medications.

**COMPETING INTERESTS DISCLAIMER:**

Authors have declared that they have no known competing financial interests OR non-financial interests OR personal relationships that could have appeared to influence the work reported in this paper.

## References

1. Taji S, Seow WK . A literature review of dental erosion in children. Aust Dent J 2010;55:358-67.
2. Maguire A, Baqir W, Nunn JH. Are Sugar-Free Medicines More Erosive Than Sugar-Containing Medicines? An In-vitro Study of Paediatric Medicines with Prolonged Oral Clearance Used Regularly and Long-Term by Children. Int J Paediatr Dent 2007;17:231-8.
3. Johansson AK, Sorvari R, Birkhed D, Meurman JH. Dental Erosion in Deciduous Teeth: An In-vivo and In-vitro Study. J Dent 2001;29:333-40.
4. Nunn JH, Ng SK, Sharkey I, Coulthard M. The dental implications of chronic use of acidic medicines in medically compromised children. Pharm World Sci 2001;23:118-19.
5. Maguire A, Baqir W, Nunn JH. Are sugars-free medicines more erosive than sugar-containing medicines? An in vitro study of pediatric medicines with prolonged oral clearance used regularly and long-term by children. Int J Paediatr Dent 2007;17:231-38.
6. Scatena C, Galafassi D, Gomes-Silva JM, Borasatto MC, Serra MC. In vitro erosive effect of Pediatric medicines on deciduous tooth enamel. Braz Dent J .2014;25:22-7.
7. Hunter L, Patel S, Rees J . The in vitro erosive potential of a range of baby drinks. Int J Pediatric Dent .2009;19:325-29.
8. Luo A, Zeng XJ. The prevalence of dental erosion . J Dent 2005;33:165-70.
9. Harding MA, Whelton H. Community Dental health. 2003;20:165-70.
10. Millward A, Shaw L, Smith A . Dental erosion in four-year-old children from differing socioeconomic backgrounds. ASDC J Dent Child. 1994;61:263-66.

11. Al-Malik MI, Holt RD, Bedi R . Erosion, caries and rampant caries in preschool children in Jeddah, Saudi Arabia. *Community Dent Oral Epidemiol* .2002;30:16-23.
12. Nunn JH, Ng SKF, Sharkey I, Coulthard M .The dental implications of chronic use of acidic medicines in medically compromised children. *Pharm World Sci*.2001;23:118-19.
13. Neves BG, Farah A, Lucas E, de Sousa VP, Maia LC . Are Paediatric medicines risk factors for dental caries and dental erosion? *Community Dent Health* .2010;27:46-51.
14. Babu KL, Rai K, Hedge AM . Pediatric liquid medicaments - do they erode the teeth surface? An in vitro study: Part I. *J Clin Pediatr Dent* .2008;32:189-94.
15. Greenwood ME, Feigal R, Messer H. Cariogenic potential of liquid medications in Rats. *Caries Res* 1984;18:447-9.
16. Maria Angelica Huebde menezes oliveria, Carolina Paestorres. Microstructure and mineral composition of dental enamel of permanent and deciduous teeth. *Microscopy Research and technique*.2010;73:572-77.
17. Featherstone JD/Melberg JR. Relative rates of progress of artificial carious lesions in bovine and human enamel. *Caries Res* .1981;15:109-14.
18. Grando LJ, Tames DR, Cardoso AC.etal. In vitro study of enamel erosion caused by soft drinks and lemon juice in deciduous teeth analyzed by Stereo microscopy and Scanning electron microscopy. *Caries Research* 1996;30:373-8.
19. Morch T, Punwani I, Greve E. The Possible Role of Complex Forming Substances in the Decalcification of the Caries Process. *Caries Res* 1971;5:135-43..
20. Onose H, Sandham HJ. PH Changes during Culture of Human Dental Plaque Streptococci on Mitis Salivarius Agar. *Arch Oral Biol* 1976;21:291-6.
21. Manton DJ, Cai F, Yuan Y, Walker GD, Cochrane NJ, et al. Effect of Casein phosphopeptide-Amorphous calcium phosphate added to acidic beverages on enamel erosion in vitro. *Aust Dent J*.2010;55:275-79.
22. Dawes C. What is the critical pH and why does a tooth dissolve in acid? *J Can Dent Assoc* 2003;69:722-24.
23. Lussi A .Dental erosion - novel Remineralizing agents in prevention or repair. *Adv Dent Res*.2009;21:13-16.

24.Arora R, Mukherjee U, Arora V . Erosive potential of sugar free and sugar containing pediatric medicines given regularly and long term to children. Indian J Pediatr.2012;79:759-63.

**Table 1: Summary of PH of Pediatric medications in different Remineralising agents.**

Type	Brand name	Generic name	Company name.	PH
1.Ironsupplement		Tonoferol.S1	Cipla	3.09
2.Antibiotics	1.Oflox-100	Ofloxacin.S2	Cipla	5.05
	2.Taxim-o	Cefixime.S3	Alkem.	4.6
	3.Pexoclav	Amoxicillin+	Apex	5.07
	4.Bendex.	Clavonic.S4 Albendazole.S5	Cipla.	5.46
3.Analgesic	1.Combiflam	Ibuprofen.S6	Sanafi	4.56
	2.P-250	Paracetamol.S7	Apex.	5.35
4.Decongesant (Antihistamine)	Maxtra	Phenylephrine HCL.S8	Zunventus.	5.7

**Table 2: Pair wise comparison of PH of Paediatric medications before & after adding Remineralising agents by Tukeys multiple posthoc procedures.**

Agents	Mean Difference	Std. Error	p-value
Control vs Tooth mouse	-1.45	0.36	0.0030*
Control vs Clinpro	-0.87	0.36	0.1380
Control vs Artificial saliva	-0.83	0.36	0.1700
Control vs 1.23% NAF	-0.67	0.36	0.3570
Tooth mouse vs Clinpro	0.58	0.36	0.5150
Tooth mouse vs Artificial saliva	0.61	0.36	0.4510
Tooth mouse vs 1.23% NAF	0.77	0.36	0.2300
Clinpro vs Artificial saliva	0.04	0.36	1.0000
Clinpro vs 1.23% NAF	0.20	0.36	0.9820
Artificial saliva vs 1.23% NAF	0.16	0.36	0.9920

<b>Agents.</b>	<b>Mean Difference</b>	<b>Std. Error</b>	<b>p-value</b>
Control vs Tooth mouse	0.39	0.41	0.8660
Control vs Clinpro	0.13	0.41	0.9980
Control vs Artificial saliva	0.12	0.41	0.9980
Control vs 1.23% NAF	0.04	0.41	1.0000

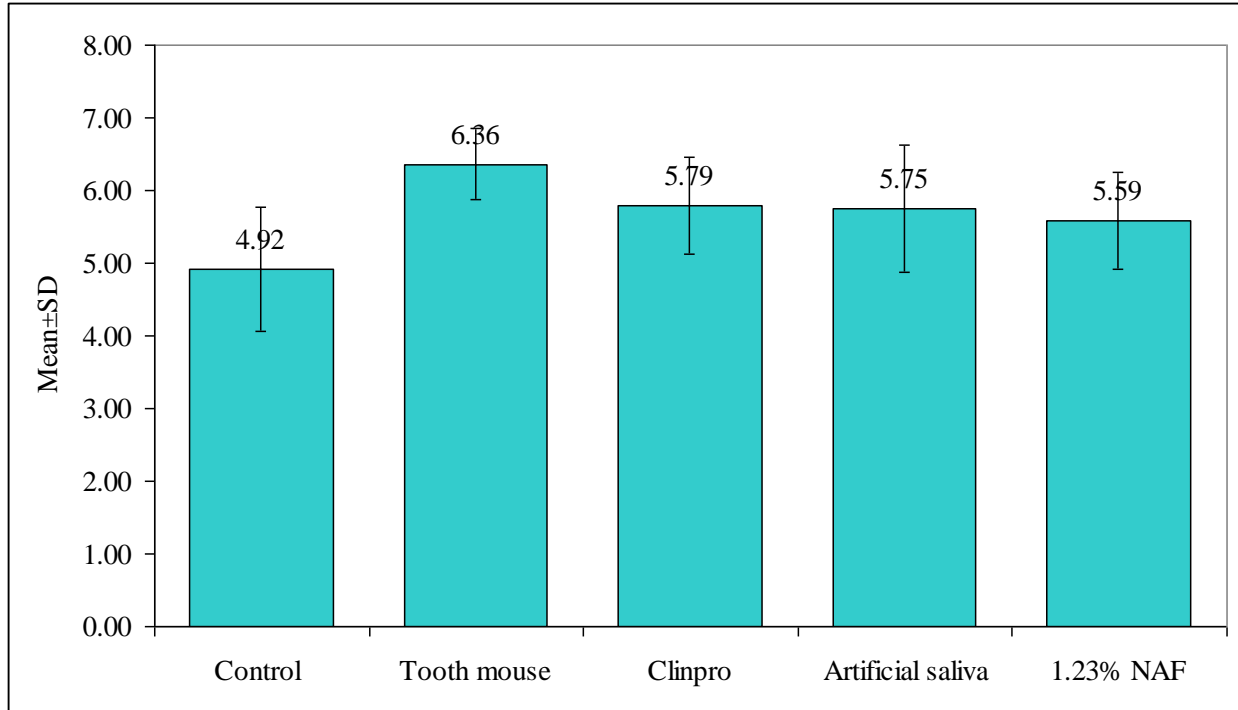
**Table 3: Pair wise comparison of TA of Paediatric medications before & after adding**

Tooth mouse vs Clinpro	-0.27	0.41	0.9650
Tooth mouse vs Artificial saliva	-0.28	0.41	0.9590
Tooth mouse vs 1.23% NAF	-0.35	0.41	0.9060
Clinpro vs Artificial saliva	-0.01	0.41	1.0000
Clinpro vs 1.23% NAF	-0.09	0.41	0.9990
Artificial saliva vs 1.23% NAF	-0.08	0.41	1.0000

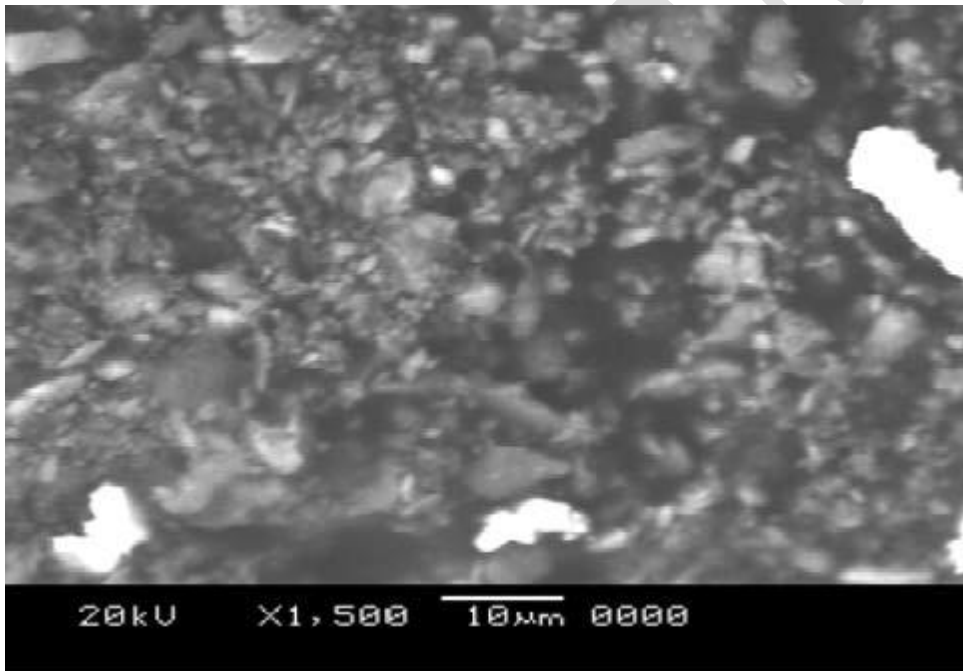
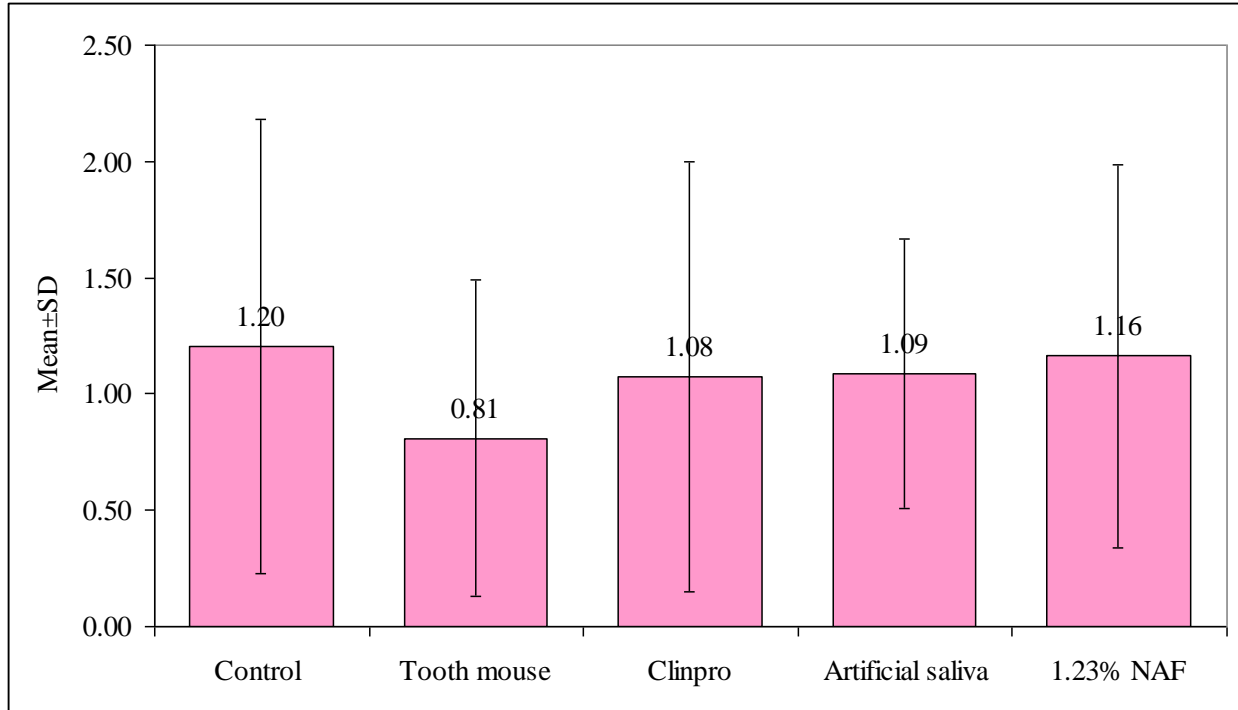
**Remineralising agents by Tukeys multiple posthoc procedures.**

**GRAPHS;**

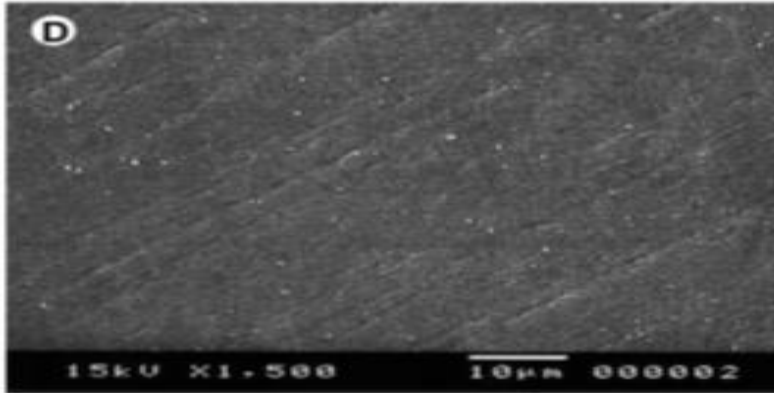
[Graph: 1]. Comparison of PH of pediatric medications before & after adding remineralising agents



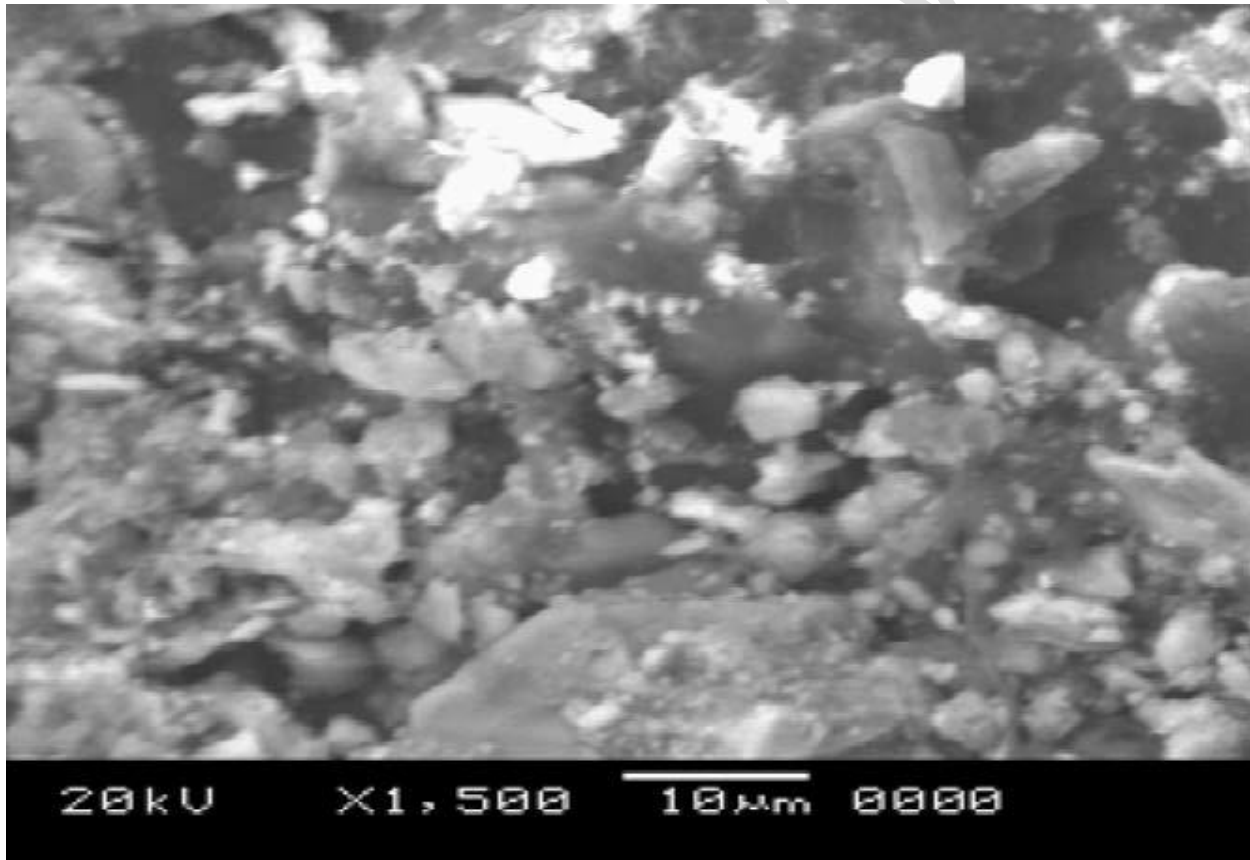
[Graph:2] Comparison of TA of paediatric medications before & after adding remineralising agents.



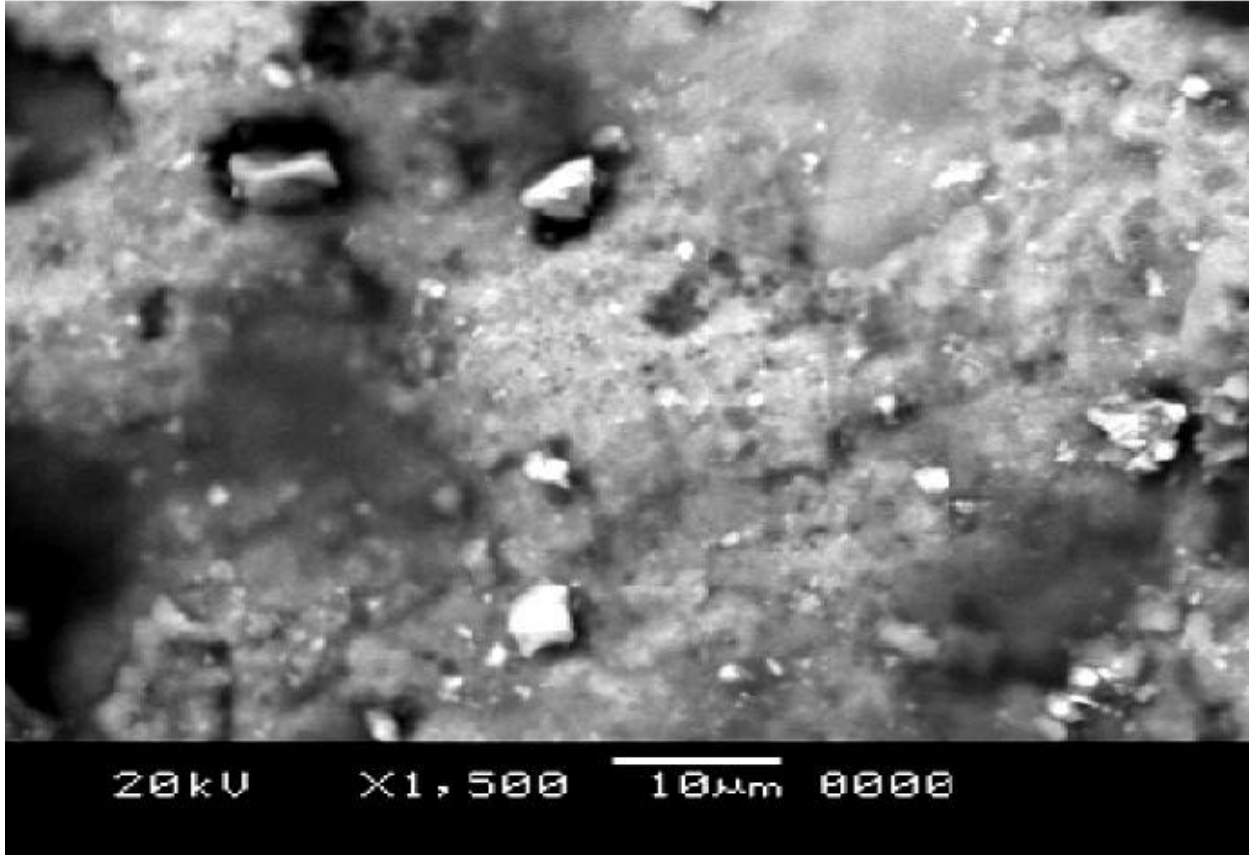
[Fig 1] Sporadic rod ends seen on primary enamel surface after immersing **Tonoferol** for 10 min.



[Fig 2] In artificial saliva, no surface changes were observed on primary enamel surface in SEM.

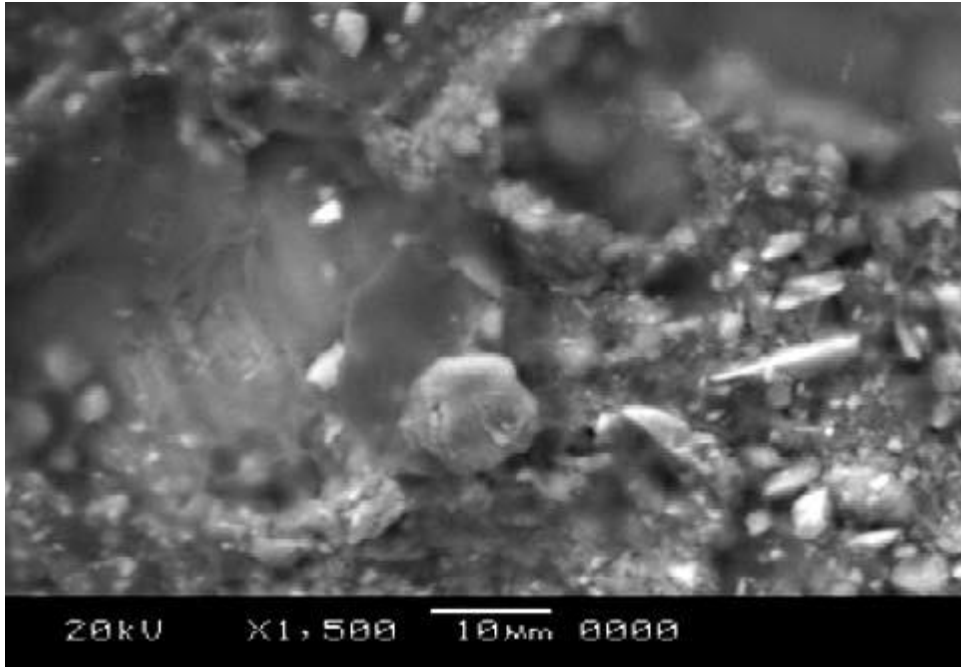


[Fig 3] Etched prism pattern seen on primary enamel surface after immersing in Ofloxacin with zinc for 1 min.

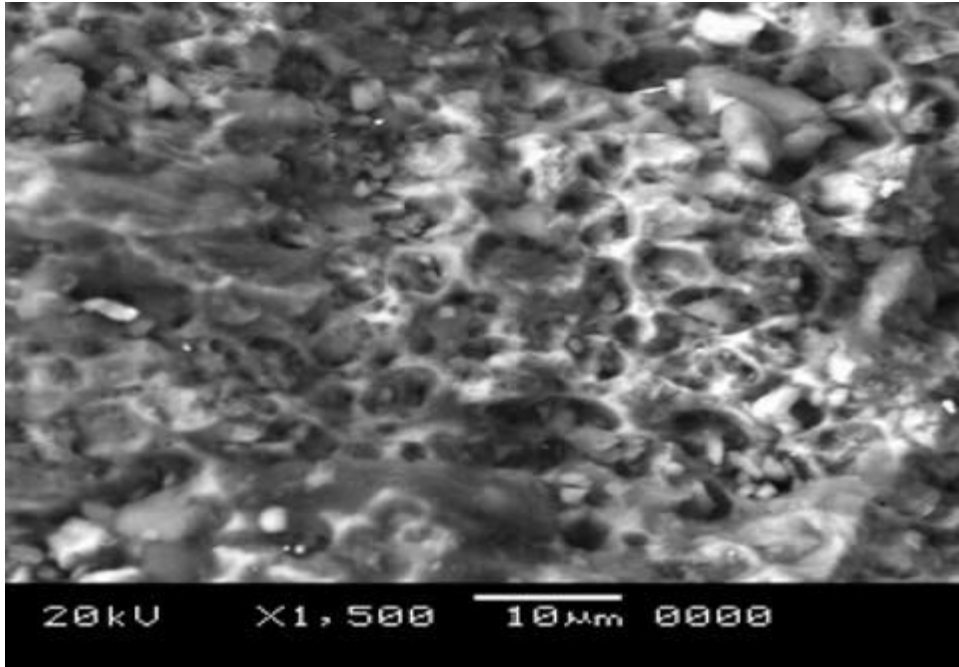


[Fig 4]. Crater formation seen on primary enamel surface after immersing in Albendazole for 10 mins.

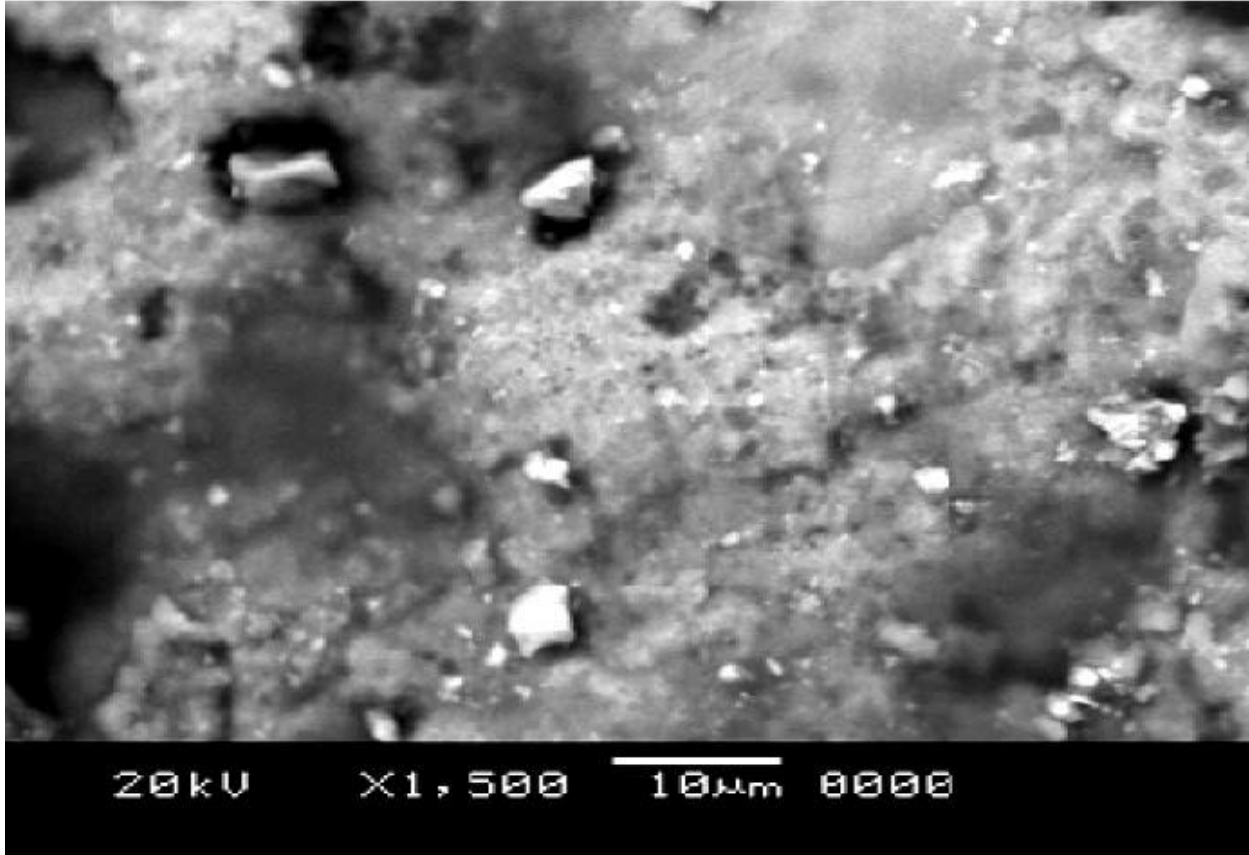
UNDER REVIEW



[Fig 5]Crater formation seen on primary enamel surface after immersing in **Albendazole** for 8 h.

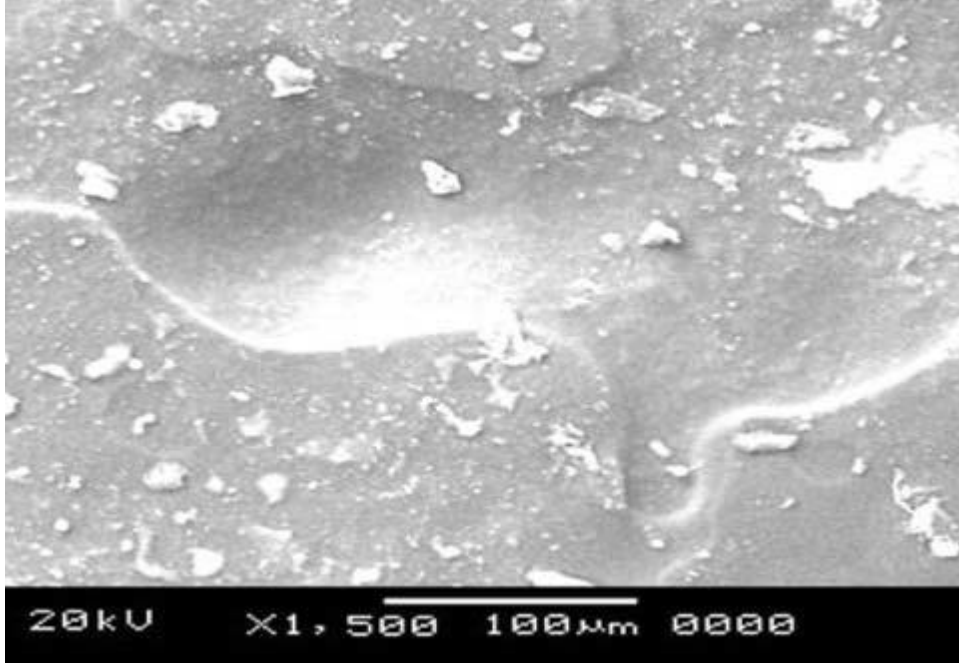


[Fig 6] Etched prism pattern seen on primary enamel surface after immersing in Crocin with Zinc for 1 min.



[Fig 7]. Crater formation seen on primary enamel surface after immersing in Nimesulide for 10 mins.

UNDER PEE



[Fig 8] Crater formation was seen after 8 h interval in Antihistamine group.