Multiple-file vs. single-file endodontics in dental practice: a study in routine care

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Background. Little is known about the differences of rotary multiple file endodontic therapy and single-file reciprocating endodontic treatment under routine care conditions in dental practice. This multicenter study was performed to compare the outcome of multiple-file (MF) and single-file (SF) systems for primary root canal treatment under conditions of general dental practice regarding reduction of pain with a visual analogue scale (VAS 100), improvement of oral-health-related quality of life (OHRQoL) with the german short version of the oral health impact profile (OHIP-G-14) and the speed of root canal preparation. **Materials and Methods.** Ten general dental practitioners (GDPs) participated in the study as practitioner-investigators (PI). In the first five-month period of the study the GDPs treated patients with MF systems. After that the GDPs treated the patients in the second five-month period with a SF system (WaveOne). The GDPs documented the clinical findings at the beginning and on completion of treatment. The patients documented their pain and OHRQoL before the beginning and before completion of treatment. **Results.** 599 patients were included in the evaluation. 280 patients were in the MF group, 319 were in the SF WaveOne group. In terms of pain reduction and improvement in OHIP-G-14 the improvement in both study groups (MF and SF) was very similar based on univariate analysis methods. Pain reduction was 34.4 (SD 33.7) VAS (MF) vs. 35.0 (SD 35.4) VAS (SF) (p=0.840) and the improvement in OHIP-G-14 score was 9.4 (SD 10.3) (MF) vs. 8.5 (SD 10.2) (SF) (p=0.365). The treatment time per root canal was 238.9 sec (SD 206.2 sec) (MF) vs. 146.8 sec. (SD 452.8 sec) (SF) (p=0.003). **Discussion.** Regarding improvement of endodontic pain and OHRQoL measure with OHIP-G-14 there were no statistical significant differences between the SF und the MF systems. WaveOne prepared root canals significantly faster than MF systems.

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17 Running head: Multiple-file vs. single-file endodontics

18 **Abstract**

19 **Background.** Little is known about the differences of rotary multiple file endodontic therapy and

20 single-file reciprocating endodontic treatment under routine care conditions in dental practice.

21 This multicenter study was performed to compare the outcome of multiple-file (MF) and single-

22 file (SF) systems for primary root canal treatment under conditions of general dental practice

23 regarding reduction of pain with a visual analogue scale (VAS 100), improvement of oral-health-

24 related quality of life (OHRQoL) with the german short version of the oral health impact profile

25 (OHIP-G-14) and the speed of root canal preparation.

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27 practitioner-investigators (PI). In the first five-month period of the study the GDPs treated

28 patients with MF systems. After that the GDPs treated the patients in the second five-month

29 period with a SF system (WaveOne). The GDPs documented the clinical findings at the

30 beginning and on completion of treatment. The patients documented their pain and OHRQoL

31 before the beginning and before completion of treatment.

32 **Results.** 599 patients were included in the evaluation. 280 patients were in the MF group, 319

33 were in the SF WaveOne group. In terms of pain reduction and improvement in OHIP-G-14 the

34 improvement in both study groups (MF and SF) was very similar based on univariate analysis

35 methods. Pain reduction was 34.4 (SD 33.7) VAS (MF) vs. 35.0 (SD 35.4) VAS (SF) (p=0.840)

36 and the improvement in OHIP-G-14 score was 9.4 (SD 10.3) (MF) vs. 8.5 (SD 10.2) (SF)

37 (p=0.365). The treatment time per root canal was 238.9 sec (SD 206.2 sec) (MF) vs. 146.8 sec.

38 (SD 452.8 sec) (SF) (p=0.003).

39   **Discussion.** Regarding improvement of endodontic pain and OHRQoL measure with OHIP-G-14

40   there were no statistical significant differences between the SF und the MF systems. WaveOne

41   prepared root canals significantly faster than MF systems.

43 **Introduction**

44 Clinical endodontic research is mainly conducted by specialists or specialized university centers

45 (Friedman et al. 2010; Ng et al. 2007). The predominant types of such studies are retrospective

46 observational studies, prospective cohort studies and a few randomized controlled trials (RCTs)

47 (Ng et al. 2007). On account of the controlled study design, these studies have greater internal

48 evidence and are classified as efficacy studies (Pfaff et al. 2011). The effectiveness of endodontic

49 interventions under everyday general dental care conditions has so far been hardly investigated

50 (Nixdorf et al. 2012). Yet, patients treated in specialized centers can differ systematically from

51 patients treated in routine care (Hulley 2013).

52 A commonality of many experimental endodontic studies is the low number of cases (Peters &

53 Wesselink 2002; Pettiette et al. 2001; Weiger et al. 2000). Larger case numbers are described for

54 retrospective observational studies and prospective cohort studies, which, however, are often

55 conducted without controls (Ng et al. 2007). Convincing results though can be obtained in

56 studies if they include an adequate number of cases (Hulley 2013). Since only few experimental

57 endodontic studies have been made and many of them are lacking sufficient patient numbers, one

58 could assume this to be an indication of considerable feasibility problems of such studies.

59 Reciprocating single-file (SF) systems are the latest stage of development of nickel-titanium

60 (NiTi) instruments for the preparation of root canals (Bürklein et al. 2013; Yared 2008). During

61 the last years several systems as Reciproc (VDW, Munich, Germany), WaveOne (Dentsply,

62 Konstanz, Germany), Genius files (Ultradent, South Jordan, UT, USA) or the Twisted Files

63 Adaptive System (Kerr, Orange, CA, USA) with a combination of rotary and reciprocating

64 movement were introduced into the market. Our knowledge of the clinical effects of using

65   different systems for root canal preparation is limited (Schäfer et al. 2004). The Swedish Council

66   on Health Technology Assessment stated in its Systematic Review of Methods of Diagnosis and

67   Treatment in Endodontics that the use of new tools facilitates the technical procedures of root

68   canal treatment and that therefore investigations are needed, which influence these techniques

69   have on everyday general practice (Bergenholtz et al. 2012).

70   Typically, new instrument systems are investigated in in-vitro-studies with extracted teeth

71   (Bürklein et al. 2013) or root canal models (Goldberg et al. 2012). In such studies, the outcomes

72   are mainly surrogate parameters, such as root canal straightening, preparation faults, preparation

73   time in a workbench situation etc. the clinical significance of which can only be estimated to a

74   limited extent (Hülsmann 2013). Most of the few clinical trials available investigated only one

75   instrument system (Fleming et al. 2010; Su et al. 2011) and rarely allow a comparison with other

76   instrument systems (Schäfer et al. 2004). Recently some randomized controlled trials were

77   published, that investigated single and multiple file systems for endodontic treatment regarding

78   pain reduction after treatment and improvement in quality of life (Kherlakian et al. 2016;

79   Pasqualini et al. 2015; Relvas et al. 2015). It is unclear if there exists an effectiveness-gap (Pfaff

80   et al. 2011) between the results of these controlled studies under the optimal treatment conditions

81   of specialized treatment providers and the use of rotary multiple-file (MF) and SF systems in

82   general dental practice.

83   Therefore research is needed when new endodontic techniques are introduced into dental

84   practice. The study we performed, investigates the effects of these endodontic techniques on

85   dental practice. For this purpose it uses the methods of health services research which studies

86   care processes under everyday conditions of dental practice (Pfaff et al. 2009). Short-term

87   Patient-relevant outcomes were in the center of the study.

88 The design we chose was a multicenter study in routine care. We started by evaluating the

89 outcome of endodontic treatment using conventional MF instrument systems for root canal

90 preparation. Then the practitioner-investigators (PIs) were trained in single-file (SF) endodontics

91 (WaveOne-Instruments, Dentsply Maillefer, Ballaigues, Switzerland). Subsequently we

92 evaluated the outcome of endodontic treatments using WaveOne.

93 The following research hypotheses were investigated in our study:

94 ***Primary outcome criterion***

95 Does root canal preparation using SF root canal instruments lead to more or less reduction of

96 patients' endodontic pain compared to using rotary MF instrument systems?

97 ***Secondary outcome criterion***

98 Does root canal preparation using SF root canal instruments lead to more or less reduction of

99 patients' oral-health-related quality of life compared to using rotary MF instrument systems?

100 ***Tertiary outcome criterion***

101 Does root canal preparation using a single-file system require less time compared to the MF

102 systems?

103 **Methods**

104 **Study Design**

105 We performed the present study as a multicenter clinical study. For the purpose of this study we

106 formed a network of ten general dental practitioners (GDPs). They acted as PIs. We conducted

107 the study in two phases (Figure 1). In the first 5-month phase the GDPs performed the

108   endodontic therapy with different rotary nickel-titanium (NiTi) MF systems (Table 1).

109   Subsequently the GDPs were trained for the use of the WaveOne SF system (Maillefer,

110   Ballaigues, Switzerland). In the second 5-month phase the PIs treated the patients with SF

111   WaveOne instruments exclusively. After each 5-month phase there was a 3-month follow-up so

112   that treatments started could be completed.

113   The authors of this study acted solely as investigator and did not treat patients.

114   The study was conducted in conformity with the Declaration of Helsinki and the Professional

115   Code for Physicians of the Medical Council of the State of Baden-Württemberg. The Ethics

116   Committee of the Baden-Württemberg Medical Council reviewed the study and approved it (AZ:

117   F-2011-034-z).

118   **Participants**

119   *Patient eligibility and recruitment*

120   All patients of the ten PIs who required endodontic therapy were consecutively assessed for

121   eligibility.

122   The following inclusion criteria were defined: patients had to be at least 18 years old and in need

123   of initial orthograde root canal treatment.

124   The following exclusion criteria were defined: Patients with hopeless teeth for periodontal or

125   restorative reasons, patients treated for emergency reasons only, more than one symptomatic

126   tooth requiring endodontic treatment at the same time in one patient, patients with other oral

127   findings causing pain, patients with craniomandibular dysfunction and communication

128   difficulties (eg, patients were not able to read, understand and complete the study questionnaires

129   in German language).

130   All patients were recorded by the assistant staff of the dental practice and asked for the reason if

131   they refused to participate. Every patient was given the study education and information

132   documents (informed consent) that had to be signed and submitted by the patient before the

133   patient was included in the study.

134   *Practitioner Investigators (PIs)*

135   The ten dentists who participated in the study were general dental practitioners with at least two

136   years of professional experience in a general dental practice and without endodontic

137   specialization. All participating dentists worked under the conditions of the German "Statutory

138   Health Insurance". The PIs were chosen as a convenient sample of dentists that wanted to change

139   their endodontic treatment to single file systems within the next 6-12 months anyway. All

140   practices were located in southwest Germany.

141   All PIs were familiar with root canal preparation using rotary NiTi instruments (Tab. 1) and used

142   them routinely in their practice. All dentists followed the "Good Clinical Practice: Root Canal

143   Treatment" Guideline of DGZMK (German Society of Dental, Oral and Craniomandibular

144   Sciences) (Hülsmann & Schäfer 2005) which contains essential key points of the Quality

145   guidelines for endodontic treatment of the European Society of Endodontology (Endodontology

146   2006).

147   *Study initiation at the PIs*

148   Before the study started, all participating PIs were visited by the principal investigator (AB) in

149   their practice. The dentists were informed about the object and purpose of the study and its

150   practical implementation. Each PI was given a copy of the study protocol and all other study

151   relevant files. The dentists were informed about the planned procedure with regard to patient

152   recruitment, education/information and treatment.

153   **Interventions**

154   In the first phase of the study from 09/2011 to 02/2012 all endodontic treatments were performed

155   with rotary NiTi MF systems (Figure 1). All MF systems were used according to the

156   manufacturer's instructions. In 03/2012 the PIs were trained for the SF system. The training

157   course explained the theoretical bases of the WaveOne System (Maillefer, Ballaigues,

158   Switzerland) and provided hands-on training on extracted teeth. After this one-day training

159   course every participating dentist was able to prepare root canals by the new method in a reliable

160   way. The training was followed by a two-week implementation phase in all participating dental

161   offices. During that time the PIs should learn to treat patients with the new instruments and gain

162   experience. In case of difficulties this procedure offered the chance of clarifying problems. In the

163   second phase of the trial from 04/2012 to 08/2012 all endodontic treatments were performed with

164   the SF system. All other variables of the practice setting and the treatment procedures remained

165   unchanged.

166   Before treatment, the affected tooth was anesthetized by local anesthesia. After local anesthesia

167   the endodontic access cavity was prepared. All teeth were isolated with a rubberdam. Root canals

168   were probed with K-steel files of ISO sizes 06, 08, 10 and 15, in order to create a glidepath up to

169   ISO 15 throughout all phases of the study. The working length was determined electrometrically

170   and/or by X-ray. The dentists prepared the root canals according to the details provided by the

171   manufacturers of the different rotary preparation systems. In the second study phase the root

172 canals were prepared with the WaveOne instruments according to the manufacturer's

173 instructions. If the dentists needed an apical preparation size that is not included in the WaveOne

174 System, the last ISO size was followed up by a single hand instrument of the desired size. During

175 rotary or reciprocating preparation the root canals were rinsed with 1-3% NaOCl between every

176 rotary instrument or in case of the SF system between every 3-4 picks with the WaveOne file.

177 After complete preparation of the root canals they were irrigated with a final irrigation of NaOCl

178 1-3% and a calcium hydroxide dressing or the root canal filling was placed. After that the tooth

179 was sealed provisionally bacteria-proof with a temporary bacteria tight seal. In the last

180 appointment the root canal filling was placed or in case of single-visit endodontics a definitive

181 coronal filling was applied.

182 **Outcomes**

183 The primary outcome of reduction of endodontic pain and the secondary outcome of

184 improvement of oral-health-related quality of life was measured with a patient questionnaire.

185 The questionnaire assessed the pain by the Visual Analog Scale (VAS 100) (Turk 2011) and the

186 oral-health-related quality of life with the items of the short version of the oral health impact

187 profile (OHIP-G-14) (John et al. 2004) wich is the German translation of OHIP-14 (Slade 1997).

188 The patients were asked about the biggest complaints (consisting of the VAS 100 and OHIP-

189 G14) without pain medication in the week before treatment and in the week before completion of

190 treatment. This was two weeks after initial treatment and in connection with either the placement

191 of the root canal filling or the definitive coronal filling of the tooth. The questionnaires were

192 filled in by the patients before treatment started or while local anesthesia was taking effect. Any

193 patient questions were answered by the dental team.

194   The time needed for root canal preparation was measured by the dental assistant staff. The

195   measurement started when the first rotating or reciprocating instrument was placed in the root

196   canal and ended when the last instrument was removed. The root canal recapitulations during

197   preparation and the irrigations were included in the time measurement. Probing and glidepath

198   preparation before using the rotary instruments were not included in the time measurement. Nor

199   were the final irrigation of the root canals and the placement of a dressing included. When a

200   tooth had several root canals, the total preparation time of all canals was measured and divided

201   by the number of root canals in order to determine the preparation time per canal.

202   **Further questionnaires and data collection**

203   In the PI questionnaires the clinical findings (dental chart, sensitivity test, percussion test, apical

204   pressure point, periodontal probing depth, radiographic presence of apical periodontitis, number

205   of prepared root canals, presence of fistulae), the time needed for root canal preparation,

206   instrument fractures and procedural events were documented. In addition, a consecutive patient

207   log was introduced to record, if possible, the patient's reason for rejecting participation. The

208   patient forms included the patient's informed consent to participate in the trial, a questionnaire

209   asking for demographic and basic medical data, and the above described pain questionnaire

210   which consisted of the VAS and the Items of OHIP-G-14.

211   All patients that qualified for participation in the study were informed about the study by the PI

212   personally. All PI forms were filled in by the assistant dental staff. The patient questionnaires

213   were filled in by the patients themselves. All patient questionnaires were pseudonymized and

214   collected in a sealed box. The PI forms were pseudonymized in the same way to be able to match

215   the patient data and the PI data in the subsequent evaluation.

216   The pain questionnaires were completed by the patients immediately before treatment started.

217   The demographic data could be provided at any time, but were requested on completion of the

218   treatment at the latest. The pain questionnaires were completed by the patients again 14 days

219   after treatment. The PIs' treatment was taken down on record. The time required for root canal

220   preparation was measured by the dental assistant staff.

221   The questionnaires were handed over to the principal investigator (AB) at the end of the first and

222   at the end of the second trial phase for evaluation.

223   **Safety measures**

224   Before treatment started, each patient participating in the study was informed about the

225   endodontic risks in the same way as it is usually done before endodontic therapy. The patient

226   was informed in particular about events, such as instrument fractures and other complications

227   that may occur in root canal preparation and could lead to the extraction of the tooth affected.

228   The patient was also informed, that root canal treatment is the last attempt to save a tooth. The

229   information was provided by the PI and an additional education and information questionnaire. If

230   in the course of the trial the complication of an instrument fracture occurred, the patient would

231   be informed about it. This information was provided by an information questionnaire for

232   instrument fractures.

233   **Sample Size Calculation and Statistics**

234   For sample size calculation we had to consider the sample design which was characterized by a

235   2-stage structure (dental practice, patient). This cluster sample made special demands on both

236   sample planning and the analysis of the results (Donner & Klar 2000).

237   To calculate the case number, the following parameters were defined: Significance level: 0.05,

238   Power: 80% and Number of clusters (dental practices): 10

239   Moreover, based on the analysis of similar studies (Pak 2012), the most realistic assumptions

240   possible were made about the clinically relevant difference of the VAS 100 (Visual Analog

241   Scale) and the ICC (Intra Cluster Correlation Coefficient) which is a measure for the

242   homogeneity in relation to a target variable of interest within the cluster: $\Delta VAS=20$ (20%) and

243   ICC=0.04.

244   For case number calculation a validated software tool was used which determined the number of

245   trial units (patients) per cluster (practice) on the basis of the parameters specified above

246   (Campbell et al. 2004).

247   The resulting number of patients was 28 per dental practice and every trial phase. This number

248   appeared realistic in terms of feasibility. In view of the basic statistical data on dental care in

249   Germany (KZBV 2011), a conservative estimate showed that one GDP performs about 60 root

250   canal treatments per year. This means that 10 participating GDPs should be able to recruit the

251   required case number in each of the trial phases.

252   The results were calculated with the SPSS (Version 21, Win x64) statistical system and SAS

253   (Version 9.2, Win x64). With the PROC MIXED procedure (Singer 1998) SAS offers options for

254   explicitly considering potential cluster effects (here: several data collection units per dental

255   office) in the overall regression model.

256   **Assessment of potential covariates**

257   Besides collecting the data for the primary outcome we assessed other dentist- and patient-

258   related as well as treatment-specific covariates. This was done with a questionnaire for

259   demographic information which also documented the patients' basic medical data. In addition,

260   the PIs recorded the dental chart and treatment-specific findings (tooth sensitivity before

261   treatment, apical translucency, percussion test, apical pressure point and fistula).

262   **Study termination criteria**

263   It was planned to terminate the study when two weeks after root canal preparation by the new

264   single-file method the patient's pain was 40% above the expected level. The study would also

265   have been terminated if single-file endodontics would have caused markedly more instrument

266   fractures than expected. If during the study more than three instrument fractures had already

267   occurred in the first 20 single-file treatment cases, the study would have been terminated.

268   **Results**

269   The ten PIs screened a total of 668 patients who met the study inclusion criteria. Of the 668

270   patients screened, 62 (9.2%) could not be included in the study primarily. In the course of the

271   study, 7 (1.0%) patients did not keep the agreed appointments for starting the treatment (Figure

272   2). The remaining 599 patients were included into the study for statistical analyses. Thus, the

273   number of patients actually included was 10.3% higher than the minimum required case number

274   of 560 determined by power analysis.

275   Table 1 shows the number of patients that were recruited by the PIs. During both trial periods the

276   individual participating GDPs recruited between 19 (PI 8) and 144 (PI 7) patients. All GDPs

277   together included 280 (46.7%) patients for MF treatment and 319 (53.3%) patients for SF

278   treatment. The distribution between the groups was nearly equal. The average age of the patients

279   was 50.2 (SD 15.7) years. The distribution of the patients to the various participating GDPs

280   differed (χ^2; P<0.001) whereas the age distribution across the study groups (MF and SF) showed

281   no statistical differences (T-Test; P=0.991) and was similar in both groups, i.e. 50.1 (SD 15.0)

282   years in MF and 50.2 (SD 16.4) in SF. More men (53.1%) were treated than women. The gender

283   distribution showed no statistical differences in the individual dental practices (χ^2; P=0.082) nor

284   in the study groups (χ^2; P=0.458). The various types of vocational qualification, as stated by the

285   study participants, differed in the individual practices (χ^2; P<0.001) but not in the study groups

286   (χ^2; P=0.102) (Table 2).

287   The return rates of the various questionnaires that had to be completed by the patients and the

288   participating GDPs were between 86% for the follow-up patient pain questionnaire and up to

289   97% for the questionnaire to be completed by the GDPs. The return rate of the patient pain

290   questionnaire before treatment was 94% and for the demographic data questionnaire it was 90%

291   (Table 3).

292   In the course of the study slightly more maxillary (53.7%) teeth were treated (Table 4). The

293   distribution of the different types of teeth showed no significant differences between the two

294   study groups (χ^2; P=0.255).

295   **Evaluation of the primary outcome criterion**

296   For the evaluation of the primary outcome, i.e. post-operative reduction of patients' endodontic

297   pain and improvement of oral-health-related quality of life, we measured pain reduction via VAS

298   100 and the OHIP-G14 score. Both values were measured before root canal treatment and 14

299   days after treatment. Then we compared the different study groups (MF and SF).

300   The mean pain score before root canal treatment for MF was 42.3 (SD 32.6) VAS and for SF

301   43.9 (SD 32.0) VAS and decreased to 10.0 (SD 18.6) VAS (MF) and 9.3 (SD 19.2) VAS (SF).

302   The mean OHIP-G 14 score before root canal treatment for MF was 12.5 (SD 10.6) and for SF

303   13.0 (SD 10.8) and decreased to 3.6 (SD 5.1) (MF) and 4.6 (SD 6.5) (SF).

304   For pain reduction and OHRQoL univariate analysis showed a very similar improvement in both

305   study groups (MF and SF)

306   (a)   Pain reduction 34.4 (SD 33.7) VAS (MF) vs. 35.0 (SD 35.4) VAS (SF) (p=0.8)

307   (b)   Improvement of oral-health-related quality of life according to OHIP-14 score: 9.4 (SD

308   10.3) (MF) vs. 8.5 (SD 10.2) (SF) (p=0.4)

309   The differences between the study groups were not significant.

310   Multivariate analysis of variance (MANOVA) taking the additional factor "single vs. multiple-

311   visit treatment" into account did not reveal any significant influence of the factors "study group

312   (MF or SF)", "single vs. multiple-visit" or an interaction of the two regarding pain reduction. For

313   improvement of OHIP-14 score there was an overall significant influence (p=0.03) with "single-

314   visit" treatments having a significantly (p=0.01) lower improvement than "multiple-visit". But

315   further analyses showed that OHIP-14 scores for "single-visit" treatments (10.7 (SD 9.9)) were

316   already significantly (p=0.06) lower before treatment than for "multiple-visit" treatments (13.6

317   (SD 10.9)) and dropped to almost the same levels before completion of treatment (4.1 (SD 5.9)

318   vs. 4.2 (SD 6.0).

319   **Evaluation of the secondary outcome criterion**

320 For the speed of root canal preparation, univariate analysis showed a significant difference

321 between the study groups MF and SF:

322 (c) Duration of treatment per root canal (in sec): 239 (MF) vs. 147 (SF) (P=0.003)

323 For (c) a multivariate analysis was made taking into consideration the dentist- and patient-related

324 as well as treatment-specific covariates:

325 ▪ Gender of dentist

326 ▪ Gender of patient

327 ▪ Age of patient

328 ▪ Comorbidities of the patient (none, hypertension, DM(I or II), asthma)

329 ▪ DMFT Index

330 ▪ Tooth

331 ▪ Tooth sensitivity before treatment (positive/negative)

332 ▪ Apical translucency (yes / no)

333 ▪ Percussion test (positive/negative)

334 ▪ Apical pressure point (yes / no)

335 ▪ Fistula (yes / no)

336 For the WO SF system a significantly shorter duration resulted in comparison to the MF systems

337 (121 sec; SD 37.40; p=0.01). The adjusted reduction in required preparation time was 32.8%

338 with the SF-System.

339 The root canal preparation with the WaveOne System produces the same results regarding

340 reduction of patients' endodontic related pain and oral health related quality of life, but the

341 preparation speed per root canal is faster.

342 **Discussion**

343 Our study showed that root canal treatment with MFs as well as with the SF WaveOne System

344 reduced the patients' endodontic related pain and improved oral health related quality of life

345 without statistically significant differences under conditions of general dental practice. The root

346 canal preparation with the SF system was faster.

347 *Measuring patients' endodontic pain and oral-health-related quality of life*

348 Pain intensity can be measured with various methods, e.g. the Visual Analog Scale (VAS), the

349 Numerical Rating Scale (NRS) or the Verbal (Categorical) Rating Scale (VRS). The VRS is an

350 easy-to-apply measuring method, but forces the study subjects to select a wording which may not

351 represent an adequate description of the pain they feel. Moreover, the VRS depends on a clear

352 and unequivocal understanding of the language (Turk 2011). The NRS and VAS are

353 uncomplicated measuring methods for the pain felt and both show good evidence of construct

354 validation (Turk 2011). In the present study, the VAS was selected because it offers a large

355 number of scores. This makes the VAS more sensitive to changes in the pain intensity felt than

356 other scales offering fewer reply categories (Turk 2011). In addition, the VAS is widely used in

357 the endodontic literature (King et al. 2012; Martin-Gonzalez et al. 2012; Pak 2012; Udoye &

358 Jafarzadeh 2011). The limitation to one scale is both feasible and adequate (Attar et al. 2008).

359 Pain assessment alone gives no information about the patients' oral-health-related quality of life

360 which represents a patient-relevant outcome (Pfaff et al. 2009). The only validated German-

361 language measuring instrument for the oral-health-related quality of life is the OHIP (John et al.

362 2003). To limit the questionnaire to a practicable length, the OHIP-G-14 was used in our study

363 (John et al. 2004). In 2011, when the study was planned, there existed two endodontic studies

364   which chose the OHIP-14 questionnaire for endodontic issues (Dugas et al. 2002; Gatten et al.

365   2011).

366   *Description of the results*

367   In the present study, we compared two clinical short-term outcome parameters of two basic

368   methods of root canal preparation.

369   We defined the most important patient-relevant outcomes as reduction of pain by endodontic

370   therapy and the improvement of oral-health-related quality of life. When we were planning this

371   study in 2011, shortly after the introduction of reciprocating SF endodontic instruments

372   (WaveOne and Reciproc), there was no information about the clinical performance of these

373   instruments. The new system should have at least a similar or an enhanced clinical outcome

374   compared to the conventional (MF) systems. This is an important condition when a new

375   technology is introduced. Moreover, we expected greater speed and thus greater efficiency of the

376   SF System as a relevant result. As far as the authors know, no studies had been made that

377   compared the clinical outcome of MF and SF systems at the time of the study planning in 2011.

378   The reduction of pain and the improvement of oral health related quality of life as a result of

379   endodontic treatment were not different in the two experimental groups (MF and SF) in the

380   second week after treatment. Both methods were equally effective in reducing endodontic pain.

381   The mean pain intensity of about 43.2 (SD 32.2) VAS before root canal treatment and about 9.5

382   (SD 19.0) VAS after treatment agrees with the results obtained by other researchers who

383   investigated pain reduction after endodontic therapy (Ehrmann et al. 2003; Pak & White 2011).

384   A review found that pain levels before endodontic therapy of 54 VAS and a standard deviation of

385   24 VAS are a common average in endodontic studies and decrease to less than 10 VAS on

386 average within 7 days (Ehrmann et al. 2003; Pak & White 2011). Also Ehrmann et al. (2003)

387 found very similar values to those in our study (44.4 (SD 26.9) VAS). The mean pain measured 4

388 days after endodontic therapy decreased to 7.5 (SD 15.5) VAS. The mean pain reduction of 36.9

389 (SD 29.0) VAS was very similar to the values in our study. The improvement of the OHIP-14

390 with mean scores of 12.8 (SD 10.6) before therapy found in our study are very comparable to

391 another study where a mean OHIP-14 score of 15.4 (SD 10.5) was found before endodontic

392 treatment (Liu et al. 2014). Also other studies report that endodontic treatment leads to an

393 improvement of the oral-health-related quality of life (Dugas et al. 2002; Hamasha & Hatiwsh

394 2013). This finding was confirmed by our study.

395 Recently a couple of randomized controlled trials (RCT) have been published, that evaluated the

396 pain reduction and/or the improvement in quality of life of single file systems (Kherlakian et al.

397 2016; Pasqualini et al. 2015; Relvas et al. 2015). In the first RCT (Kherlakian et al. 2016) two SF

398 reciprocating systems (Reciproc (VDW, Munich, Germany) and WaveOne (Dentsply)) and one

399 MF system (ProTaper Next (Dentsply)) were compared. Only asymptomatic vital teeth were

400 treated with the different systems. Therefore patients did not have pain before treatment. Also

401 after treatment, pain rates on a categorized VAS 100 score were also very low and showed no

402 significant differences between the systems. The second RCT (Relvas et al. 2015) compared one

403 reciprocating SF system (Reciproc (VDW, Munich, Germany) with a MF system (ProTaper

404 (Dentsply)). Only asymptomatic teeth with apical periodontitis were included in the trial.

405 Therefore patients were pain-free before treatment. Pain measurement was not performed with

406 the VAS. Therefore results can be hardly compared with our study. The different instrument

407 systems showed no statistically significant differences in postoperative pain scores after 24h and

408 72h. The third RCT (Pasqualini et al. 2015) investigated the ProTaper MF and the WaveOne SF

409   system. Compared were primary root canal treatments of every clinical condition (symptomatic,

410   asymptomatic, vital and non-vital cases). Mean pain on VAS was 35.2 for SF before treatment

411   and 24.6 for MF decreasing to very low rates of 1.3 (SF) and 0.9 (MF) after seven days. This is

412   different to our study, where the mean pain scores were higher before treatment, but more equal.

413   And also our mean pain scores in the second week after treatment were higher than the score

414   found in the above mentioned study. If this can be interpreted as an effectiveness gap regarding

415   the success of these instruments in general dental practice compared to specialized care providers

416   remains unclear, because our initial VAS scores were higher and therefore perhaps naturally

417   need a longer time period to drop to low scores. Recently our research group published a smaller,

418   but very similar study comparing hand instrumentation with Reciproc for root canal preparation

419   under conditions of general dental practice (Bartols et al. 2016). The mean pain score before root

420   canal treatment with hand instruments was 43.6 (SD 30.7) VAS and with Reciproc it was 41.2

421   (SD 27.7) VAS, which is perfectly comparable with the initial values of the present study where

422   the initial VAS scores were 42.3 (SD 32.6) VAS for MF and 43.9 (SD 32.0) VAS for SF. Within

423   the same period of time the scores decreased in all four groups to values in the range between 9.3

424   and 11.5 (with SDs of 16.5-19.2) VAS with therefore only minimal differences. Also regarding

425   OHRQoL the OHIP-G-14 scores of all four groups show only minimal differences before

426   treatment and before completion of therapy (hand instruments 9.2 (SD 9.6) decreasing to 3.4 (SD

427   5.4), Reciproc 10.4 (SD 9.6) decreasing to 3.5 (SD 6.1), MF 12.5 (SD 10.6) decreasing to 3.6

428   (SD 5.1) and SF WaveOne 13.0 (SD 10.8) decreasing to 4.6 (SD 6.5). Therefore it can be

429   concluded, that all four techniques investigated show the same clinical outcome regarding pain

430   reduction and improvement in OHRQoL under routine care conditions.

431   As the main outcome parameter was the reduction of endodontic pain after treatment, the

432   different root canal preparation techniques and their influence on postoperative pain has to be

433   considered. Recently it was demonstrated that different root canal preparation techniques lead to

434   the expression of different levels of inflammatory neuropeptides in the periapical periodontal

435   ligament linked with the possible emergence of symptomatic apical periodontitis (Caviedes-

436   Bucheli et al. 2013). It is believed that this is connected to the different amounts of extruded

437   debris beyond the apical foramen (Caviedes-Bucheli et al. 2016). Since nearly all root canal

438   instrumentation techniques including hand instrumentation as well as engine driven instruments

439   lead to apical extrusion of debris (Al-Omari & Dummer 1995; Bürklein & Schäfer 2012; Capar

440   et al. 2014; De-Deus et al. 2010) in most cases there will be an inflammatory response to a

441   certain extent. In an in-vitro studies reciprocating instruments extruded more debris than rotary

442   instruments (Bürklein & Schäfer 2012) with Reciproc producing most debris while another in-

443   vitro studies found Reciproc to produce significantly less extruded debris compared to rotary

444   techniques (Kocak et al. 2013). The only clinical studies measuring the expression of

445   inflammatory neuropeptides in the periodontal ligament found that the instrument design of

446   engine driven root canal instruments has a greater impact on expression of neuropeptides than

447   the instrumentation technique (Caviedes-Bucheli et al. 2016). Because of this contradictory data

448   situation and the limited knowledge if the amount of expressed neuropeptides can be directly

449   correlated to the perceived pain it remains unclear if there is an impact on the postoperative pain

450   levels of patients after root canal treatment. In our study the preoperative levels of pain, their

451   improvement and the postoperative VAS pain levels were very similar and very much

452   comparable to our previously published study (Bartols et al. 2016). Therefore in the

453 heterogeneous situation of clinical cases, the impact of the root canal preparation systems used

454 seems to be limited regarding postoperative pain relief.

455 A significant difference was found in the speed of root canal preparation. The preparation time

456 required when using WO instruments was on average 92 seconds shorter than the time required

457 with MF systems. This time is probably saved because the WO system does not require any

458 instrument changes. As changing instruments cannot be avoided with MF systems, the time

459 needed for it was included in the time measurement. An in-vitro study reported that root canal

460 preparation with WaveOne instruments in contrast to MF systems is about 100 seconds faster

461 (Bürklein et al. 2012). This time benefit per canal was also observed in our study. In both study

462 designs the instrumentation time included instrument changes, cleaning of instruments and

463 irrigation of the root canal. Therefore results are comparable. Thus, there is nearly no

464 effectiveness gap of the method. This was not necessarily to be expected as unlike root canal

465 preparation in the laboratory the preparation in the patient's mouth is more complicated due to

466 patient-related factors, such as mouth opening, restlessness of the patient etc. The time saved in

467 canal preparation can be beneficially reinvested in additional root canal disinfection (van der

468 Sluis et al. 2009).

469 In general, the endodontic literature proves that pain that existed before endodontic therapy will

470 be reduced by root canal therapy (Ehrmann et al. 2003; Genet et al. 1986; Pak & White 2011).

471 Comparative studies on endodontic pain have so far mainly compared different types of pain

472 medication (Attar et al. 2008; Ryan et al. 2008), different types of root canal dressings (Ehrmann

473 et al. 2003; Torabinejad et al. 1994) and differences between single-visit vs. multiple-visit

474 treatment (Prashanth et al. 2011; Su et al. 2011). For single- versus multiple-visit treatment,

475 studies found no differences for one week postoperative pain levels (Figini et al. 2008; Prashanth

476   et al. 2011). This suits our results, because we also did not find differences in our analyses

477   regarding single- versus multiple-visit treatments regarding pain reduction. For OHRQoL there

478   was a difference in improvement of OHIP-14 scores between single- and multiple-visit

479   treatments. But as the initial OHIP-14 scores were significantly lower in the single-visit group

480   than the initial scores in the multiple-visit group, we conclude that the PIs primarily treated

481   "safe" cases with low initial OHIP-14 scores as single-visit.

482   Clinical trials comparing pain after root canal preparation with different instrument systems are

483   rare (Gambarini et al. 2013; Kherlakian et al. 2016; Pasqualini et al. 2015; Relvas et al. 2015)

484   and have mostly low case numbers (N=30-70 per experimental group). The authors do not know

485   of any large-scale clinical comparative studies with high case numbers reliably reflecting the

486   dental practice reality. Research in practice networks offers an environment which allows to

487   generate case numbers high enough for clinical trials (Nixdorf et al. 2012). In this way new

488   research opportunities are created that can also be applied to other issues of endodontics or other

489   fields of dentistry.

490   All three outcome parameters reflect patient relevant short term success criteria, that are not

491   necessarily connected to the long term success of the treatments performed. To the knowledge of

492   the authors until now there is only one study that investigated also the long term success of

493   technological change from stainless steel instrumentation to rotary instrumentation in a general

494   dental practitioner situation in the Swedish Public Dental Service (Koch et al. 2015). While tooth

495   survival was higher in teeth treated post-education with rotary instruments there was no

496   improvement in periapical health. Only surrogate parameters like the technical quality of the

497   treatments improved. Studies investigating the long term outcome after technological change to

498   reciprocating technique in endodontics are not known to the authors. Therefore further research

499   regarding long term results of reciprocating techniques in general dental practice is needed. *Study*

500   *design and feasibility*

501   Generally this study was planned as a health services research study. Therefore it was never

502   intended to compare two treatment groups in a classical clinical trial setting. We wanted to

503   investigate the effects of technological change in endodontic treatment in general dental practices

504   and chose therefore a study design in a timeline sequence and not a study design with parallel

505   treatments groups. We accompanied the technological change in endodontic treatment methods

506   in everyday dental practice to uncover possible "shortcomings" or effectiveness gaps by

507   structured observation, which were basically not found for short-term outcomes as pain reduction

508   and improvement of OHRQoL.

509   In the present study, the number of recruited patients agreed with the initial case number

510   planning. The planning therefore seemed to be based on realistic assumptions. The return rates of

511   the collected study data and questionnaires were high and the patients were adequately followed

512   up. Judged by these requirements, research can be conducted in dental practices in an adequate

513   way. (Kohout et al. 2015)

514   Ten GDPs agreed to participate in the study as PIs. This exactly equaled the number underlying

515   the power analysis. The recruitment of the minimum number of required GDPs poses the risk

516   that the case numbers aimed at cannot be reached. As studies of this type are rare in endodontics,

517   there are no broadly-based typical figures available on the experience regarding the recruitment

518   of PIs. There is only one study pursuing a similar approach by observing the results obtained in

519   dental practices (Nixdorf et al. 2012). That study was designed as an observational study to

520   measure pain and burden connected with initial orthograde root canal treatment. 62 GDPs

521   participated in the study, whereas 48 had been aimed for in case number planning. This

522   corresponds to an over-recruitment rate of 29% (Nixdorf et al. 2012).

523   Contrary to that study (Nixdorf et al. 2012), the present study takes an approach to compare

524   different treatment methods, which makes considerably higher demands on the participating

525   GDPs. The GDPs had to undergo training to learn how to prepare the root canals with the SF

526   WaveOne instruments and, at the same time, they had to care for two therapeutic groups and to

527   recruit themselves the patients for each. The GDPs did not get any financial support. As an

528   incentive they were offered a payment of EUR 5 for every evaluable/analyzable case which,

529   however, most colleagues did not take. The training for the use of WaveOne instruments was

530   provided free of charge to the GDPs. In addition, in the second study phase Dentsply Maillefer

531   (Ballaigues, Switzerland) made available the required WaveOne files free of charge and loaned

532   the GDPs the Wave-One motors. In view of the fact that the literature describes serious

533   resentments of German physicians against practice-based clinical trials (Hummers-Pradier et al.

534   2012; Hummers-Pradier et al. 2008; KZBV 2011), it is a special success to recruit 10 GDPs.

535   Moreover a recently published similar study of our research group showed, that 3 of 9 PIs could

536   not cope with the organizational demands of a study like this and could not contribute any cases

537   for evaluation (Bartols et al. 2016).

538   The participating GDPs documented treatments that were required anyway. The practice routine

539   had to be changed for the documentation requirements of the study, but the organizational work

540   with the study participants was mainly delegated to the assistant dental staff. This certainly is one

541   reason for the good feasibility of the study. Moreover, there were no special demands on the

542   patients, so that their willingness to participate in the study was very high. The GDPs screened

543   668 patients and actually enrolled 599 in the study so that on average every GDP screened about

544 1.1 patients to include one in the study. Compared with the study of Nixdorf et al. (2012) who

545 screened 1.5 patients for each subject included in the study, this is a high rate of inclusion and

546 shows the patients' great willingness to participate in a clinical trial of the extent described here.

547 Altogether 599 participants were recruited, whereas 560 would have been needed. This is an

548 over-recruitment of not quite 7%, so that, on average, the case number aimed for was reached.

549 However, the individual case numbers differed very much (Table 1).

550 The two-phase study design split into separate periods increased the GDPs' willingness to

551 participate in the clinical study because it limited the organizational effort for the study.

552 Although this means that the present study was not based on randomization, generally considered

553 the optimum study design (Friedman et al. 2010; Hulley 2013), the clear time split of the study

554 groups prevented randomization errors and selection bias at the level of the participating dental

555 practices in the sense of manipulating the patient randomization to each of the study groups and

556 was also used in another study investigating endodontic technological change in general dental

557 practice (Koch et al. 2015). The consecutive sample used in the present study also counteracted

558 the volunteer bias (volunteerism) (Hulley 2013). Additionally the broad inclusion criteria for the

559 participating patients made recruitment feasible for the PIs and reflects in this way the conditions

560 of everyday general dental practice.

561 **Conclusion**

562 Concerning the reduction of endodontic pain and improvement of oral health related quality of

563 life, the WaveOne SF system shows no statistical difference to MF systems under the conditions

564 of general dental practice. The speed of preparation of root canals appears to be higher with the

565 WaveOne SF instruments.

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744

Number of patients recruited by practitioner-investigator (P-I) and study group distribution

1   Table 1 Number of patients recruited by practitioner-investigator (P-I) and study group distribution

P-I ID	Instrument type						Total N	
	MF	Target	Actual	SF	Target	Actual	Target	Actual
1	BioRaCe	28	28	WaveOne	28	28	56	56
2	RaCe	28	45	WaveOne	28	56	56	101
3	BioRaCe	28	30	WaveOne	28	23	56	53
4	BioRaCe	28	22	WaveOne	28	23	56	45
5	Alpha Kite	28	38	WaveOne	28	32	56	70
6	BioRaCe	28	26	WaveOne	28	20	56	46
7	Mity Roto Files	28	49	WaveOne	28	95	56	144
8	BioRaCe	28	15	WaveOne	28	4	56	19
9	BioRaCe	28	10	WaveOne	28	17	56	27
10	BioRaCe	28	17	WaveOne	28	21	56	38
Total N		280	**280**		280	**319**	560	**599**

2   MF=multiple-file system, SF= single-file system (WaveOne)

Table 2

1 Table 2 Important socio-demographic characteristics and DMF-T of study participants by study center and study group

Socio-demographic data of participants	Total	Study center										Study group	
		1	2	3	4	5	6	7	8	9	10	MF	SF
Age (yr)													
Mean	**50**	52	46	55	53	49	52	47	59	49	61	50	50
SD	**16**	15	13	17	16	13	17	16	16	15	14	15	16
Range	**18-88**	20-83	18-79	19-88	19-79	23-85	23-84	19-80	35-84	19-74	29-79	18-85	19-88
Total N	**548**	55	96	50	45	48	37	141	18	25	33	245	303
Gender (female)													
N (%)	**269**	28 (51)	42 (42)	35 (66)	16 (36)	23 (43)	21 (48)	68 (48)	7 (37)	16 (59)	13 (37)	126 (48)	143 (46)
Total N	**573**	55	101	53	44	53	44	142	19	27	35	259	314
DMF-T													
Mean	**12.0**	15.5	7.7	16.2	17.0	7.8	11.4	9.3	17.4	10.0	20.7	12.0	12.0
Total N	**550**	56	96	50	45	50	45	137	15	18	38	249	301
Highest education													
Completed apprenticeship N (%)	241	31 (58)	40 (40)	15 (41)	22 (65)	17 (35)	15 (37)	57 (43)	13 (87)	20 (74)	11 (34)	95 (41)	146
Technical/Vocational school N	97	10 (19)	12 (12)	12 (32)	7 (21)	13 (27)	9 (22)	20 (15)	1 (7)	4 (15)	9 (28)	52 (22)	45 (16)
University/College N (%)	92	7 (13)	33 (33)	1 (3)	0 (0)	10 (21)	12 (29)	17 (13)	1 (7)	0 (0)	11 (34)	47 (20)	45 (16)
Other N (%)	40	1 (2)	9 (9)	3 (8)	1 (3)	4 (8)	4 (10)	17 (13)	0 (0)	1 (4)	0 (0)	19 (8)	21 (7)
No N (%)	51	4 (8)	6 (6)	6 (16)	4 (12)	4 (8)	1 (2)	23 (17)	0 (0)	2 (7)	1 (3)	21 (9)	30 (11)
Total N	521	53	100	37	34	48	41	134	15	27	32	234	287

3 MF multiple-file system SF single-file WaveOne

Important socio-demographic characteristics and DMF-T of study participants by study center and study group

Table 2 Important socio-demographic characteristics and DMF-T of study participants by study center and study group

Socio-demographic data of participants	Total	Study center											Study group	
		1	2	3	4	5	6	7	8	9	10	MF	SF
Age (yr)														
Mean	**50**	52	46	55	53	49	52	47	59	49	61	50	50
SD	**16**	15	13	17	16	13	17	16	16	15	14	15	16
Range	**18-88**	20-83	18-79	19-88	19-79	23-85	23-84	19-80	35-84	19-74	29-79	18-85	19-88
Total N	**548**	55	96	50	45	48	37	141	18	25	33	245	303
Gender (female)														
N (%)	**269**	28 (51)	42 (42)	35 (66)	16 (36)	23 (43)	21 (48)	68 (48)	7 (37)	16 (59)	13 (37)	126 (49)	143 (46)
Total *N*	**573**	55	101	53	44	53	44	142	19	27	35	259	314
DMF-T														
Mean	**12.0**	15.5	7.7	16.2	17.0	7.8	11.4	9.3	17.4	10.0	20.7	12.0	12.0
Total *N*	**550**	56	96	50	45	50	45	137	15	18	38	249	301
Highest education														
Completed apprenticeship N (%)	**241**	31 (58)	40 (40)	15 (41)	22 (65)	17 (35)	15 (37)	57 (43)	13 (87)	20 (74)	11 (34)	95 (41)	146 (51)
Technical/Vocational school N (%)	**97**	10 (19)	12 (12)	12 (32)	7 (21)	13 (27)	9 (22)	20 (15)	1 (7)	4 (15)	9 (28)	52 (22)	45 (16)
University/College N (%)	**92**	7 (13)	33 (33)	1 (3)	0 (0)	10 (21)	12 (29)	17 (13)	1 (7)	0 (0)	11 (34)	47 (20)	45 (16)
Other N (%)	**40 (8)**	1 (2)	9 (9)	3 (8)	1 (3)	4 (8)	4 (10)	17 (13)	0 (0)	1 (4)	0 (0)	19 (8)	21 (7)
No N (%)	**51**	4 (8)	6 (6)	6 (16)	4 (12)	4 (8)	1 (2)	23 (17)	0 (0)	2 (7)	1 (3)	21 (9)	30 (11)
Total N	**521**	53	100	37	34	48	41	134	15	27	32	234	287

MF multiple-file system SF single-file WaveOne

Questionnaire return rates for the enrolled participants

1   Table 3 Questionnaire return rates for the enrolled participants

Description	Timing	N (received)	N (expected)	%
Patient survey demographic data	before root filling	538	599	90
Pain and OHIP-14 survey before treatment	1st appointment	565	599	94
Dentist survey for treatment parameters	all appointments	582	599	97
Patient 2 weeks follow-up survey	2 weeks after RCT	518	599	86

Table 4

1   Table 4 Descriptive data of teeth treated by location, study center (PI) and study group

Tooth type		Study Center 1 Mean	(SD)	N	2 Mean	(SD)	N	3 Mean	(SD)	N	4 Mean	(SD)	N	5 Mean	(SD)	N	6 Mean	(SD)	N
Maxillary anteriors	No. of treated root canals (mean)	1.0	0.0	7	1.0	0.0	13	1.0	0.0	11	1.0	0.0	5	1.0	0.0	2	1.0	0.0	7
	Time for treatment per tooth (sec.)	194.0	140.4	7	139.7	83.1	13	114.2	53.1	11	132.8	77.7	5	169.0	58.0	2	78.7	38.2	7
	Time for treatment per root canal (sec.)	225.4	145.6	7	139.7	83.1	13	114.2	53.1	11	132.8	77.7	5	169.0	58.0	2	163.6	193.8	7
Maxillary premolars	No. of treated root canals (mean)	1.6	0.5	11	1.4	0.5	17	1.7	0.5	7	2.0	0.0	9	2.0	0.0	5	1.7	0.5	6
	Time for treatment per tooth (sec.)	265.3	117.9	11	188.1	104.8	17	274.7	116.9	7	150.6	42.6	10	267.8	154.0	5	134.8	50.1	6
	Time for treatment per root canal (sec.)	180.0	75.0	11	141.6	69.4	17	171.1	105.5	7	74.3	22.4	9	161.1	22.9	5	97.6	31.2	6
Maxillary molars	No. of treated root canals (mean)	3.2	0.4	13	2.9	0.4	26	3.1	0.7	11	3.0	0.7	9	3.3	0.5	14	3.1	0.4	8
	Time for treatment per tooth (sec.)	905.3	1618.0	12	305.5	141.4	26	354.3	162.0	11	174.8	86.7	9	303.1	269.3	14	103.8	90.7	9
	Time for treatment per root canal (sec.)	252.3	397.1	12	109.3	53.2	26	113.0	42.1	11	70.3	17.0	9	151.0	47.5	14	63.7	63.7	8
Mandibular anteriors	No. of treated root canals (mean)	1.0		1	1.0	0.0	3	1.0	0.0	7	1.0	0.0	3	1.0	0.0	2	1.0	0.0	2
	Time for treatment per tooth (sec.)	114.0		1	122.7	76.2	3	113.7	38.4	7	55.0	35.5	3	229.5	113.8	2	50.0	31.1	2
	Time for treatment per root canal (sec.)	114.0		1	122.7	76.2	3	113.7	38.4	7	55.0	35.5	3	229.5	113.8	2	50.0	31.1	2
Mandibular premolars	No. of treated root canals (mean)	1.0	0.0	5	1.0	0.0	15	1.0	0.0	6	1.0	0.0	7	1.0	0.0	3	1.0	0.0	13
	Time for treatment per tooth (sec.)	162.2	101.9	5	99.7	74.7	15	162.8	65.2	6	99.3	55.8	7	327.0	96.0	3	60.7	45.9	13
	Time for treatment per root canal (sec.)	162.2	101.9	5	104.9	74.0	15	162.8	65.2	6	99.3	55.8	7	327.0	96.0	3	122.5	128.2	13
Mandibular molars	No. of treated root canals (mean)	3.0	0.0	19	2.8	0.5	25	2.7	0.7	10	2.8	0.8	11	3.2	0.5	28	2.9	0.9	9
	Time for treatment per tooth (sec.)	368.1	205.1	19	271.9	167.1	27	247.9	92.2	10	198.4	48.5	11	361.0	220.9	28	154.1	106.1	9

	Time for treatment per root canal (sec.)	126.3	65.2	19	98.6	57.3	25	96.6	42.4	10	71.8	13.1	11	135.3	53.1	28	93.7	53.8	9
Totals	No. of treated root canals (mean)	2.3	1.0	56	2.0	1.0	99	1.9	1.0	52	2.1	1.0	44	2.8	0.9	54	1.8	1.0	45
	Time for treatment per tooth (sec.)	419.2	790.9	55	219.4	147.8	101	217.8	134.9	52	150.8	71.2	45	323.5	217.9	54	99.3	76.8	46
	Time for treatment per root canal (sec.)	180.2	200.7	55	115.9	66.0	99	123.8	61.7	52	82.2	41.3	44	157.1	69.2	54	106.1	109.7	45

Descriptive data of teeth treated by location, study center (PI) and study group

Table 4 Descriptive data of teeth treated by location, study center (PI) and study group

Tooth type		Study Center 1 Mean	(SD)	N	2 Mean	(SD)	N	3 Mean	(SD)	N	4 Mean	(SD)	N	5 Mean	(SD)	N	6 Mean	(SD)	N	7 Mean	(SD)	N	8 Mean	(SD)	N	9 Mean	(SD)	N	10 Mean	(SD)	N	Study Group Multiple file Mean	(SD)	N	Single file Mean	(SD)	N	Total Mean	(SD)	N
Maxillary anteriors	No. of treated root canals (mean)	1.0	0.0	7	1.0	0.0	13	1.0	0.0	11	1.0	0.0	5	1.0	0.0	2	1.0	0.0	7	1.1	0.3	23	1.0	0.0	2	1.0	0.0	2	1.0	0.0	8	1.0	0.0	37	1.1	0.3	43	1.0	0.2	80
	Time for treatment per tooth (sec.)	194.0	140.4	7	139.7	83.1	13	114.2	53.1	11	132.8	77.7	5	169.0	58.0	2	78.7	38.2	7	155.0	201.6	22	50.0	14.1	2	93.0	52.3	2	1500.0	2342.3	8	252.9	227.7	37	293.4	1116.2	42	274.4	824.2	79
	Time for treatment per root canal (sec.)	225.4	145.6	7	139.7	83.1	13	114.2	53.1	11	132.8	77.7	5	169.0	58.0	2	163.6	193.8	7	155.0	201.8	22	50.0	14.1	2	93.0	52.3	2	1500.0	2342.3	8	274.9	229.6	37	293.4	1116.2	42	284.7	824.2	79
Maxillary premolars	No. of treated root canals (mean)	1.6	0.5	11	1.4	0.5	17	1.7	0.5	7	2.0	0.0	9	2.0	0.0	5	1.7	0.5	6	1.8	0.4	20	2.0	0.0	2	1.6	0.5	8	1.3	0.5	7	1.7	0.5	53	1.6	0.5	39	1.7	0.5	92
	Time for treatment per tooth (sec.)	265.3	117.9	11	188.1	104.8	17	274.7	116.9	7	150.6	42.6	10	267.8	154.0	5	134.8	50.1	6	296.6	273.8	19	87.5	17.7	2	115.1	128.4	8	950.0	435.0	6	277.0	210.9	53	248.5	329.3	38	265.1	265.5	91
	Time for treatment per root canal (sec.)	180.0	75.0	11	141.6	69.4	17	171.1	105.5	7	74.3	22.4	9	161.1	22.9	5	97.6	31.2	6	162.1	130.8	19	43.8	8.8	2	148.2	70.0	8	710.0	210.1	6	192.5	154.3	52	164.4	197.8	38	180.7	173.5	90
Maxillary molars	No. of treated root canals (mean)	3.2	0.4	13	2.9	0.4	26	3.1	0.7	11	3.0	0.7	9	3.3	0.5	14	3.1	0.4	8	3.1	0.3	37	2.3	0.6	3	3.2	0.8	6	3.0	0.4	11	3.1	0.5	58	3.0	0.5	80	3.1	0.5	138
	Time for treatment per tooth (sec.)	905.3	1618.0	12	305.5	141.4	26	354.3	162.0	11	174.8	86.7	9	303.1	269.3	14	103.8	90.7	9	463.9	640.6	35	255.0	336.6	2	198.7	248.2	6	2100.0	523.8	11	795.1	744.8	57	322.7	757.9	78	522.1	785.3	135
	Time for treatment per root canal (sec.)	252.3	397.1	12	109.3	53.2	26	113.0	42.1	11	70.3	17.0	9	151.0	47.5	14	63.7	63.7	8	164.9	226.7	35	86.4	110.2	2	133.7	46.9	6	700.9	141.5	11	273.1	246.1	57	120.3	204.8	77	185.3	235.0	134
Mandibular anteriors	No. of treated root canals (mean)	1.0		1	1.0	0.0	3	1.0	0.0	7	1.0	0.0	3	1.0	0.0	2	1.0	0.0	2	1.1	0.4	14	1.0	0.0	3	1.0	0.0	2	1.0	0.0	2	1.1	0.3	16	1.0	0.2	23	1.1	0.2	39
	Time for treatment per tooth (sec.)	114.0		1	122.7	76.2	3	113.7	38.4	7	55.0	35.5	3	229.5	113.8	2	50.0	31.1	2	271.1	268.9	14	45.0	5.0	3	172.0	1.4	2	1320.0	1018.2	2	276.1	239.8	16	195.6	421.0	23	228.6	356.2	39
	Time for treatment per root canal (sec.)	114.0		1	122.7	76.2	3	113.7	38.4	7	55.0	35.5	3	229.5	113.8	2	50.0	31.1	2	249.3	259.6	14	45.0	5.0	3	172.0	1.4	2	1320.0	1018.2	2	259.2	229.2	16	194.1	421.5	23	220.8	353.1	39
Mandibular premolars	No. of treated root canals (mean)	1.0	0.0	5	1.0	0.0	15	1.0	0.0	6	1.0	0.0	7	1.0	0.0	3	1.0	0.0	13	1.1	0.3	28	1.0	0.0	2	1.0	0.0	6	1.0	0.0	2	1.0	0.2	35	1.0	0.2	52	1.0	0.2	87
	Time for treatment per tooth (sec.)	162.2	101.9	5	99.7	74.7	15	162.8	65.2	6	99.3	55.8	7	327.0	96.0	3	60.7	45.9	13	188.0	291.0	28	75.0	21.2	2	187.0	168.1	6	810.0	466.7	2	233.3	251.3	35	110.4	175.1	52	159.8	216.4	87
	Time for treatment per root canal (sec.)	162.2	101.9	5	104.9	74.0	15	162.8	65.2	6	99.3	55.8	7	327.0	96.0	3	122.5	128.2	13	162.7	188.8	27	75.0	21.2	2	204.2	149.4	6	810.0	466.7	2	237.7	156.6	34	114.2	174.4	52	163.0	177.4	86
Mandibular molars	No. of treated root canals (mean)	3.0	0.0	19	2.8	0.5	25	2.7	0.7	10	2.8	0.8	11	3.2	0.5	28	2.9	0.9	9	3.3	0.5	20	2.8	0.4	6	2.7	0.6	3	3.3	0.5	8	3.0	0.6	61	3.0	0.6	78	3.0	0.6	139
	Time for treatment per tooth (sec.)	368.1	205.1	19	271.9	167.1	27	247.9	92.2	10	198.4	48.5	11	361.0	220.9	28	154.6	106.1	9	589.8	824.3	19	113.2	103.1	6	181.0	158.2	3	2002.5	501.9	8	626.8	673.7	62	257.3	326.8	78	421.0	540.6	140
	Time for treatment per root canal (sec.)	126.3	65.2	19	98.6	57.3	25	96.6	42.4	10	71.8	13.1	11	135.3	53.1	28	93.7	53.8	9	181.9	263.5	19	53.9	22.2	5	108.6	33.4	3	635.6	168.4	8	219.6	205.0	60	93.6	107.3	77	148.8	169.1	137
Totals	No. of treated root canals (mean)	2.3	1.0	56	2.0	1.0	99	1.9	1.0	52	2.1	1.0	44	2.8	0.9	54	1.8	1.0	45	2.0	1.0	142	1.9	0.9	18	1.9	1.0	27	2.1	1.1	38	2.1	1.0	260	2.1	1.0	315	2.1	1.0	575
	Time for treatment per tooth (sec.)	419.2	790.9	55	219.4	147.8	101	217.8	134.9	52	150.8	71.2	45	323.5	217.9	54	99.3	76.8	46	332.5	506.1	137	102.9	120.4	17	159.6	159.0	27	1650.8	1214.3	37	464.6	554.9	260	248.4	608.0	311	346.8	593.8	571
	Time for treatment per root canal (sec.)	180.2	200.7	55	115.9	66.0	99	123.8	61.7	52	82.2	41.3	44	157.1	69.2	54	106.1	109.7	45	173.5	211.9	136	57.2	34.8	16	150.7	86.3	27	900.4	1114.8	37	238.9	206.2	256	147.1	453.5	309	188.7	365.5	565

Project plan

MF: Multiple-file system, SF: Single-file system WaveOne

	Year 1 (09/2011 – 09/2012)											Year 2 (from 10/2012)		

MI Inclusion phase | Follow-up phase

SF Training

SF Inclusion phase | Follow-up phase

Evaluation

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15

Flow Diagram

Assessed for eligibility (n=668)

Enrollment

Excluded (n=62)
- Declined to participate (n=9)
- Other reasons (n=33)
- Practitioner forgot to ask (n=2)
- Communication problems (n=5)
- Patient did not have time (n=4)
- No reason stated (n=3)
- Patient <18 years (n=2)
- Emergency treatment only (n=4)

Allocated (n=606)

Allocation

Allocated to MF RCT (1st phase) (n=284)
- Received allocated intervention (n=283)
- Did not receive allocated intervention (patient did not return) (n=1)

Allocated to SF RCT (2nd phase) (n=322)
- Received allocated intervention (n=319)
- Did not receive allocated intervention (patient did not return) (n=3)

Follow-Up

Lost to follow-up (n=0)

Discontinued intervention (patient did not return) (n=3)

Lost to follow-up (n= 0)

Discontinued intervention (n=0)

Analysis

Primary outcome
Analysed (n=198)
- Excluded from analysis (n=82)
  Did not return questionnaire (n=77)
  Missing data (n=5)
Secondary outcome
Analysed (n=203)
- Excluded from analysis (did not return questionnaire) (n=77)
Tertiary outcome
Analysed (n=256)
- Excluded from analysis (missing data) (n=28)

Primary outcome
Analysed (n=289)
- Excluded from analysis (n=30)
  Did not return questionnaire (n=24)
  Missing data (n=6)
Secondary outcome
Analysed (n=299)
- Excluded from analysis (did not return questionnaire) (n=24)
Tertiary outcome
Analysed (n=310)
- Excluded from analysis (missing data) (n=9)