

Single Versus Multi-visit Endodontic Treatment of Teeth with Apical Periodontitis: An *In vivo* Study with 1-year Evaluation

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Abstract

Background: Apical periodontitis (AP) is one of the most prevalent diseases of the teeth. Treatment of AP is based on the removal of the cause, i.e., bacteria from the root canals. Achievement of adequate bacterial eradication in one appointment treatment remains a controversy. **Aim:** This prospective study was conducted with the objective to compare the periapical healing of teeth with AP treated in (a) single visit versus (b) two visits, either with or without Vitapex as an intracanal medicament. **Subjects and Methods:** Patients were selected randomly from the Department of Conservative Dentistry and Endodontics. Forty-three patients (81 teeth) met the inclusion criteria, i.e., AP (both symptomatic and asymptomatic) visible radiographically size $\geq 2 \text{ mm} \times 2 \text{ mm}$, not suffering from any immune-compromising disease, age between 16 and 65 years and tooth not accessed previously. Patients were randomly divided into three groups, i.e., single-visit group (Group 1), multi-visit group without any intracanal medicament (Group 2), and multi-visit group with Vitapex as interim intracanal medicament (Group 3). Comparison was done radiographically using periapical index (PAI). The primary outcome measure was the change in periapical radiolucency after 1 year assessed by PAI scores. The Mann–Whitney U-test was used to evaluate differences between groups at baseline (immediate postoperative) and at the 12-month follow-up evaluation. Change in PAI score for each group from baseline to 12-month follow-up evaluation was tested with the Wilcoxon signed rank test. The secondary outcome measures, proportion of teeth in each group that could be considered improved (decreased PAI score) or healed (PAI < 2), were assessed with the Chi-square test. **Results:** No statistically significant difference in periapical healing was found between three groups. **Conclusion:** After 1-year evaluation, no difference in periapical healing was found between single-visit treatment and multi-visit treatment groups with the given sample size.

Keywords: 1-year evaluation, Apical periodontitis, Multi-visit treatment, Single visit, Vitapex

Introduction

Apical periodontitis (AP) is one of the most prevalent diseases of the teeth with prevalence as high as one radiolucency per patient noted in a recent study.^[1] It is the main prognostic factor in initial endodontic treatment, healed rate in teeth with preoperative

radiolucency is significantly lower than teeth without AP.^[2] Ørstavik and Pitt Ford have stated that the “ultimate biological aim of root canal treatment as either to prevent or cure AP (in cases where disease is already present).”^[3] Treatment of AP aims at the removal of cause, i.e., bacteria present in the canal.

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Role of bacteria in the causation and the persistence of AP are well-established.^[4,5] Mechanical debridement combined with antibacterial irrigation (0.5–6% sodium hypochlorite) can render 40–60% of the treated teeth bacteria-negative.^[6,7] In addition to mechanical debridement and antibacterial irrigation, use of calcium hydroxide as intracanal medicament between canals has been shown to reduce the bacterial count further by 80–100%.^[8,9] This has been the basis of the multi-visit treatment of AP. However, many studies have questioned the effectiveness of the calcium hydroxide in the healing of AP.^[6,10-12] Moreover, the tooth may also be susceptible to reinfection through the temporary filling and dressing during the interim period in case of multiple visits.

Single-visit root canal treatment of the AP relies on mechanical debridement, disinfection with antimicrobial irrigating solution, and the antimicrobial activity of the sealer or the zinc (Zn²⁺) ions of Gutta-percha for eradication of bacteria.^[12-14] Remaining bacteria may be deprived of nutrition and space to multiply.^[15] It has become common practice and offers several advantages such as high patient acceptance^[16] and reduced flare-up rate.^[17,18]

Achievement of adequate bacterial eradication in one appointment treatment is the main controversial question. Although there might be a reasonable biologic argument to prefer multiple appointment root canal therapy for infected teeth with AP, clinical research has been equivocal.^[19-21] The direct evidence comparing the healing rates following single- and multiple-visit root canal treatment should provide insight as to which regimen is more effective for the treatment of such a prevalent problem.

The main objective of the present study was to compare the periapical healing of teeth with AP treated in (a) single visit and in (b) multiple visits with and without intracanal medicament (Vitapex, J. Morita Co., Tokyo, Japan) using periapical index (PAI).

Subjects and Methods

Patient selection

After informed consent, study subjects were selected randomly from among the patients referred to the Department of Conservative Dentistry and Endodontics for initial nonsurgical endodontic treatment. The study was conducted between 2010 and 2013 in Conservative Dentistry and Endodontics Department. Ethical clearance for the study was obtained from the Institutional Review Board.

The primary criterion for inclusion of subjects in the study was the presence of radiographically demonstrable AP (minimum size - 2.0 mm × 2.0 mm) in maxillary anterior or mandibular anterior tooth/teeth. Both symptomatic and asymptomatic patients were included in the study. Patients were excluded from the study if (a) they had a diagnosis of diabetes, (b) they

had a diagnosis of immune-compromising disease, (c) they were <16 or >65-year-old, or (d) the tooth had been previously accessed or treated. Once the eligibility was confirmed, the study was explained to the patient, and the patient was invited to participate. After written and verbal informed consent was obtained, the patients were randomly assigned to the treatment groups by throwing of a dice viz., Group 1 (single-visit treatment group), Group 2 (multi-visit treatment group without any intracanal medicament), and Group 3 (multi-visit treatment group with Vitapex as an intracanal medicament).

Clinical procedure

All patients were treated by one investigator (gsg) according to a standard regimen including elements of access, rubber dam, and establishment of asepsis. Local anesthesia (2% lidocaine with 1:100,000 epinephrine) was administered to the patient. Initial caries excavation was performed, rubber dam isolation was obtained, and a standard access cavity was prepared. Initial canal working length was established by using the Root ZX electronic apex locator (J. Morita Co., Tokyo, Japan) and a #15 stainless steel K-file. Working length was confirmed and adjusted as needed by using radiographs. Canals were instrumented using stainless steel K-files and H-files using step-back technique. Canals were irrigated with 5.0 mL 2.5% sodium hypochlorite after each instrumentation cycle, and canal patency was maintained by passing a #10 stainless steel file up to working length. After the completion of canal instrumentation, all canals were irrigated with 5.0 mL, 17% ethylenediaminetetraacetic acid (EDTA) (Glyde, Dentsply Maillefer, Ballaigues, Switzerland) for 1 min followed by a final irrigation with 5.0 mL 2.5% sodium hypochlorite. Canals were then dried with sterile paper points. Obturation was done with Gutta-percha and zinc oxide eugenol sealer using lateral condensation technique.

Teeth in Group 1 were obturated at the same appointment by using the lateral condensation technique using Gutta-percha and access cavities were restored using anterior composite restorations.

In Group 2, instrumentation was completed at the first appointment. The canal was left empty, and access cavity was closed using Cavit-G (3M ESPE) for at least 1 week before the second appointment. On the second appointment, the obturation was completed using lateral condensation technique and tooth/teeth restored with composite restoration.

In Group 3, instrumentation was completed at the first appointment. Vitapex was placed in the canal to remain for at least 1 week before second appointment. At the second appointment, the Vitapex paste was removed by using circumferential filing with Hedstrom-files and copious irrigation with 2.5% sodium hypochlorite followed by 5.0 mL 17% EDTA and a final rinse of 5.0 mL 2.5% sodium hypochlorite. Complete removal of the Vitapex paste was confirmed radiographically. The canals were dried with

sterile paper points, and obturation was performed with the same technique described for Group 1. After completion of treatment, the teeth were restored with a composite restoration.

Radiographic technique

All radiographs were done using Kodak RVG5000 size#2 sensor and Gnatus X-ray machine operating at (70 Kvp-7 mA-0.08-1540VA) with exposure settings kept constant at 0.32 s. Immediately after obturation, digital radiographs were taken using Rinn XCP sensor holder (Rinn Corp., Elgin, IL, USA) stabilized with polyvinyl siloxane impression material (AFFINIS-Coltene Whaledent) to standardize image geometry. Impression of each patient was preserved for follow-up radiograph in a polythene packet. Follow-up radiographs after 1 year recall were done with the individual custom index and recorded exposure settings [Figure 1].

Outcome measures and data analysis

The primary outcome measure for this study was change in apical bone density (radiographically) at 12 months. Secondary outcome measures were the presence of clinical symptoms or abnormal findings at 12 months (spontaneous pain, presence of sinus tract, swelling, mobility, or sensitivity to percussion or palpation).



Figure 1: RVG sensor with Rinn XCP sensor holder with addition silicone impression

Radiographic evaluation [Figure 2] was done using the PAI scoring system.^[22,23] This is a 5-point scale radiographic interpretation designed to determine the absence, presence, or transformation of a disease state. Radiographs to be assessed were compared with a set of five radiographic images with corresponding line drawings and their associated score on a photographic print (derived from Brynolf's histologic-radiographic co-relation study).^[24]

Radiographic images were evaluated blindly and independently by 4 experienced endodontists by assigning a score according to which of the reference images it appears to match. Before evaluation of the study images, each examiner graded a series of twenty radiographic images not associated with the study sample and representing a wide range of periapical bone densities. Instructions for grading images with the PAI scoring system were adapted from Orstavik *et al.*^[22] and are presented in Figure 2. The following specific written instructions were given to the examiners:

- Find the reference radiograph where the periapical area most closely resembles the periapical area you are studying. Assign the corresponding score to the observed root
- When in doubt, assign a higher score
- All teeth must be given a score.

Approximately, 1 week after the calibration session, each examiner independently scored the randomly assembled study images. All study images were coded by the investigator and evaluated randomly and individually on a 15.6" LCD monitor by the four experienced endodontists. The examiners then met as a group and reviewed all scores to enhance calibration and inter-rater agreement. Consensus was reached on images that were not initially scored the same by all examiners.

To assess intrarater agreement, 4 weeks after the first session the examiners again scored all study images. This method generated a total of 6 PAI scores for each image, 2 from each of the three examiners. Agreement between and within examiners was determined by using the intraclass correlation coefficient (ICC). Intrarater reliability was measured with the single measure ICC (SPSS 13 for Windows, SPSS Inc., Chicago, IL, USA), and inter-rater agreement was measured

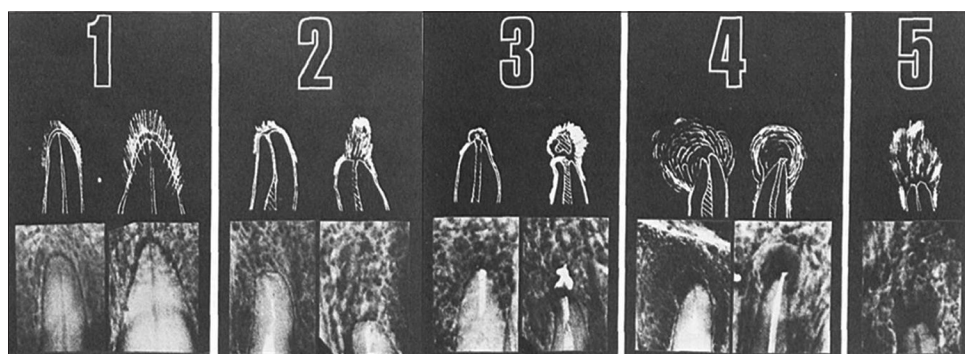


Figure 2: Periapical index (adapted from Orstavik *et al.*^[22])

with the average measure ICC (also known as the inter-rater reliability coefficient).

After the second independent scoring session, the examiners met as a group to reach consensus on cases that did not receive unanimous agreement on the PAI score. Consensus was reached on all images. The consensus score for each image was considered the true score and was used for statistical analysis. The identifying code for each image was not broken until after the consensus score was determined. Computations of necessary sample sizes for comparisons among treatments indicate that sample sizes of 55 per group would be sufficient to detect differences in rates of 5% or more, with a power of $P = 0.20$.^[25]

The Mann–Whitney U-test was used to evaluate differences between groups at baseline (immediate postoperative) and at the 12-month follow-up evaluation. Change in PAI score for each group from baseline to 12-month follow-up evaluation was tested with the Wilcoxon signed rank test. The secondary outcome measures, proportion of teeth in each group that could be considered improved (decreased PAI score) or healed ($PAI < 2$), were assessed with the Chi-square test. Clinical symptoms and abnormal findings at the 12-month follow-up examination were recorded but not subjected to statistical analysis.

Results and Statistical Analysis

The conventional procedure should produce a success rate of approximately 80%.

Eighty-one teeth in forty-three patients met the inclusion criteria and were included in the study after getting consent from the patients. There were two treatment failures (five teeth) before the 1 year period, one each in Group 2 and Group 3. Periapical surgery was done in one patient due to persistent draining sinus, within 3 months of obturation, other patient had an accident and both teeth had to be extracted because of fracture. Ten patients (16 teeth) were lost to follow-up. We defined failure as the need for any additional treatment before or at the 12-month follow-up evaluation period.

In this study, 60 teeth (Group 1 = 21, Group 2 = 18, Group 3 = 21) presented for the 1 year follow-up examination. Because these numbers were unlikely to be large enough for statistical differences between the groups, a power analysis evaluation was performed.

Agreement between and within examiners was determined by using the ICC. Single and average measure inter-reliability score of 0.91 and 0.95, in this study, represent a very high level of agreement between the examiners.^[26]

Mean PAI score for Group 1 was 3.42 at baseline and decreased to 2 by the end of 1 year, a decrease of 1.42. Similarly, the

decrease in PAI for Group 2 and Group 3 was 1.28 and 1.67, respectively. The decrease in PAI was statistically significant in all groups ($P < 0.001$) [Table 1 and Graph 1].

In Group 1 and 3, 76.2% (16/21) teeth could be considered healed ($PAI \leq 2$), whereas in Group 2, only 66% (12/18) teeth could be considered healed. There was no statistically significant difference in healing among the groups [Table 2 and Graph 2].

Clinical symptoms and abnormal findings at the 12-month follow-up examination were recorded but not subjected to statistical analysis. One tooth in Group 2 was discolored after 1 year except that there were no abnormal findings.

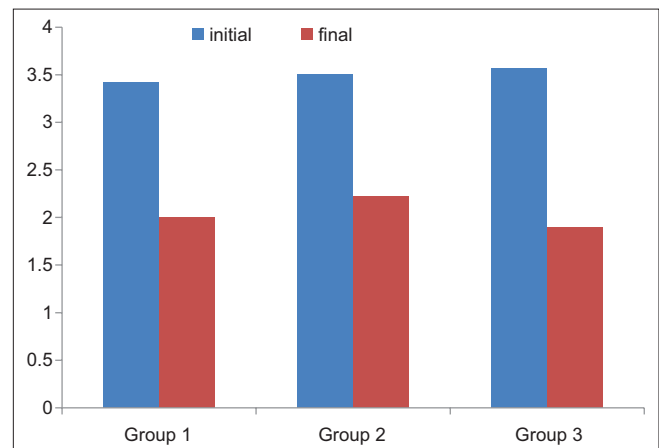
Discussion

AP is one of the most prevalent diseases affecting the teeth. Prevalence increases with age and by the age of 50 years, it is more than 50%.^[27] It is equally well established that prognosis for complete healing of teeth with pretreatment diagnosis of AP is approximately 10–15% lower than for teeth without AP.^[28,29] With billions of teeth affected with the disease, a treatment regimen, which can result in successful and predictable healing can help save many teeth. This prospective study was done to compare the effectiveness of two commonly used nonsurgical primary endodontic treatment regimens for the treatment of AP, i.e., single visit and traditional multi-visit treatment regimen.

Table 1: Periapical index score at immediate postoperative and 12-month evaluation

Groups	Immediate postoperative mean PAI (SD)	1 year recall mean PAI (SD)	Change in PAI score with 95% CI
1	3.42 (0.67)	2 (1.224)	1.42
2	3.5 (0.67)	2.22 (1.35)	1.28
3	3.57 (0.7859)	1.90 (1.09)	1.67

PAI: Periapical index, SD: Standard deviation, CI: Confidence interval

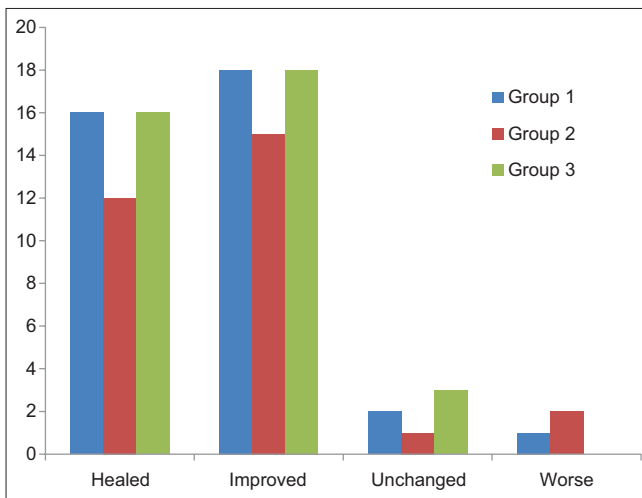


Graph 1: Periapical index score at immediate postoperative and 12-month evaluation

Table 2: Proportion of teeth healed, improved, unchanged, or worse in each group at 12-month evaluation

Status	Group 1	Group 2	Group 3
Healed % (PAI <2)	76.19 (16/21)	66.67 (12/18)	76.19 (16/21)
Improved % (decreased PAI)	85.7 (18/21)	71.4 (15/21)	85.7 (18/21)
Unchanged % (same PAI)	9.52 (2/21)	5.56 (1/18)	14.29 (3/21)
Worse % (increased PAI)	4.76 (1/21)	11.11 (2/18)	0.00 (0/21)

PAI: Periapical index

**Graph 2:** Proportion of teeth healed, improved, unchanged, or worse in each group at 12-month evaluation

Only teeth with definite periapical lesions (AP) were included in the study making the comparison between treatment protocols viable. The expectation that teeth treated in two visits with an interappointment dressing of Vitapex paste would result in improved healing when compared with one-visit root canal therapy was not supported by this study. No statistically significant differences were found between the three treatment groups in the healing of AP; our findings are consistent with the majority of well-controlled clinical studies.^[11,12,30-32] When comparison was done between initial and final (1 year recall) PAI score in each group, statistically significant improvement in PAI score was found in every group (including Group 2), which corroborates the study of Gesi *et al.*^[33] that with proper use of aseptic operating procedures, proper instrumentation, and filling, an inter-appointment dressing with calcium hydroxide does not seem to influence outcome.

Leaving the canal empty without obturation or use of any additional medicament (Group 2) was numerically (though not statistically significant with given sample size) the worst method of treatment, which are consistent with the study of Trope *et al.*^[11]

Main problems with prospective studies are the recall rate, adequate sample size, and control of the baseline status. Sixty

teeth out of the original sample of 81 teeth were included in the final analysis with a recall rate of 74% which is more than the median recall rate of 52.7% Ng *et al.*^[34] and Penesis *et al.*^[31] but less compared to the studies by Weiger *et al.*^[12] and Paredes-Vieyra and Enriquez.^[32]

Although the sample size in this study was small, it is typical when compared with similar studies,^[11,12,30] but the sample size of the present study was not large enough for statistical significance. Because even inferior protocols result in about 60% success, it requires a large sample size to show statistically significant differences in results for different treatment protocols. These numbers are extremely difficult to achieve in a prospective study like this one considering the time constraints, limited resources, and subject dropouts associated with long recall period. To compensate for this problem, a power analysis^[35] of the results was performed to calculate the adequate sample size required to make the results clinically significant. In general, a sample size of 49 per group would have resulted in significant differences between the single visit and multiple visit groups.

To minimize the bias, patients were randomly assigned to treatment groups, and root canal therapy was performed according to a standardized regimen. Conventional hand instrumentation was done as it was planned as a baseline study so that subsequently studies using rotary instruments can be done and influence of different techniques compared with the study.

Only anterior teeth were included in the study similar to Sjögren *et al.*^[6] To control the baseline apical status, only teeth with visible radiolucent area of 2.0 mm × 2.0 mm were included similar to Penesis *et al.*,^[31] which assured an initial PAI score ≥3. The only known independent variable was number of visits and use of Vitapex as intracanal medicament. Fortunately, the number of teeth in each group was reasonably evenly distributed during treatment and after 1 year recall, neither group varied significantly from the study dropouts.

Calcium hydroxide is currently acknowledged as the best intracanal medicament in endodontic treatment procedures. It is effective against most endopathogens but has a limited activity against *Candida albicans* and *Enterococcus faecalis*. Leonardo *et al.*^[36] recommended the addition of other substances to the paste to increase or maintain the paste consistency of the material which does not harden or set in the root canal, to improve flow, maintain pH of calcium hydroxide, to improve radio-opacity, to make clinical use easier and not alter the excellent biological properties of calcium hydroxide itself. Hence, Vitapex was used as an intracanal medicament which contains iodoform (synergistic action), silicone oil (silicone oil based calcium hydroxide is effective in the elimination of *E. faecalis*),^[37] and radiopacifiers, which increase the paste radiopacity, which helps in confirming insertion and complete removal from the root canal using radiograph.

Vitapex was placed in the root canals of Group 3 for a period of 7 days before obturation as in the studies by Law and Messer,^[9] Siqueira *et al.*^[38] for calcium hydroxide dressings.

After root canal treatment the probability of healing increases over time,^[12,39] however periapical changes are evident by 1 year.^[40] Orstavik^[41] concluded that 1-year follow-up predicts long-term success and the risk of disease development at 2 years or later is not greater than general risk based on epidemiological studies. As the number of dropouts increases with the time and recall rate falls, many studies have used 1-year recall period like the present study.^[11,31,42]

Evaluation of results was done using PAI scoring system. PAI scoring system given by Orstavik *et al.*^[22] is based on a histological correlation study of Brynolf^[24] with a PAI score of 1–2 defined as healed or minimally inflamed and scores 3–5 defined as diseased. This index was used by 58 studies from 1986 to 2009 and is accepted as a valid tool to determine outcome and reveal changes in bone density after root canal treatment.^[43]

However, PAI is still a subjective outcome and is prone to observer variation. To overcome this disadvantage as much as possible, calibration of examiners was done and consensus score was taken as a true score. Agreement between and within examiners was determined by using the ICC. Single and average measure inter-reliability score of 0.91 and 0.95, respectively, in this study, represent a very high level of agreement between the examiners, even before meeting for a consensus score for each image similar to that in the study by Penesis *et al.*^[31]

In most studies, PAI scores are generally dichotomized as healed (PAI 1, 2) and diseased (PAI 3, 4, 5), but it may obscure the small amount of change that may be clinically significant. The process of calibrating observers and arriving at a consensus is also very time-consuming.

Measurement of density as a continuous variable can be more expressive than the PAI scoring system^[44,45,46]. Use of digital radiography has added advantages of less radiation exposure, time saving, and freedom from processing errors. However, variations in soft tissue superimposition and physiologic modeling over time may influence the measurements, producing erroneous information.

Nevertheless, the sample and operator may not accurately represent the true population of patients and clinicians. Results can be influenced by many unknown and uncontrolled variables like on an average, every patient contributed two teeth in the study with some patients contributing three or four teeth which increased the dependency of results on subjects. Similarly, diabetic and immune-compromised patients were excluded from the study, but smokers were not despite the evidence that they also experience poorer

treatment outcomes.^[47,48] All cases in the study were done during the period of postgraduation course (2010–2013) under faculty supervision with the opportunity for consultation and assistance as needed; therefore, the treatment environment might not be truly representative of either specialty or general dental practice setting.

Conclusion

From the present study, it can be concluded that 12 months after initial nonsurgical root canal therapy on teeth with AP, there was no statistically significant difference in periapical healing between single visit treatment and multi-visit treatment with the given sample size.

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Conflicts of interest

There are no conflicts of interest.

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