

Acoustic rhinometric evaluation of the nasal cavity after rapid maxillary expansion

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ABSTRACT

Objective: Because of the anatomic proximity between the nasal cavity and the maxilla, much has been investigated regarding changes in nasal geometry after this procedure. In this study, we propose to evaluate the repercussion of RME in the nasal cavity in the patient during the growth phase.

Materials and methods: For this, we evaluated 19 patients with transverse maxillary deficiency and indication for RME. The patients were evaluated using acoustic rhinometry in 3 moments (pre-RME, post-RME, post-restraint).

Results: There was no mean change in MCA1 M1, M2 and M3 ($p = 0.122$). MCA2 measurement appears to increase in M2. VOL 1 is suffering a mean increase in M2 compared to M1 ($p = 0.025$) and continues higher in M3 ($p = 0.271$). There is little variation of VOL 2 between the evaluated moments.

Conclusion: The results allow us to affirm that RME significantly increases the anterior region of the nasal cavity immediately to the procedure, however, after the period of containment there is a tendency of recurrence of this increase returning to values close to the initial cross-sectional area of the nasal cavity.

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INTRODUCTION

The nasal breath is the only one considered normal and physiological because it is present at birth. However, there may be transference to the predominantly oral respiratory pattern when there is nasal obstruction, which may lead to alterations in the development of dentofacial structures.¹

Many authors¹⁻⁴, consider that oral breathing is one of the etiological factors of transverse maxillary deficiency. However, this cause-effect relationship continues to be much discussed due to controversies in the literature.^{5,6}

Rapid maxillary expansion (RME) is an excellent method for the correction of transverse maxillary deficiency through the opening of the palatine suture.⁷⁻¹¹ Due to the anatomical proximity between the nasal cavity and the maxilla, much has been investigated regarding the changes in nasal geometry after this procedure (RME). Some authors^{9,12-14} have demonstrated

that the benefits of this procedure in patients during the growth phase go beyond the correction of malocclusion and may have repercussions in the nasal cavity.

Some researchers^{11,12,14,15-17} have associated RME with a reduction in nasal respiratory resistance, increased airflow, increased nasal cavity size and a change in respiratory mode from oral to nasal. However, other studies did not observe significant changes in the nasal cavity after RME^{18,19}, so this subject is not yet fully clarified in the literature, thus requiring additional studies.

Acoustic rhinometry was introduced by Hilberg et al.²⁰ and has been described as an accurate and reproducible method of evaluation of the anterior nasal cavity, being considered an objective, easily performed, non-invasive and well-tolerated examination by adults or children.^{21,22} When compared with other methods of evaluation such as computed tomography, rhinomanometry and magnetic resonance, the results obtained with acoustic rhinometry are equivalent in determining the nasal geometry of the anterior portion of the nasal cavity.^{23,24,25}

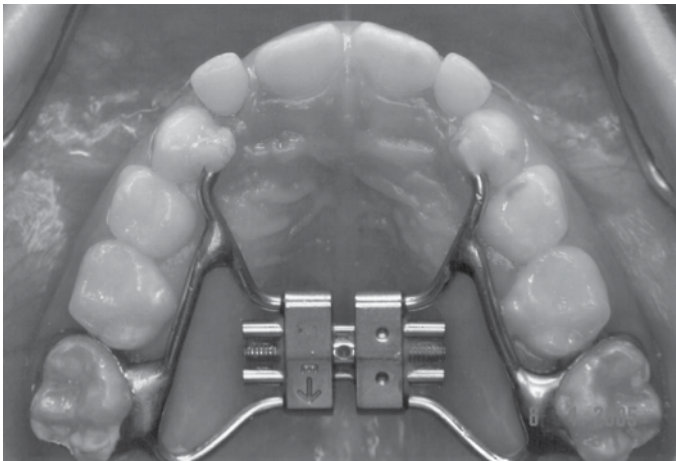
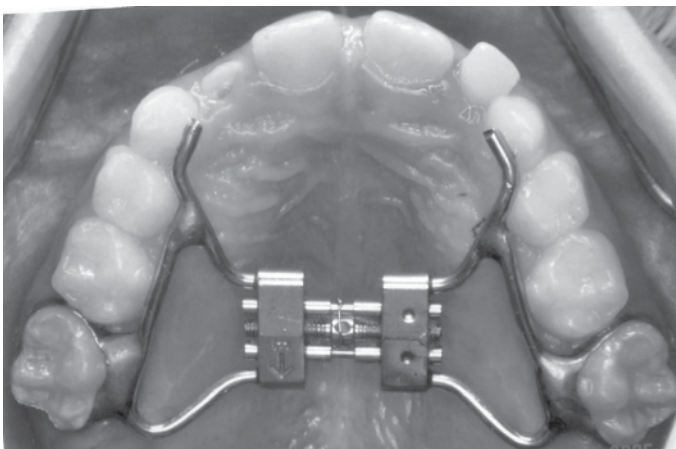
In this study we proposed to evaluate the alterations in the nasal cavity, using acoustic rhinometry, of patients with transverse maxillary deficiency before, immediately after and 6 months after Rapid Maxillary Expansion (RME) treatment.

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Table 1. Distribution of age and gender.

Patient	Age	Gender
1. BCF	7	F
2. CVSL	9.5	M
3. DHB	8.6	M
4. FCL	7.5	F
5. GSP	9.4	M
6. LMP	7.6	F
7. LRA	12.5	M
8. MPFM	9.3	M
9. MFS	9	M
10. MCS	7	F
11. PMG	8	F
12. PSO	8.4	F
13. PCS	8.3	M
14. PPS	8.6	F
15. VQC	12.6	M
16. VAP	8.4	M
17. WBC	7.5	M
18. APA	11	F
19. CCV	9.6	F

*Figure 1. Biederman Modified appliance installed before RME.**Figure 2. After RME with correction of the maxillary deficiency.*

MATERIALS AND METHODS

This study was approved by the Research Ethics Committee of the Faculty of Dentistry of the University of São Paulo, under protocol 21/01.

19 patients were selected (9 females - 47.3% and 10 males - 52.7%), with a mean age of 8.9 years (Table 1). All the patients presented transversal maxillary deficiency and, based on the data obtained in the diagnosis, were submitted to rapid maxillary expansion in the Preventive Orthodontics Clinic of the Faculty of Dentistry of the University of São Paulo, from January to December 2009. As inclusion criteria, patients could not have undergone previous orthodontic or otorhinolaryngologic treatment. For ethical reasons, it was not possible to evaluate a control group, for they would be subjected to unnecessary radiation, since they would not be treated orthodontically. In this way, we use data already published in the literature for comparison purposes with the selected sample.

The rapid maxillary expansion was performed using a modified Biederman dento-supported expander type. After a week of installation of the device, the activations were initiated, with one complete lap performed, and the patient responsible was instructed to activate $\frac{1}{4}$ turn every 12 hours. The activation time varied according to the need of expansion for correction of transverse maxillary deficiency (Figure 1 and 2).

All patients selected were submitted to a three-stage acoustic rhinometry test: before RME (M1), one week after the end of the activations (M2) and six months after the end of the activations (M3). The rhinometric examination was performed in each individual after it was acclimatized to the ambient conditions of the examination room, using RhinoScan apparatus from RhinoMetrics A/S - Denmark.

The procedure followed the determinations of the Standardization Committee of Acoustic Rhinometry.²⁰ In each of the times, three measurements were performed in each nostril before and after the use of topical vasoconstrictor (0.5 mg/ml - Pedrin Afrin) oxymetazoline hydrochloride. The measurements followed the following protocol: obtaining the measurements in the right nasal cavity and then in the left nasal cavity, both without the use of the vasoconstrictor. Thereafter, the application of three drops of topical vasoconstrictor was followed for a period of 10 to 15 minutes and the measurements were repeated following the same sequence.

Two area measurements were evaluated in each nostril (left and right): minimum cross-sectional area between 0mm and 22 mm (MCA1), and minimum cross-sectional area between 22 mm and 54 mm (MCA2). Similarly, two volume measurements were evaluated on each side: Nasal space volume between 0 mm and 22 mm (VOL1) and Nasal space volume between 22 mm and 54 mm (VOL2). The results were reported in cm² and cm³, respectively. The data were automatically calculated by the RhinoScan® software and were considered averages of the 3 measurements (Figure 3).

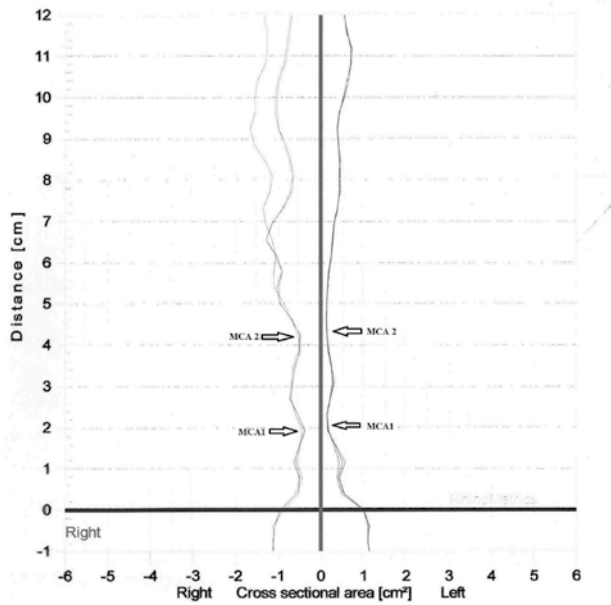


Figure 3. Graph automatically generated by RhinoScan® software with three curves obtained on each side. The central line representing the nasal septum and the curves representing the lateral walls of the nasal cavity on the right and left sides. From this graph the program provides the measures of MCA1 and MCA2 and volume 1 and 2.

RESULTS

In order to respond to the objectives of the study, first the measurements of nasal cavity area were recorded according to the use of vasoconstrictor, nasal cavity side and moment of the examination, using summary measures (mean, standard deviation). The results obtained were illustrated with the use of graphs of means with the respective standard errors. Subsequently, repeated measures (ANOVA) were performed with three factors, vasoconstrictor, nasal cavity side and time of the examination, followed by multiple Tukey comparisons to verify the level of significance between the mean measures found.

The tests were performed at a significance level of 5%.

Table 2 shows that the MCA1 measurements do not vary between the right and left sides, with or without vasoconstrictor, but there seems to be a change between the moments of evaluation. The MCA2 measurement is lower without vasoconstrictor use and appears to increase in M2.

Table 3 shows that there was no mean change in MCA1 between the right and left sides (p=0.299), between M1, M2 and M3 (p=0.122) or with vasoconstrictor use (p=0.510). But, for mean MCA2 there was statistically change between the moments (p=0.043) and with vasoconstrictor use (p=0.008) independent of the side.

Table 4 shows that the use of the vasoconstrictor increases the mean MCA2 (p=0.009) and the mean value of MCA2 is statistically higher in M2 when compared to M1 (p=0.045).

Table 2. Description of MCA1 and MCA2 (cm²) according to use of vasoconstrictor, right and left sides and moments M1, M2 and M3.

Vasoconstrictor	Slide	Moment	MCA1		MCA2	
			Average	SD	Average	SD
With vasoconstrictor	Right	M1	0.31	0.11	0.41	0.17
		M2	0.36	0.10	0.50	0.15
		M3	0.34	0.11	0.44	0.16
	Left	M1	0.38	0.47	0.43	0.16
		M2	0.30	0.11	0.47	0.18
		M3	0.25	0.10	0.38	0.17
No vasoconstrictor	Right	M1	0.30	0.12	0.36	0.15
		M2	0.35	0.09	0.43	0.14
		M3	0.31	0.13	0.42	0.29
	Left	M1	0.25	0.11	0.32	0.14
		M2	0.40	0.47	0.38	0.13
		M3	0.24	0.10	0.35	0.24

Table 3. Results of ANOVA for MCA1 and MCA2 measurement. (* statistically significant).

Factor	fd num.	fd den.	MCA1		MCA2	
			Value F	P	Value F	P
Vaso constrictor	1	18	0.45	0.510	8.75	0.008*
Side	1	18	1.14	0.299	3.01	0.100
Moment	2	36	2.23	0.122	3.44	0.043*
Vaso*Side	1	18	0.01	0.935	0.39	0.538
Vaso*Moment	2	36	1.70	0.198	0.63	0.540
Side*Moment	2	36	1.12	0.337	0.74	0.482
Vaso*Side *Moment	2	36	1.68	0.201	0.14	0.869

Table 4. Results of Tukey's multiple comparisons between moments M1, M2 and M3 and use of vasoconstrictor in MCA2. (* statistically significant) WV – With vasoconstrictor / NV – No vasoconstrictor.

Factor	Comparison	Estimate	Standard error	Value T	fd	p
Vaso constrictor	WV-NV	0.063	0.022	2.91	18	0.009*
	M1-M2	-0.066	0.026	-2.49	36	0.045*
Moment	M1-M3	-0.017	0.026	-0.64	36	0.798
	M2-M3	0.049	0.026	1.84	36	0.170

Table 5 suggests that the measure of VOL 1 is smaller on the left side than on the right side. The VOL 2 is lower without vasoconstrictor use and there is little variation of this measure between the evaluated moments.

Table 5. Description of VOL1 and VOL2 according to the use of vasoconstrictor, right and left sides and moments M1, M2 and M3.

Vasoconstrictor	Slide	Moment	VOL1		VOL2	
			Average	SD	Average	SD
With vasoconstrictor	Right	M1	1.15	0.18	3.14	3.14
		M2	1.23	0.19	3.18	3.18
		M3	1.22	0.20	3.43	3.43
	Left	M1	1.06	0.15	3.27	3.27
		M2	1.17	0.34	3.37	3.37
		M3	1.01	0.29	2.87	2.87
No vasoconstrictor	Right	M1	1.13	0.28	2.43	2.43
		M2	1.22	0.17	2.69	2.69
		M3	1.23	0.20	2.36	2.36
	Left	M1	1.07	0.23	2.32	2.32
		M2	1.10	0.25	2.70	2.70
		M3	1.09	0.25	2.15	2.15

Table 6 shows that VOL 1 varies between the right and left sides ($p < 0.001$) and between moments M1, M2 and M3 ($p = 0.028$), regardless of whether or not vasoconstrictor is used. For VOL 2 we observe a statistically significant increase only for the use of the vasoconstrictor ($p < 0.001$) independent of the side or the moment of evaluation and, therefore, Tukey's multiple comparisons test was not performed for this measure.

Table 6. Result of ANOVA for measurement of VOL1 and VOL2. (* statistically significant).

Factor	fd num.	fd den.	VOL1		VOL2	
			Value F	P	Value F	P
Vasoconstrictor	1	18	0.00	0.963	44.56	<0.001*
Side	1	18	25.66	<0.001*	0.63	0.439
Moment	2	36	3.94	0.028*	2.06	0.142
Vaso*Side	1	18	0.21	0.652	0.01	0.931
Vaso*Moment	2	36	1.24	0.302	0.71	0.499
Side*Moment	2	36	1.80	0.180	1.68	0.200
Vaso*Side*Moment	2	36	0.76	0.475	0.65	0.527

In Table 7 it can be seen that the mean VOL1 is statistically higher on the right side ($p < 0.001$) and suffers a mean increase in M2 compared to M1 ($p = 0.025$) and continues higher in M3 ($p = 0.271$).

Table 7. Results of Tukey multiple comparisons between right and left sides and moments M1, M2 and M3 for the measurement of VOL 1.

(* statistically significant).

Factor	Comparison	Estimate	Standard error	Value T	fd	P
Side	Right-Left	0.114	0.023	4.97	18	<0.001*
	M1-M2	-0.077	0.028	-2.75	36	0.025*
Moment	M1-M3	-0.033	0.028	-1.17	36	0.476
	M2-M3	0.044	0.028	1.57	36	0.271

DISCUSSION

According to the literature, according to the literature, many studies attempt to evaluate the relationship between rapid maxillary expansion and increased nasal cavity size by different methods.²⁶ The purpose of this study was to evaluate the nasal cavity before and after RME using acoustic rhinometry, which provides Minimal crosssection areas and nasal volumes. When analyzing the results, we can observe in Table 2 the variation of the mean values of MCA1 and MCA2 between moments M1, M2 and M3, with their respective standard errors of the right and left sides. In MCA1 for the left side, without the use of vasoconstrictor, the MCA1 ranged from 0.25 cm² in M1 to 0.40 cm² in M2, but there was a recurrence in M3 to 0.24 cm². For the right side, without the use of vasoconstrictor we observe a variation of the same form, being that, in M1, the average value of MCA1 was 0,30 cm², in M2 0,35 cm² and in M3 0,31 cm². However, this observed increase was not statistically significant, possibly by sample size (Table 3).

Similar results were found by Enoki et al.¹⁸ who investigated ²⁹ patients undergoing RME and observed a significant increase of MCA1 immediately after RME. The authors found that in M1 (Pre-RME) the mean MCA1 of the left and right nasal cavities was 0.987 cm². After RME, this measure increased to 1.006 cm² and after the use of restraint, these authors also observed a recurrence of this measure to 0.973 cm².

Bicakci et al.²⁷ reported an increase of 8.7% in MCA1 in a group of patients treated with RME between M1 (Pre-RME) and M3 (Post-retention). The authors also observed that there was a recurrence of 6.3% between M2 (Post-RME) and M3.

As shown in Table 3, in this study no difference in behavior was observed between the right and left sides of the nasal cavity, similarly to the study of Cappellette et al.¹², however, these authors observed a statistically significant increase in MCA1 immediately after RME.

De Fellippe et al.¹⁶ observed increased values in MCA1 after RME. In the pre-RME phase (M1) in the left nasal cavity, they found MCA1 of 0.38 cm², in the post-RME phase (M2) increased to 0.48 cm², three months after RME (M3) the measurement was 0.47 cm² and nine months after RME (M4) was 0.51 cm². For the right nasal cavity, the behavior was similar, with values of MCA1 in M=0.41 cm², M2=0.48 cm², M3=0.48 cm² and M4=0.53 cm². Statistical significance was found between M1 and M2, M1 and M3 and M1 and M4, but between the moments M2 and M3 there was no change with statistical significance that, according to the authors, represented a moment of stability after removing the appliance.

Tables 2 and 3 show the MCA2 values and it is noted that there was a change in the measurements between the evaluated moments. For the left nasal cavity in M1 we found 0.36 cm², in M2 0,43 cm² and in M3 0,42 cm². For the right nasal cavity the value found in M1 was 0.32 cm², in M2 0,38 cm² and in M3 0,35 cm². After the statistical analysis (Table 3) we noticed that the observed increase was significant in M2 when compared to M1, however, between M2 and M3 no significant change was found, which can be analyzed as a moment of stability as stated by De Fellippe et al.¹⁶

The results found by Enoki et al.¹⁸ show variation of the MCA2 values of the right and left nasal cavities. In M1 (Pre-RME) the value found was 0.732 cm², in M2 (Post-RME) 0.780 cm² and in M3 (3 months post-retainer) 0.763 cm². However, none of the changes presented statistical significance. Cappellette et al.¹² reported a statistically significant increase in MCA2 immediately after RME.

Our results show that when MCA2 means between M1 e M3 is compared, there is no statistically significant difference. Showing that after the period of containment there is a tendency of relapse to the initial measurements.

Regarding the nasal volume, it can be observed in Tables 5, 6 and 7 that the mean VOL1 increased between M1 and M2 (p=0.025), values statistically significant for the right nasal cavity. The mean variation was 1.13 cm³ in M1, to 1.22 cm³ in M2, remaining stable in M3 (1.23 cm³). In the left nasal cavity there was no significant variation in the moments of evaluation. Already, Cappellette et al.¹² reported an increase in VOL 1, after RME, on both sides of the nasal cavity. The mean increase was 0.07 cm³ for both right and left the nasal cavity.

De Fellippe et al.¹⁵ also observed increased nasal cavity volume after RME. The increase between M1-M2, M1-M3 and M1-M4 moments were significant, but no significant change was observed between M2-M3 and M3-M4. These results, in the opinion of the authors, demonstrate the stability of the volume increase achieved. Likewise, Babacan et al.²⁸ observed

a statistically significant increase of 14.48% in the nasal volume between M1 (Pre-RME) and M2 (post-RME).

For the measurements of nasal volume 2, we observed that there was an increase of 0.26 cm³ in the right nasal cavity between moments M1 and M2, without statistical significance. On the left side, an increase of 0.38 cm³ was observed between the moments M1 and M2, which were not statistically significant (Tables 5 and 6). Similarly, Cappellette et al.¹² reported an increase, between M1 and M2, of 0.34 cm³, statistically significant (p=0.023), from VOL 2 to the right nasal cavity, and on the left side the increase between these moments of evaluation was 0.21 cm³, not statistically significant.

CONCLUSION

The results found in this study allow us to affirm that RME increases significantly the anterior region of the nasal cavity immediately to the procedure, however, after the period of containment there is a tendency of recurrence of this increase returning to values close to the initial cross-sectional area of the nasal cavity. Therefore, the indication of this procedure should be performed only for correction of dentoalveolar problems and no expectation of correction of respiratory function should be generated.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

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