Comparison of two early treatment protocols for anterior dental crossbite in the mixed dentition: *A randomized trial*

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ABSTRACT

Objective: To evaluate and compare two treatment protocols to correct anterior dental crossbite in the mixed dentition.

Materials and Methods: Thirty children, 8–10 years of age, participated. Individuals were divided into two groups. Group 1 consisted of 15 children treated with an upper removable appliance with finger springs; group 2, 15 children treated by bonding resin-reinforced glass ionomer cement bite pads on the lower first molars. The 30 participants were evaluated before treatment (T1) and 12 months after treatment began (T2). The variables evaluated included overjet, perimeter of the maxillary arch, intercanine distances in the maxilla and mandible, SNA, SNB, ANB, and U1.NA. Data analysis included descriptive statistics, paired *t*-test and Student's *t*-test. Effect sizes and confidence intervals were also calculated.

Results: Group 1 showed a significant increase in overjet (P < .001), intercanine distance in the maxilla (P = .006), intercanine distance in the mandible (P = .031), and U1.NA (P = .002). Group 2 showed a significant increase in overjet (P = .008), intercanine distance in the mandible (P = .005), and U1.NA (P < .001). For all the evaluated variables, no statistically significant differences were observed between the two groups.

Conclusions: No significant differences were observed between the two protocols: use of a removable maxillary biteplate with finger springs and bonding of resin-reinforced glass ionomer cement bite pads on the lower first molars, for the correction of anterior crossbite in the mixed dentition. (*Angle Orthod.* 2018;88:144–150.)

KEY WORDS: Child; Malocclusion; Anterior crossbite; Mixed dentition; Interceptive orthodontics

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INTRODUCTION

Anterior crossbite is defined as a malocclusion in the sagittal dimension resulting in a lingual position of the maxillary anterior teeth in relation to the mandibular anterior teeth. It has great clinical significance that is both esthetic and functional.¹ It can be found in the deciduous, mixed, and permanent dentitions. Based on the individual presentation, an anterior crossbite can be dental, functional, or skeletal. In the literature, the prevalence of all types of anterior crossbite varies from 2.2% to 11.9%, depending on whether the edge-to-edge relationship of the incisors is considered and on the racial characteristics of the individuals evaluated.²

Anterior crossbite occurs because of a change in the vestibular-lingual relationship of one or more anterior teeth, with the maxillary incisor(s) lingually positioned and the mandibular teeth more facial. It has been reported that traumatic occlusion may be present and, if this problem does not receive early treatment, that it

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may result in periodontal problems in the mandibular incisors,³ the occurrence of pain, changes in the anterior-posterior positioning of the mandible, and development of problems in the temporomandibular joint (TMJ).^{4,5}

Interceptive orthodontic treatment is defined as any procedure that eliminates or reduces the severity of a developing malocclusion.² Such an intervention during the mixed dentition may allow the clinician to correct an anterior crossbite, thus favoring more harmonious growth of the bones^{6–8} and perhaps preventing the crossbite to persist in the permanent dentition. In this sense, when the orthodontist acts in an interceptive manner, comprehensive orthodontic treatment with fixed appliances may be simplified or reduced.⁹

A wide range of treatment protocols can be used to correct an anterior crossbite in the mixed dentition.² However, there is little evidence to indicate which treatment method is the most efficient.¹⁰ Therefore, the present study aimed to compare two of these protocols: an upper removable appliance with finger springs and the bonding of resin-reinforced glass ionomer cement bite pads on the lower first molars. The null hypothesis was that the early treatment of anterior crossbite with either of these two protocols is equally efficient.

MATERIALS AND METHODS

This study is reported according to the Consolidated Standards of Reporting Trials (CONSORT) guide-lines.¹¹

Participants, Study Location, and Eligibility Criteria

The sample consisted of 30 individuals 8-10 years of age who presented with anterior crossbite in the mixed dentition. The participants were divided into two groups. Group 1 consisted of 15 children who were treated with an upper removable appliance with finger springs. Group 2 consisted of 15 children treated by bonding resin-reinforced glass ionomer cement bite pads on the lower first molars. The distribution of the 30 individuals into the two groups was performed in a randomized manner as follows: a sealed envelope was prepared with 30 cards containing the names of the two treatment protocols on 15 cards each. For each participant, one card was drawn from the envelope to indicate to which group the patient would be assigned. This process was carried out by an assistant until all patients had been placed in a group. The 30 children were treated by one orthodontist.

The sample was selected from the medical records of patients receiving treatment at the Children's Clinic of the Federal University of the Valleys of Jequitinhonha and Mucuri (UFVJM), Diamantina, Brazil. The study included individuals from 8 to 10 years of age who presented with an anterior crossbite in the mixed dentition with all four permanent first molars erupted and at least one permanent incisor in crossbite. Exclusion criteria were (1) any compromised condition of the child's overall health (including craniofacial anomalies and cognitive disorders) according to the child's medical record and physical examination as reported by the parents, (2) children with functional crossbites, (3) individuals with skeletal anterior crossbites (ANB <0°) or a posterior crossbite associated with the anterior crossbite, (4) children with sucking habits or cessation of a sucking habit within less than 1 year, and (5) individuals with a prior history of orthodontic treatment.

Ethical Considerations

The study was approved by the UFVJM Human Research Ethics Committee (protocol 525.056). The children and their guardians were informed about the objectives of the study and that their participation was voluntary. For those agreeing to participate, the children and their guardians signed an informed consent form. After 12 months of follow-up, the children who did not exhibit a full correction of the anterior crossbite either continued in treatment or underwent a new type of treatment.

Sample Calculation

Considering an alpha significance level = 0.05 and a statistical power of 95%, the study required at least nine individuals in each group to detect an average difference of 2.0 mm (\pm 3.0) in overjet between the treatment protocols. To compensate for possible losses, six additional participants were included in each group. Therefore, there were 15 individuals assigned to each group (Figure 1).

Upper Removable Appliance with Finger Springs

The device had two Adams clasps on the permanent first molars, two arrow clasps between the deciduous molars, and a double finger spring adapted to the palatal surfaces of the teeth to be uncrossed, in addition to the labial bow. The posterior region included an occlusal splint in an attempt to obtain sufficient disocclusion to allow for moving the teeth in crossbite. The patients were advised to remove the appliance only to eat or during oral hygiene.

Resin-reinforced Glass Ionomer Cement Bite Pads

Resin-reinforced glass ionomer cement bite pads (Riva Light Cure, Bayswater, Australia) were placed on the occlusal surface of the mandibular permanent first



Figure 1. Flow chart of the study.

molars. These devices were thick enough to disclude all the anterior teeth, which allowed enough space for the movement of the teeth in crossbite by tongue pressure. Appointments were scheduled every 3–4 weeks for patients in both groups. Treatment for the 30 participants was conducted by an orthodontic specialist.

Evaluated Variables

The assessor of the outcomes was blinded. The 30 participants were evaluated before treatment (T1) and 12 months after implementation of the protocols (T2). The following outcomes were measured on study casts:

Overjet (Correction of the Crossbite)

The therapeutic effect of the two treatment protocols was evaluated, using a metal ruler, by measuring the overjet increase in millimeters, that is, the difference of overjet between T1 and T2.

Perimeter of the Maxillary Arch

Evaluation of the maxillary arch perimeter was performed with an initial and final plaster model using a brass wire, beginning at the distal surface of the deciduous second molar (or the mesial surface of the permanent first molar), passing around the arch over the contact points of the posterior teeth and incisal edges of the anterior teeth to the distal surface of the Number (%) Number (%)

Group 2

Table 1. Characteristics of the Children in Both Groups and

Group 1

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07 (46.7) 0.264* 11 (73.3) 04 (26.7) 08 (53.3) 05 (33.3) 08 (53.3) 0.462* 10 (66.7) 07 (46.7) 07 (46.7) 07 (46.7) 0.999* 08 (53.4) 08 (53.3) Mean (SD^a) Mean (SD) 9.07 (0.79) 9.00 (0.84) 0.826** Number of teeth crossed 1.60 (1.06) 1.67 (0.61) 0.834** ^a SD indicates standard deviation. Pearson chi-square test. deciduous second molar (or the mesial surface of the permanent first molar) on the opposite side.¹² The increase in arch perimeter was calculated by the difference between the perimeter of the arch in T1

Intergroup

Comparison

(P Value)

Intercanine Distance

Intergroup Comparisons

Full crossbite correction

Children's age (years)

** Student's t-test.

Gender

Boys

Girls

Crowding

No

Yes

No

Yes

and T2.

The intercanine distance in the maxilla and mandible was measured with a digital caliper (Digital 6. 8M007906, Mauser-Messzeug GmbH, Oberndorf/ Neckar, Germany) as the shortest linear distance between the canine cusp tips on the plaster models. Intercanine expansion was calculated by the difference between the intercanine distances at T1 and T2.13

Cephalometric Analysis

The cephalometric angles evaluated were SNA, SNB, and ANB, to evaluate the position of the maxilla and mandible relative to the cranial base and the

Table 2.	Comparison of	Pretreatment	Measures	Between	Groups
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	Group 1	Group 2	
	Mean (SD)	Mean (SD)	P Value*
Overjet	-1.13 (0.35)	-1.27 (0.45)	.379
Arch perimeter ^a	92.20 (6.61)	90.73 (5.24)	.506
Dist IC (Mx)	42.93 (2.12)	42.20 (3.52)	.496
Dist IC (Md)	36.80 (2.04)	35.07 (2.57)	.051
SNA	82.14 (3.74)	81.77 (4.41)	.805
SNB	78.34 (3.76)	78.42 (4.36)	.961
ANB	3.80 (1.74)	3.35 (2.75)	.600
U1.NA	20.60 (4.88)	19.73 (6.11)	.671

^a Indicates maxillary arch perimeter; Dist IC (Mx), intercanine distance in the maxilla; Dist IC (Md), intercanine distance in the mandible; SD, standard deviation.

Indicates Student's *t*-test. Significance level of P < .05.

position of the maxilla and mandible to each other. The upper incisor inclination (U1.NA) was also evaluated. The change in cephalometric angles was determined by the difference in the values between T1 and T2.

Statistical Analysis

Statistical analysis was conducted using the Statistical Package for the Social Sciences (SPSS for Windows, version 17.0, SPSS Inc, Chicago, III). Application of the Shapiro-Wilk test demonstrated that the data were normally distributed. Therefore, parametric tests were used. The analysis of the data included descriptive tests (chi-square and Student's ttest) to characterize the sample. Paired *t*-tests were used to evaluate the effects (changes occurring during treatment, T2-T1) of the two treatment protocols for correcting anterior crossbite. A Student's t-test was used to compare the changes occurring during treatment (T2-T1) between the two groups. Values of P < .05 were considered statistically significant. Effect sizes with 95% confidence intervals were also calculated by dividing the difference between the means of both groups by the pooled standard deviation.14,15 Effect sizes were interpreted according to the following values: 0.20, small; 0.50, medium; and 0.80, large.14

RESULTS

The average age of the children in group 1 was 9.07 years (± 0.79), while in group 2, it was 9.00 years (± 0.84) . Characteristics of the participants and intergroup comparisons are described in Table 1. Comparison of the pretreatment measures between groups is displayed in Table 2. Table 3 shows the effects (T2-T1) of the two treatment protocols for correcting the anterior crossbite. Group 1 showed a significant increase in overjet (P < .001), maxillary intercanine distance (P=.006), mandibular intercanine distance (P= .031), and upper incisor inclination (P = .002). Group 2 showed a significant increase in overjet (P = .008), mandibular intercanine distance (P = .005), and upper incisor inclination (P < .001). Table 4 compares the changes during treatment between the two protocols and the effect sizes. No statistically significant differences were observed between the two groups for any of the variables evaluated.

DISCUSSION

The orthodontic literature on early treatment protocols for anterior crossbite has been sparse. Recently, a systematic review suggested that clinical trials should be conducted to evaluate the efficiency of different treatment protocols for this type of malocclusion.² Taking into account the lack of statistically significant

	Group 1			Group 2		
	Mean (SD) T1	Mean (SD) T2	P Value*	Mean (SD) T1	Mean (SD) T2	P Value*
Overjet	-1.13 (0.35)	0.27 (0.88)	<.001	-1.27 (0.45)	-0.27 (0.96)	.008
Arch perimeter ^a	92.20 (6.61)	91.73 (6.60)	.250	90.73 (5.24)	90.27 (5.50)	.396
Dist IC (Mx)	42.93 (2.12)	44.33 (2.71)	.006	42.20 (3.52)	43.33 (1.98)	.084
Dist IC (Md)	36.80 (2.04)	38.40 (2.66)	.031	35.07 (2.57)	37.27 (2.37)	.005
SNA	82.14 (3.74)	83.20 (3.76)	.114	81.77 (4.41)	81.63 (4.61)	.794
SNB	78.34 (3.76)	79.12 (3.69)	.276	78.42 (4.36)	78.42 (4.45)	1.000
ANB	3.80 (1.74)	4.09 (2.48)	.589	3.35 (2.75)	3.21 (1.46)	.795
U1.NA	20.60 (4.88)	23.87 (4.67)	.002	19.73 (6.11)	23.60 (5.08)	<.001

Table 3. Effects of the Two Treatment Protocols in Correcting Anterior Crossbite

^a Indicates maxillary arch perimeter; Dist IC (Mx), intercanine distance in the maxilla; Dist IC (Md), intercanine distance in the mandible; SD, standard deviation; T1, before treatment; T2, 12 months after beginning treatment.

* Paired *t*-test. Significance level of P < .05.

differences between protocols, the null hypothesis could not be rejected in the present study. Anterior crossbite affecting one or more incisors was corrected efficiently by both an upper removable appliance with finger springs and bonded resin-reinforced glass ionomer cement bite pads on the lower first molars. Thus, both treatment protocols could be recommended for correcting this type of malocclusion. Consequently, this study offers relevant information to practitioners because early treatment of anterior crossbite has been widely advocated.^{16,17}

The findings reported in this study showed that both of the early protocols investigated led to a significant increase in overjet and mandibular intercanine distance after the 12-month treatment period. Moreover, improvement in overjet and intercanine distance in the mandible was not different between the upper removable appliance and the bite pads on the lower first molars. One prior study that compared the efficiency of fixed appliances and upper removable appliances with finger springs demonstrated that an anterior crossbite in the mixed dentition can also be corrected successfully using either of those two protocols.¹⁰ Additionally, long-term, posttreatment stability was similar in those two modes of treatment. For both the fixed and the removable appliance, the success rate remained high at the 2-year follow-up.¹⁸

One randomized clinical trial that evaluated the early correction of unilateral posterior crossbite revealed that the success rate was superior with a fixed device (Quad-helix) compared with treatment using a removable appliance with an expansion screw.¹⁹ The average treatment time was also significantly shorter and cheaper with the bonded device.²⁰ This finding may be attributed to low compliance with the removable device. It is well-known that when therapy with removable appliances is prescribed, patient compliance is a determining factor in the efficiency of treatment.²¹ The level of compliance with treatment can partly explain the prolonged treatment time observed with removable appliances. However, if the patients had cooperated, perhaps there would have been a more favorable outcome.²² It is likely that patients with anterior crossbite have greater awareness of their malocclusion since the condition is esthetically obvious in contrast to patients with posterior crossbite.¹⁸ Hence, the individuals from the current study should have been highly motivated and willing to comply with treatment recommendations.

Despite being a nonsignificant difference, the increase in overjet was marginally higher for the

 Table 4.
 Comparison of the Changes During Treatment (T2–T1) Between the Two Groups

	Group 1 T2–T1	Group 2 T2–T1				
	Mean (SD)	Mean (SD)	Difference Between Groups	P Value*	Effect Size**	CI (95%)
Overjet	1.40 (0.91)	1.00 (1.25)	0.40	.326	0.37	-0.33-1.07
Arch perimeter ^a	-0.47 (1.50)	-0.47 (2.06)	0.00	1.000	0.00	-0.70-0.70
Dist IC (Mx)	1.40 (1.68)	1.13 (2.35)	0.27	.724	0.13	-0.57-0.83
Dist IC (Md)	1.60 (2.58)	2.20 (2.56)	-0.60	.529	0.23	-0.47-0.93
SNA	0.74 (2.55)	-0.14 (2.03)	0.88	.306	0.38	-0.32-1.08
SNB	-3.30 (6.68)	-0.40 (1.59)	-2.90	.458	0.24	-0.46-0.94
ANB	0.29 (2.05)	-0.14 (2.04)	0.43	.568	0.21	-0.49-0.91
U1.NA	3.27 (3.41)	3.87 (3.14)	-0.60	.620	0.18	-0.52-0.88

^a indicates maxillary arch perimeter; Dist IC (Mx), intercanine distance in the maxilla; Dist IC (Md), intercanine distance in the mandible; T1, before treatment; T2, 12 months after beginning treatment; SD, standard deviation; CI, confidence interval.

* Student's *t*-test. Significance level of P < .05.

** Difference between means of both groups by the pooled standard deviation.

individuals treated with the upper removable fingerspring appliance compared with those treated with the resin-reinforced glass ionomer cement bite pads. This effect size was 0.37 mm. Indeed, taking into account the analysis of the effect size, the benefit of a particular protocol may be suggested by a small trial such as this one with nonstatistically significant results. It has been advocated that statistical outcomes give relevant information but, sometimes the statistical significance might not reflect the size of the treatment effect.²³

The present study had weaknesses that should be acknowledged. This study should have included a control group of individuals with untreated anterior crossbite of the mixed dentition.18 However, this would have been unacceptable for ethical reasons²⁴ and, also, the use of historical control groups has been a subject of much criticism in orthodontic research.25 Second, as previously mentioned, in any removable appliance therapy, patient compliance with treatment is a significant determinant of treatment efficiency. Therefore, one could argue that the present clinical trial may have been sensitive to the risk of the Hawthorne effect, through which the awareness of being evaluated could have had a positive impact on childrens' behavior.²⁶ Consequently, they may have cooperated better with the prescribed treatment regimen. However, this trial was strengthened by the random allocation of participants and the prospective longitudinal design. The former helped to minimize bias in the assignment of individuals to each treatment protocol, resulting in two groups that were comparable for known or unknown confounding variables.²⁷ The latter allowed investigation of causal associations between the interventions and the outcome.28

The variables included in this study were highly relevant measures for evaluating treatment effectiveness.^{29,30} Nevertheless, it is important to note that the literature has also advocated that other important aspects of early intervention should be evaluated in mixed dentition treatment. These include cost-benefit31 and complications during treatment (displacement, breakage, and loss of appliances),²⁰ in addition to other variables, such as the perception of pain and discomfort associated with treatment.32 Those outcomes should be analyzed in future studies. Patientreported measures, such as quality of life assessments have been underrepresented in orthodontic clinical trials. Thus, future research should evaluate individuals' perceptions of the physical and psychological consequences of such orthodontic protocols.33 Moreover, as the early treatment of anterior crossbite is performed in individuals who are still growing, it is also important to evaluate the stability after correction and the changes observed from a long-term perspective. $^{\scriptscriptstyle 10}$

CONCLUSIONS

 No significant differences were observed between the two protocols: use of a removable maxillary biteplate with finger springs and bonding of resinreinforced glass ionomer cement bite pads on the lower first molars, for correcting anterior crossbite in the mixed dentition.

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