

Evaluation of Pain and Tenderness in Endodontic Treatment of Deciduous Teeth using (LSTR) 3-Mix MP therapy

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Abstract:

Aim and objective: The purpose of this study was to evaluate the prognosis of pain and tenderness in endodontic treatment of deciduous teeth using LSTR 3-MixMP therapy.

Materials and method: The treatment was performed on selected 60 patients of 60 teeth which were divided into two groups; Group-A(n=30) and Group-B(n=30). In group-A patients were diagnosed initial pulpitis or reversible pulpitis (vital inflamed pulp) and in group-B patients were diagnosed as non-vital necrosed pulp with or without inter-radicular lesion. In both groups carious lesion and previous restorative materials were removed and the cavity was prepared in such a way which was termed as "Medication cavity". The cavity was cleaned and dried and then LSTR 3Mix MP therapy was placed at the orifice of root canals or on the bottom of the pulp chamber and then sealed with glass-ionomer cement and further reinforced with composite resin. Resolution of clinical signs and symptoms; pain and tenderness to percussion were evaluated at one week, one month, six months and one year interval.

Result: In all cases clinical symptoms such as pain and tenderness to percussion disappeared after treatment in both groups but group B showed better clinical success. Finally all the cases were evaluated as successful.

Conclusion: Based on the present study it can be concluded that endodontically involved deciduous teeth in both vital (inflamed) and non-vital teeth associated with pain and tenderness were treated successfully by LSTR 3-Mix MP therapy.

Key Words: Endodontic treatment: Deciduous tooth, 3-Mix MP therapy.

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Introduction

Endodontically involved deciduous teeth due to carious lesion are associated with numerous bacteria. Bacteria present in Carious lesions, together with reactive inflammatory and immunological host responses, lead to the clinical development of pulpitis, pulp necrosis, periapical infections, abscess. The combination of inflammation and infections causes significant pain.

Pediatric dentists are often faced with the management of primary teeth that show signs of irreversible pulpitis or necrosis. Traditionally accepted treatment options for teeth with infected root canals/periradicular tissue are pulpectomy or extraction¹. Although extraction with space maintenance remain a viable treatment option, a successfully restored primary tooth is a superior space maintainer than an appliance²⁻³.

With pulpectomy treatment necrotic or abscessed primary teeth have acceptable outcomes approximately 85 percent of the time⁴, but the technique can be very challenging given the continuous changes in the apical foramen as a result of physiologic and pathologic resorption. Over instrumentation may injure the developing permanent tooth bud⁴⁻⁶. Sjogren et al. found that, even after thorough mechanical and chemical cleaning of the canals, up to 40 percent of the canals may still exhibit positive bacterial cultures⁷.

These findings led researchers to investigate antibiotic option for disinfecting root canal systems. As a result, the cariology research unit of the school of dentistry, Nigata University, Japan developed the concept of lesion sterilization and tissue repair (LSTR) therapy, which is a noninstrumentation endodontic treatment that employs a mixture of antibacterial drugs in a propylene glycol vehicle for the disinfection of dentinal, pulpal and periapical lesions. If lesions are completely disinfected tissue repair can be expected⁸.

Further affected dentine that is clinically softened upon probing could be recalcified after disinfection⁹ and so, affected dentin could be intentionally left in a carious lesion. In addition, when 3-Mix (ciprofloxacin, metronidazole and minocycline) was mixed with macrogol (M) and propylene glycol (P) the combination has been demonstrated to penetrate efficiently dentinal lesions, via dentinal tubules,¹⁰ suggesting that bacteria in pulpal lesions could be killed by placement of 3-Mix MP at the bottom of a carious lesion. Thus one important prediction of the LSTR hypothesis is that local delivery of effective antibiotics in an

appropriate vehicle to a carious lesion may lead to healing of cases of pulpal lesion without a need for conventional pulpectomy procedure. The aim of this study was to evaluate the prognosis of pain and tenderness observed in endodontically involved like reversible pulpitis (vital inflamed pulp) and non-vital necrotic pulp with or without periapical lesion in deciduous teeth using (LSTR) 3-Mix MP therapy.

Methodology

Materials and Methods:

A total of 60 children aged between 4-8 years old who were having 60 infected or inflamed primary molars teeth were selected from the out patient department of conservative dentistry and Endodontics in Bangabandhu Sheikh Mujib Medical University and Pedodontics Department of Dhaka Shishu (Children) Hospital. A general examination of the children was done prior to beginning of the study, the children who were free of any contraindication of endodontic treatment were included in the study. An informed written consent was taken from patient's parents prior to start the study.

The teeth were selected by taking history, clinical and radiographic examination. Mild to moderate pain was assessed by VAS (visual analogue system) and tenderness to percussion was performed by gentle tapping with blunt handle of mouth mirror on all surface of adjacent teeth first then to the offending tooth. Tenderness was assessed as mild, moderate. Vitality and sensitivity test were done by hot and cold stimuli and electric pulp tester.

Teeth with perforated pulpal floor, radiographical evidence of excessive root resorption, deciduous molar which are near to normal shedding, nonrestorable teeth were excluded from the study.

The entire number of respondents was divided into two groups according to diagnosis of the patients.

Group-A: Vital inflamed pulp(initial pulpitis or reversible pulpitis)

Group-B: Non-vital necrotic pulp(with or without periapical lesion).

Preparation of 3-Mix Mp therapy

Commercially available three antibacterial drugs as ciprofloxacin, metronidazole and Minocycline were used in this study. After removal of the enteric coating and the capsules that enclose the drug products, each of the drugs were pulverized and kept separately- in tightly capped porcelain containers to prevent exposure to the light and moisture. These drugs were

stored in refrigerator. Care was taken not to open the containers before they had reached room temperature when used. Before start the treatment the powdered drugs were taken in the ratio of 1:1:1 ciprofloxacin, metronidazol, minocycline and were mixed thoroughly which called. 3-Mix and was kept. One part of macrogol (M) and one part of propylene glycol (P) was taken and then mixed well to make up MP.

One part of MP against 7 parts of 3-Mix (powder) is needed to make 3 mix-MP. These 3 Mix MP were mixed properly and final preparation was small ball like structure of 1 mm of diameter (Fig-2).

Clinical procedure of LSTR

Clinical symptoms such as pain, tenderness to percussion were recorded prior to treatment start. Access cavity was performed, carious lesions and any restoration if present was removed with bur but in group-A care was taken not to cleaned pulpal floor with bur to avoid pulp exposure. Pulpal floor was cleaned with excavator if any soften dentin present. The cavity was prepared such a way to create a medication cavity. [Fig-3]. The cavity was cleaned with sodium hypochloride and normal saline and dried, then 3-Mix MP therapy was placed at the orifice of root canal in group-B and on the bottom of the pulp chamber in group-A and sealed by glassionomer (Fuji-IX) and finally reinforced by composite resin [Fig-4]. The whole procedure was completed in one visit.



3Antibiotics, propylene glycol & macrogol



3 MIX MP

Figure 2



After Cavity preparation



Placement of 3 MIX MP

Figure 3



After treatment



After 1 month follow-up

Figure 4



Patients picture

Pre operative picture of affected tooth

Figure 1

Evaluation of pain and tenderness to percussion

The tooth was considered success depending upon clinical evaluation. Each patient was examined intra-operatively and post operatively at one week-1st follow-up, one month- 2nd follow-up, 6 months 3rd follow-up, and one year final follow-up according to following criteria.

Clinical evaluation

Variables	Success	Failure
Pain	Absence of pain	No remission or increase of pain compare to preoperative status
Tenderness to percussion	Absence of tenderness to percussion	No remission or increase of tenderness to percussion compare to preoperative status

Result:

The observations were based on clinical evaluation, the data were tabulated and subjected to statistical analysis by chi-square test or two sided Fisher's exact test. The results were summarized as follows. Preoperative, post operative clinical findings were recorded.

Table - I

Distribution of samples by affected teeth (n=60)

Position of the tooth	Group A	Group B
Mandibular first deciduous molar	2 (6.6)	8 (26.6)
Mandibular second deciduous molar	26 (86.6)	18 (60.0)
Maxillary first deciduous molar	2 (6.6)	0 (0.0)
Maxillary second deciduous molar	0 (0.0)	4 (13.3)
Total	30 (100.0)	30 (100.0)

Figure within parentheses indicates in percentage. Group A= Vital tooth (inflamed), Group B= Non vital tooth, N=Number of patients

To know about the infected tooth of the patient's it was found that highest affected teeth were Mandibular Second deciduous molar 26(86.6%) the next Mandibular first deciduous molar 2(6.6), Maxillary first deciduous molar 2 (6.6) and Maxillary second deciduous molar 0(0.0) respectively in group-A. In group-B highest affected teeth were mandibular second deciduous molar 18(60%), the next mandibular 1st deciduous molar 8(26.6%) and maxillary second deciduous molar 4(13.3%) respectively.

Table - II
Post-operative pain evaluation (n=60)

Groups			
Pain (VAS)	Group-A (n=30)	Group (n=30)	P value*
Pre operative			
• Mild	18 (60%)	16 (53.3%)	0.999 ^{NS}
• Moderate	12 (40%)	14 (46.7%)	
1st follow-up			
• Absent	26(86.70%)	20 (66.7)	0.235 ^{NS}
• Mild	2 (6.7%)	10 (33.3%)	
• Moderate	0 (0%)	0 (0%)	
• Severe	2 (6.7%)	0 (0%)	
2 nd follow-up			
• Absent	30(100%)	30 (100%)	Note done
3 rd follow-up			
• Absent	30 (100%)	30 (100%)	Note done
• Mild	0 (0)	0 (0)	
Final follow-up			
• Absent	30 (100%)	30 (100%)	Note done

* Fisher's exact test was done to measure the level of significance, Figure within parentheses indicates in percentage, Group A= Vital tooth (inflamed), Group B= Non vital tooth, N= Number of patients, NS= Non significant

Table-2:- Shows the post-operative pain evaluation. In group A all patients were presented pre-operative pain (VAS) mild 18(60%), moderate 12(40%) and in group-B mild 16(53.3%), moderate 14(46.7%) cases, 1st follow-up in group-A, absent, mild, moderate, severe 26(86.7%), 2(6.7%),0(0%), 2(6.7%) and in group-B absent, mild, moderate, severe, 20(66.7%),10(33.3%), 0(0%), 2nd follow-up in group-A absent 30(100%), and in group -B absent 30(100%), 3rd follow-up in group-A absent 30(100%) and in group -B absent 30(100%) and Final follow-up in group-A absent, 30(100%) and in group -B absent, 30 (100%) cases respectively. This is statistically non significant.

Table - III
Post-operative Tenderness to percussion (n=60)

Groups			
Tenderness to percussion	Group-A (n=30)	Group (n=30)	P value*
Pre operative			
• Mild	20 (66%)	16 (53.3%)	0.710 ^{NS}
• Moderate	0 (0%)	14 (46.7%)	
1st follow-up			
• Absent	24(80%)	16(53.3)	0.148 ^{NS}
• Mild	4(13.3%)	14(46.7%)	
• Moderate	2(6.7%)	0(0%)	
2nd follow-up			
• Absent	30(100%)	30(100%)	Not done
3rd follow-up			
• Absent	30(100%)	30(100%)	999 ^{NS}
• Mild	0(0)	0(0%)	
Final follow-up			
• Absent	30(100%)	30(100%)	Note done

*Fisher's exact test was done to measure the level of significance, Figure within parentheses indicates in percentage, Group A= Vital tooth, Group B= Non vital tooth, n= Number of patients, NS= Non significant.

Table-3: Shows the post-operative tenderness to percussion. In group-A 20 patients were presented tenderness to percussion mild, 20(66.7%), and in group- B mild, moderate 16(53.3%), 14(46.7%) cases. In 1st follow-up in group- A absent, mild, moderate 24(80.0%), 4(13.3%),2(6.7%) and in group- B absent, mild, moderate 16 (53.3%), 14(46.7%), 0(0%), 2nd follow- up in group- A absent 30(100%) and in group- B absent 30(100%), 3rd follow- up in group- A absent, mild 30(100%), 0(0%), and in group- B absent, mild 28(93%), 2(6.7%) and final follow- up in group- A absent 30(100%) and in group- B absent 30(100%) cases respectively. This is statistically non significant.

Discussion

Conventional endodontic treatment (pulpectomy) in deciduous molar teeth is very difficult and some times is impracticable due to difficulty in obtaining adequate access to root canals in small mouth , ribbon shaped canal, curve root, presence of permanent tooth bud and physiologic root resorption. Endodontic treatment at the stage of physiologic root resorption is contraindicated by some authorities.¹¹⁻¹⁴

Therefore it can be considered that elimination of bacteria from the root canal system by local application of antimicrobial agents as the LSTR 3mix-MP

therapy might be an effective and alternative method in endodontic treatment of deciduous teeth. This treatment is simple, no need of mechanical preparation, and does not need long chair time or multiple visit. This treatment is also called as non instrumentation endodontic treatment (NIET).

In the present study 60 patients of their 60 deciduous teeth were successfully treated by LSTR 3Mix-MP therapy. All of the patients were suffering from pain. For further assessment the patient's teeth were divided into two groups: vital inflamed pulp (Initial pulpitis or reversible pulpitis) and non-vital necrosed pulp (with or without periapical lesion). In both groups teeth were treated in same method by using LSTR 3Mix-MP therapy. Finally clinical outcome was evaluated at one week (1st follow-up), one month (2nd follow-up), six months (3rd follow-up) and one year (final follow-up) interval.

In group-A all the cases exhibited pain and 20 cases exhibit mild tenderness to percussion. Carious lesion extends to the pulp. The teeth were response to cold stimuli and electric pulp tester. In group-B all cases presented with pain and tenderness to percussion with large carious lesion and pulpless cavity in which no teeth response to hot or cold stimuli and electric pulp tester. The present study showed that in all cases clinical symptoms such as pain and tenderness to percussion were disappeared in both groups finally. Although four cases in vital group were retreated in the same method at the first follow-up period in which pain and tenderness were increased which may be due to accumulation of inflammatory exudation. After retreatment pain and tenderness gradually disappeared and finally no complications were observed and were considered as success.

In group-A (vital group 22 cases pulp remained vital which were response to stimuli at the end of study. The result is corresponded to the previous study Toyohiko Takushige et al.¹⁵. So we can call this pulpitis treatment as LSTR 3Mix-MP save pulp therapy, because most inflamed pulps were saved to be alive instead of the removal of pulp tissue in the conventional endodontic therapy.

The result of this prospective clinical study indicate that LSTR therapy using 3 Mix MP mixture of metronidazole, ciprofloxacin and minocyclin (3 Mix) and macrogol and propylene glycol (MP) provided excellent clinical outcomes in prognosis of pain and tenderness in endodontic treatment of deciduous teeth.

Conclusion:

The following conclusion was drawn within the limitation of this study. Prognosis of pain and tenderness to percussion in endodontic treatment of deciduous teeth using LSTR 3-Mix MP therapy has shown good clinical success and better in non-vital necrosed pulp when compared with vital inflamed pulp.

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