Effectiveness of the EndoActivator System in Removing the Smear Layer after Root Canal Instrumentation

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Abstract

Introduction: Elimination of the smear layer after root canal instrumentation requires the use of irrigating solutions. This cleaning can be completed with passive ultrasonic or sonic irrigation. The aim of this study was to evaluate the effectiveness of the EndoActivator System in removing the smear layer after rotary root canal instrumentation, with and without a final flush of 17% ethylenediaminetetraacetic acid (EDTA) solution, in coronal, middle, and apical thirds. Methods: Forty single-canal teeth were decoronated and randomly divided into 4 groups (n = 10). The groups were instrumented by using Mtwo System. EndoActivator was used with a final rinse of 1 mL of 17% EDTA or 4% NaOCl for 1 minute. The roots were longitudinally split and were grooved in the coronal, middle, and apical thirds. Scanning electron microscopy digital photomicrographs at 400x were taken to evaluate the amount of smear layer in each third. Results: The NaOCl/EndoActivator group did not remove any smear layer of the root canal wall (100% in the coronal, middle, and apical thirds). In the groups that used 17% EDTA (with or without EndoActivator), the smear layer was eliminated completely in the coronal third, but the amount of removal was less in the other two thirds. The comparisons between NaOCl versus NaOCl/EndoActivator groups and EDTA/NaOCl versus EDTA/EndoActivator/NaOCl groups showed no significant differences in root canal thirds. Conclusions: The EndoActivator System did not enhance the removal of smear layer as compared with conventional Max-I-Probe irrigation with NaOCl and EDTA. (J Endod 2009;35:1–4)

Key Words
EDTA, EndoActivator System, scanning electron microscopy, smear layer, sodium hypochlorite

Materials and Methods

Specimen Preparation

Forty single-canal maxillary human teeth extracted for periodontal reasons were stored in a 2% thymol solution until use. The specimens were decoronated to obtain a standardized root length of 15 mm by using a diamond disk in an Accutom 50 cutting machine. The roots were longitudinally split and grooved in the coronal, middle, and apical thirds. Each root was divided into 4 groups (n = 10). The groups were instrumented by using Mtwo System. EndoActivator was used with a final rinse of 1 mL of 17% EDTA or 4% NaOCl for 1 minute. The roots were longitudinally split and were grooved in the coronal, middle, and apical thirds. Scanning electron microscopy digital photomicrographs at 400x were taken to evaluate the amount of smear layer in each third. Results: The NaOCl/EndoActivator group did not remove any smear layer of the root canal wall (100% in the coronal, middle, and apical thirds). In the groups that used 17% EDTA (with or without EndoActivator), the smear layer was eliminated completely in the coronal third, but the amount of removal was less in the other two thirds. The comparisons between NaOCl versus NaOCl/EndoActivator groups and EDTA/NaOCl versus EDTA/EndoActivator/NaOCl groups showed no significant differences in root canal thirds. Conclusions: The EndoActivator System did not enhance the removal of smear layer as compared with conventional Max-I-Probe irrigation with NaOCl and EDTA. (J Endod 2009;35:1–4)

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machine (Struers, Ballerup, Denmark). The external surface of roots was sealed with nail polish to prevent the extrusion of irrigants through the apical foramen (23, 24).

Root Canal Instrumentation

The working length was determined by subtracting 1 mm from the length at which a #10 K-file (Dentsply/Maillefer, Ballaigues, Switzerland) tip extruded apically. Mtwo instruments (Sweden & Martina, Padova, Italy) were used to the full length in permanent rotation with a gentle in-and-out motion, with a 4:1 reduction handpiece (WD-66 EM; W & H, Buermoos, Austria) powered by a torque limited electric motor (Endo IT motor; VDW, Munich, Germany), according to the manufacturer’s instructions. The instrumentation sequence was as follows: #10/04, #15/05, #20/06, #25/06, #30/05, #35/04, and #40/04.

The root canals were flushed with 1 mL of 4% NaOCl solution between files by using a plastic syringe with a closed-end needle (Hawe Max-I-probe; Dentsply Rinn) inserted as deep as possible into the root canal without binding. The teeth were randomly divided into 4 groups (n = 10). In NaOCl group, the canals were irrigated with a final flush of 3 mL of 4% NaOCl. In EDTA/NaOCl group, the canals were irrigated with a final flush of 3 mL of 17% EDTA, after which the EndoActivator System was used to activate the 4% NaOCl for 60 seconds at 10,000 cycles per minute with a #15/02 polymer tip. In EDTA/NaOCl group, the canals were irrigated with a final flush of 1 mL of 17% EDTA for 60 seconds, followed by a final rinse of 3 mL of 4% NaOCl. In NaOCl/EndoActivator group, the canals were irrigated with a final flush of 3 mL of 4% NaOCl. The EndoActivator System was used to activate the 4% NaOCl for 60 seconds at 10,000 cycles per minute with a #15/02 polymer tip. In NaOCl/EndoActivator group, the canals were irrigated with a final flush of 1 mL of 4% NaOCl. The EndoActivator System was used to activate the 4% NaOCl for 60 seconds at 10,000 cycles per minute with a #15/02 polymer tip. Finally, a rinse of 3 mL of 4% NaOCl was applied.

Scanning Electron Microscopy Evaluation

The roots were longitudinally split in the buccolingual plane and were grooved to 3 levels at 4, 8, and 12 mm from the root apices by using a diamond bur to define the coronal, middle, and apical thirds. Canal halves were secured on metal stubs, desiccated, sputter-coated with gold, and viewed with scanning electron microscopy (LEO 1430 VP; Carl Zeiss NTS GmbH, Oberkochen, Germany).

Digital photomicrographs at 400× for smear layer evaluation were taken at the center of each third. The scoring procedure was carried out by 2 independent examiners who were trained in the punctuation procedure, resulting in a sufficient interobserver reproducibility, by using the criteria reported by Torabinejad et al (25), who measured the presence of smear layer as follows: Score 0 = no smear layer; absence of smear layer on the surface of the root canal; all tubules clean and open. Score 1 = moderate smear layer; no smear layer on the surface of the root canal, but tubules contain debris. Score 2 = heavy smear layer; smear layer covers the root canal surface and the tubules.

The final result of the smear layer analysis of each root canal specimen was obtained by calculating the percentage of each score on the images.

Statistical Analysis

The significant differences in the amount of smear layer removal by the irrigating solutions were analyzed by using the χ² Pearson test. The level of statistical significance was set at P < .05. All statistical analyses were performed by means of SPSS 15.0 software (SPSS Inc, Chicago, IL).

Results

The results of the 3 categories of smear layer amount in each of the 4 groups of irrigating solutions or combination of irrigating solutions and sonic system (EndoActivator) appear in Table 1 in the form of percentage distribution. Fig. 1 shows examples of smear layer removal in the coronal, middle, and apical thirds.

In both EDTA groups, the entire dentin wall of the coronal third was free of smear layer (100%), yet this percentage diminished to 80% of the dentin wall surface in the middle third and 20% in the apical third in EDTA/NaOCl group. In EDTA/EndoActivator/NaOCl group, the percentages of dentin wall free of smear layer were 50% and 10%, respectively, for the middle and apical thirds.

When comparing the amount of smear layer removal by all 4 groups in each root third, there were statistically significant differences in the results of the coronal third (P < .05), middle third (P < .05), and apical third (P < .05). Only NaOCl group versus NaOCl/EndoActivator group and EDTA/NaOCl group versus EDTA/EndoActivator/NaOCl group
comparisons were not significantly different in the apical third, middle, or coronal third.

When comparing the results among the root thirds, NaOCl group and NaOCl/EndoActivator group showed no statistically significant differences, because the smear layer amount was equal in the three thirds (the entire surface was covered by smear layer). The other groups, EDTA/NaOCl and EDTA/EndoActivator/NaOCl, exhibited significant differences among thirds \( (P < .05) \). In the pairwise comparisons, EDTA/NaOCl group showed statistically significant differences in the comparisons of coronal third versus apical third \( (P < .05) \) and middle third versus apical third \( (P < .05) \). In EDTA/EndoActivator/NaOCl group, all 3 comparisons among thirds were significant (coronal versus middle, \( P < .05 \); coronal versus apical, \( P < .05 \); and middle versus apical, \( P < .05 \)).

**Discussion**

In the present study focused on evaluating the effectiveness of a new sonically-driven system (EndoActivator) in removing the smear layer after rotary instrumentation of root canals, scanning electron microscopy was used.

Removal of the smear layer during or after root canal instrumentation \( (4, 26) \) requires the use of irrigants that can dissolve both organic and inorganic components. The method recommended for this purpose involves a final rinse with 15% or 17% EDTA solutions followed by 1%–6% of NaOCl. However, there is no consensus with respect to the optimal volume \( (6, 27) \), time of application \( (5, 7) \), or the activation method of irrigating solutions \( (16, 28–31) \).

Ultrasound devices have been used in the removal of smear layer in several studies, although their results are scarcely conclusive \( (9) \). Some authors have shown that the use of PUI for 3–5 minutes with NaOCl concentrations of 3% or 5% \( (15, 18) \) is sufficient for the complete removal of the smear layer in instrumented root canals. However, a time of application less than 1 minute did not allow for complete removal of the smear layer with 1% NaOCl \( (32) \). In this study, the use of a 4% NaOCl solution activated with EndoActivator for 1 minute did not achieve better elimination of smear layer than 4% NaOCl delivered by syringe irrigation. This result could be due to the fact that sonic frequency ranges are much lower than ultrasonic irrigation, and therefore the acoustic microstreaming would be lower, as would be the cleaning efficacy \( (16, 17) \).

The results of this study showed that the most effective smear layer elimination might be related to the use of a final flush with 17% EDTA solution. Without a final flush with 17% EDTA solution, the smear layer was seen to cover the root canal surface in the coronal, middle, and apical thirds, even when the EndoActivator System was used. These
results are in agreement with previous studies underlining the necessity of chelating or acid solutions to remove smear layer in root canal preparation (4, 29, 33, 34). Recently, a study showed the efficacy of a final rinse with 1 mL of 17% EDTA for 1 minute followed by a 3 mL rinse with 6% NaOCl for removing the smear layer after root canal instrumentation (7). Time exposures over 1 minute with EDTA might, however, produce an excess of chelating effect, which could affect the adhesion of endodontic sealer or adhesive cements and decrease the microhardness of the dentin walls (8, 35).

In the present study, a final rinse with 1 mL of 17% EDTA for 1 minute, with or without the use of EndoActivator, eliminated a greater amount of the smear layer in coronal and middle thirds than in the apical third, which showed the worst results. Meanwhile, de Gregorio et al (21), by using ultrasonic and sonic (EndoActivator) activation in simulated lateral canals, found better irrigation in the apical third (at 4.5 and 2 mm from working length) than with traditional needle irrigation alone, and the addition of EDTA did not result in better penetration of irrigants into the lateral canals. However, Mancini et al (34), in agreement with our results, showed that the smear layer was not removed completely in the apical third of the root canal by using 1 mL of BioPure MTAD (Dentsply Tulsa Dental) or 1 mL of 17% EDTA, followed by 3 mL of 5.25 NaOCl. This finding might be attributed to the lesser volume of final rinse used (1 mL 17% EDTA and 3 mL 4% NaOCl) and/or the major circulating volume of irrigant present in coronal and middle thirds as opposed to the apical third. In this sense, it was demonstrated that when the apical foramen was sealed, the apical thirds failed to become clear of smear layer (23, 24). A recent study indicated that a greater volume of 17% EDTA (5 mL) for 1 minute with ultrasonics followed by a greater volume of 1% NaOCl (5 mL) would prove efficient for smear and debris removal at the apical region of the instrumented root canal (30).

Chopra et al (31) irrigated root canals with a final flush of 10 mL 17% EDTA activated with a similar device, the F-file (PlasticEndo, Buffalo Grove, IL), for 30 seconds at 600 rpm in the electric slow-speed rotary handpiece (Dentsply Tulsa Dental Specialties) followed by 10 mL 6% NaOCl; they concluded that the F-file was no more beneficial in removing smear layer, and that smear layer removal appeared to be mostly influenced by the introduction of an EDTA rinse. Within the limitations of our study, the EndoActivator System did not enhance the removal of smear layer as compared with conventional Max-I-Probe irrigation with NaOCl and EDTA. A final irrigation with 1 mL of 17% EDTA solution was necessary to remove the smear layer after rotary instrumentation of root canal with or without the use of the EndoActivator System. The removal of smear layer was more complete in coronal and middle thirds than in the apical third. Further research entailing different solutions, volumes, and activation times of the irrigant would be necessary to fully evaluate the effectiveness of the EndoActivator System in smear layer removal after root canal instrumentation.  

References