

Modified Dental Crowding Treatment Protocol during Mixed Dentition

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Submitted: 09-01-2023	Accepted: 19-01-2023

Annotation.Dentalcrowdinginmixeddentitionisone of the most common or tho dontic pathologies now adays

. According to modern literature database it's prevalence reaches 77% and present at all occlusion periods. which is a significant sign of malocclusion's severity. The aim of our research is to develop rational treatment protocol of patients with dental crowding during mixed dentition period according to facial skeleton growth patterns and also to make comparative analysis of treatment efficiency by using traditional and suggested protocol. Patients in mixed dentition period with dental crowding (n=164) were examined over a period of last three years at the base of NMU O.O. Bogomolets Dental Center, Kyiv. A total of 328 CBCT sections of facial skeleton (medium FOV) before and after treatment were submitted to the general analysis. The results and conclusions of treatment efficiency analysis were carried out by using the suggested protocol of dental crowding treatment with different facial skeleton growth patterns. Results indicated an improvement in treatment efficiency of this pathology and significant treatment time shortening. The results obtained after 16 months in patients with a horizontal growth pattern indicate that treatment efficiency value of clinical group (CG) II patients was 58.1±1.3%; after 17 months in patients with vertical and neutral growth patterns, treatment efficiency value of CG III was 66.7±1.6%, CG I was 52.3±0.9%. The algorithm proposed by us to shorten treatment duration by 3-4 allows months.

Keywords:<u>malocclusion</u>, <u>mixeddentition</u>, growthpattern, facial skeleton, palatalexpansiontechnique.

I. INTRODUCTION.

Modern orthodontic literature database indicates a consistently high frequency of malocclusions and dentognathic deformities that appear in children and adolescents. A huge increase in their prevalence is observed in children during the mixed dentition stage, which reaches 80%.[1-3]. Moreover, the most common are class I malocclusions, which according to various authors range from 50.6% to 84.4%[1-3]. It is also scientifically proven that with age no selfregulation of dental crowding is observed and in 80-90% of all cases it's likely to be observed during the permanent dentition period[5,7]. Determination of facial skeleton growth pattern is of significant practical importance, as it allows to make the most optimal choice for treatment start, to choose correct treatment method, to predict treatment's duration and consequences[5,9]. Dental crowding is one of the most common issues of orthodontics nowadays. According to worldwide literatureit's prevalence reaches 77% [2,3,8] and present at all occlusion periods, which is a significant sign of malocclusion's severity. Literature describes many methods of dental crowding treatment during mixed dentition period, which is caused by both maxillary and mandibular constriction. The most modern is usage of Rapid Maxillary Expansion protocol (RME) with Marco-Rosa appliance [10,11]. While the advantages of this aplliance are well known and scientifically proven, this appliance doesn't allow to directly expand maxillary frontal area and to create enough amount of space for anterior dental crowding regulation. That's why a new appliance for dental crowding treatment during mixed dentition was suggested by us. It's not only transversally expanding constricted maxilla a but also equally expands maxillary frontal area (patent of Ukraine № 149170, 21.10.2021).

Aim of research. To develop rational treatment protocol of patients with dental crowding during mixed dentition period according to facial skeleton growth patterns and also to make comparative analysis of treatment efficiency based on conebeamed computed tomography (CBCT) data by using traditional and suggested treatment protocol.

Materials and methods.For three years (2020-2022), we examined and treated patients with dental crowding at the Dental Medical Center of the Bogomolets National Medical University named. The research was carried out in compliance with the main provisions of the "Rules of Ethical Principles of Scientific Medical Research with Human Participation" approved by the Helsinki



Declaration (1964-2013), ICH GCP (1996), EU Directive No. 609 (from November 24, 1986). orders of the Ministry of Health of Ukraine No. 690 dated September 23, 2009, No. 944 dated December 14, 2009, No. 616 dated August 3, 2012.All participants were informed about the purpose and methods of the study and signed an informed consent to participate in it, and all measures were taken to ensure patient anonymity. The criteria for randomization of patients were next: mixed dentition period (7-11 years), the presence of dental crowding in maxillary and/or mandibular frontal area, erupted first permanent molars, the absence of general somatic diseases. Research included 164 people, 64 (39.1%) patients were male, and 100 (60.9%) patients were female.The distribution of examined patients according to the facial skeleton growth pattern is shown in Table 1. According to the algorithm developed by us, all patients who entered the examination groups were subjected to diagnostics before and after treatment. A total of 328 CBCT slices of the facial skull (medium FOV) of the patients at the beginning and after the treatment were analyzed.On CBCT sliceswe evaluated changes in width of both maxilla and mandible at basal arches (in the projection of the first permanent molars between the most convex points of the cortical plate, departing from the enamelcement junction by 8 mm in the direction of the apex of the root) and alveolar arches (in the projection of the first permanent molars between the most convex points of the alveolar process, receding from the enamel-cement junction by 3 mm in the direction of the apex of the root) levels before and after treatment, and changes of dental crowding severity were also evaluated according to the Little's IrregularityIndex values.

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Growth	Group of o	control, n=	Clinical	groupI,	Clinical	group II,	Clinical	group III,
pattern	20		n=44		n=48		n=52	
	male	female	male	female	male	female	male	female
horizontal	2	3	-	-	18	30	-	-
neutral	6	5	12	32	-	-	-	-
vertical	3	1	-	-	-	-	21	31

Table 1. Distribution of patients according to facial skeleton growth pattern and gender.

The generally accepted algorithm for dental crowding treatment is applying RME protocol has 2 phases, consisting fixation of Marco-Rosa appliance on maxilla and the activation of a 10 mm screw, once every 2 days at 90°, the active phase of screw activation is 64 days (2 months), after the end of the active phase this appliance remains in the oral cavity for 6 months as a retention . After the RME protocol, if necessary, a myofunctional trainer is additionally prescribed for 12 months with a wearing regime of 12 hours per day.

All patients had clinically significant dental crowding and were distributed into the clinical groups according to their facial skeleton growth patterns.

- 1. First clinical group -44 patients(30,5%)
- 2. Second clinical group -48 patients(33,3%)
- 3. Third clinical group -52 patients(36,2%)

The first clinical group consisted of patients with a neutral type of growth of the bones of the facial skull, the second clinical group included patients with a vertical type of growth of the bones of the facial skull, and the third clinical group included patients with a horizontal type of growth of the bones of the facial skull. The patients of each clinical group were treated according to our proposed algorithms: the first phase of orthodontic treatment consisted of suggested appliance fixationon maxillary dental arch with existing beams adjacent to the lateral group of teeth and protracting archesin the frontal area, together with a fixation of Williams fixed mandibular expander; the second phase consisted in prescription of myofunctional appliance depending on the presented malocclusion.

In the 1st clinical group, 30 people were treated according to the algorithm proposed by us: the appliance's screw proposed by us is activated once a day, the active phase is 32 days (1 month). fixed mandibular expanderby Williams is installed 2 weeks after the start of treatment and is activated once per 3 days, active phase - 1.5 months. Both devices remain in the oral cavity for a retention period of 6 months. After immediate removal of the devices, a myofunctional trainer is prescribed for 10 months: the mode of use is 14 hours a day, every day. The effectiveness of treatment was 52.3±0.9%, total duration of treatment was 17 months; 14 people were treated according to the standard algorithm: the screw of Marco-Rosa appliance is activated once per 2 days, the duration of the active phase of treatment is 64 days (2



months), the retention period is 6 months. After 3 weeks from the start of treatment, an installation of fixed mandibular expander byWilliams with an activation scheme once per 4 days, the duration of the active phase of treatment is 1.5 months, and the retention period is 5 months. After immediate removal of the devices, a myofunctional trainer is prescribed for 12 months. Mode of use: 12 hours a day. The effectiveness of treatment was $25.6 \pm 1.7\%$, the total duration of treatment was 20 months. Our proposed algorithm is more effective than the standard one in clinical group I patients by $26.9 \pm 1.2\%$.

In the II clinical group, 36 people were treated according to the algorithm proposed by us: the appliance's screw proposed by us is activated once a day, the active phase is 32 days (1 month), fixed mandibular expander by Williams is installed 2 weeks after the start of treatment and is activated once per 3 days, active phase - 1.5 months. Both devices remain in the oral cavity for a retention period of 6 months. After immediate removal of the devices, a myofunctional trainer is prescribed for 9 months: the mode of use is 15 hours per day, every day. The effectiveness of the treatment was 58.1±1.7%, total duration of treatment was 16 months; 12 people were treated according to the standard algorithm: the screw of Marco-Rosa appliance is activated once per 2 days, the duration of the active phase of treatment is 64 days (2 months), the retention period is 6 months. After 3 weeks from the start of treatment, an installation of fixed mandibular expander by Williams with an activation scheme once per 4 days, the duration of the active phase of treatment is 1.5 months, and the retention period is 5 months. After immediate removal of the devices, a myofunctional trainer is prescribed for 12 months. Mode of use: 12 hours per day. The effectiveness of treatment was $22.3\pm2.1\%$, the total duration of treatment was 20 months. Our proposed algorithm is $35.8 \pm 0.9\%$ more effective than the standard one in clinical group II patients.

In the III clinical group, 30 people were treated according to the algorithm proposed by us: the appliance's screw proposed by us is activated once a day, the active phase is 32 days (1 month), fixed mandibular expander by Williams is installed 2 weeks after the start of treatment and is activated once per 3 days, active phase - 1.5 months. Both devices remain in the oral cavity for a retention period of 6 months. After immediate removal of the devices, a myofunctional trainer is prescribed for10 months: the mode of use is 14 hours per day, every day. The effectiveness of the treatment was $66.7\pm1.6\%$, the total duration of treatment was 17 months; 16 people were treated according to the standard algorithm: the screw of Marco-Rosa appliance is activated once per 2 days, the duration of the active phase of treatment is 64 days (2 months), the retention period is 6 months. After 3 weeks from the start of treatment, an installation of fixed mandibular expander by Williams with an activation scheme once per 4 days, the duration of the active phase of treatment is 1.5 months, and the retention period is 5 months. After immediate removal of the devices, a myofunctional trainer is prescribed for 12 months. Mode of use: 12 hours a day. The effectiveness of treatment was $29.5\pm2.4\%$. total duration of treatment was 20 months. Our proposed algorithm is 37.3 ±0.7% more effective than the standard one in patients of the III clinical group. The control group consisted of 20 people.

The data we received were analyzed, interpreted and statistically processed. Statistical processing of these data included a number of parametric and non-parametric criteria of statistical methods. The analysis was performed using statistical packages MedStat and EZR v. 1.35 (Saitama Medical Center, Jichi Medical University, Saitama, Japan 2017). Statistical analysis of materials. summarization of results, and generalization of conclusions were performed using the method of variational statistics, taking into account average values (mode, median, arithmetic mean) and average error (M) with evaluation of reliable values according to the Wilcoxon t-test, as well as with the determination of the correlation coefficient using the paired Pearson's method to identify relationships between the obtained indicators. A value of p<0.05 was taken as the minimum probability threshold.

To compare these indicators obtained before and after treatment, appropriate comparison criteria for related samples were used. Comparison of qualitative features was carried out using the Chi-square test. If more than two clinical groups were compared, univariate analysis of variance (if a normal distribution) or Kruskel–Wallis test (if a non-normal distribution) was used for quantitative indicators. During the statistical analysis, criteria with a two-sided critical area were used, with the critical level of significance being p=0.05.

II. RESULTS.

According to research results (**table 2**), it was established that when using the proposed protocol in CG 1, the skeletal effect of the maxillary expansion (BAMxW) is 4.8 ± 1.1 mm, the alveolar effect of the expansion of maxilla (AAMxW) is 4.9 ± 0.8 mm., expansion of mandible at the basal level (BAMdW) was 3.0 ± 0.7 mm,



while at the alveolar level (AAMdW) 5.9 ± 1.2 mm was reached; at the same time, we managed to reduce the Little's Irregularity Index value of upper teeth (LIIMx) by 12.2 ± 1.5 mm, the Little's Irregularity Index value of lower teeth(LIIMd)by 9.3 ± 0.8 mm, i.e., we were able to transfer the severity of crowding from severe to mild on both maxilla and mandible. The effectiveness of the treatment was $52.3\pm0.9\%$.

When using the proposed protocol in CG 2, the skeletal effect of the maxillary expansion (BAMxW) is4,8±0,6 mm, the alveolar effect of the expansion of maxilla (AAMxW) – 4,2±0,6 mm, expansion of mandibleat the basal level (BAMdW) was 3,2 ±0,4 mm, while at the alveolar level (AAMdW) –4,1 ± 0,7 mm; at the same time, we managed to reduce the Little's Irregularity Index value of upper teeth (LIIMx) by 13,1 ± 1,2 mm, the Little's Irregularity Index value of lower teeth(LIIMd)by $6,9\pm1,4$ mm, i.e., we were able to transfer the severity of crowding from severe to mild on both maxilla and mandible. The effectiveness of the treatment was $58,1\pm1,7\%$.

When using the proposed protocol in CG 3, the skeletal effect of the maxillaryexpansion (BAMxW) is $6,3\pm0,7$ mm, the alveolar effect of the expansion of maxilla(AAMxW) – $5,2\pm0,9$ mm,, expansion ofmandibleat the basal level (BAMdW) was $3,6\pm0,8$ mm, while at the alveolar level (AAMdW) – $4,7\pm1,1$ mm; at the same time, we managed to reduce the Little's Irregularity Index value of upper teeth (LIIMx) by $11,9\pm1,7$ mm, the Little's Irregularity Index value of lower teeth(LIIMd)by $6,9\pm1,4$ mm, i.e., we were able to transfer the severity of maxillary crowding from severe to moderate, from severe to mild on mandible. The effectiveness of the treatment was $66,7\pm1,6\%$.

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CBCT data	CGI						
	Before	After	Treatment efficiency	p-value			
	treatment	treatment					
BAMxW	58,7 ±1,6 mm	$62,1 \pm 0,9 \text{ mm}$. ± 0,9 mm				
AAMxW	$56,2 \pm 2,3 \text{ mm}$	$60.9 \pm 1.3 \text{ mm}$ $52.3 \pm 0.9\%$		p<0,05			
BAMdW	56,8 ± 1,2 mm	$58,7 \pm 0,6 \text{ mm}$		p<0,05			
AAMdW	$55,5 \pm 2,1 \text{ mm}$	$59,6 \pm 1,4 \text{ mm}$		p<0,05			
LIIMx	18,1±3,5 mm	$6,4 \pm 1,5 \text{ mm}$		p<0,05			
LIIMd	13,9±2,4 mm	4,6±3,2 mm		p<0,05			
CBCT data	CGII						
	Before	After	Treatment efficiency	p-value			
	treatment	treatment					
BAMxW	58,9 ±1,8 mm	$63,2 \pm 1,7 \text{ mm}$		p<0,05			
AAMxW	$56,5\pm 2,1$ mm	$61,4 \pm 1,3 \text{ mm}$	58,1±1,7%	p<0,05			
BAMdW	$57,1 \pm 1,4 \text{ mm}$	$59,4 \pm 0,9 \text{ mm}$		p<0,05			
AAMdW	$55,9 \pm 2,6 \text{ mm}$	$60,8 \pm 2,4 \text{ mm}$		p<0,05			
LIIMx	17,8±2,9 mm	$4,7 \pm 1,8 \text{ mm}$		p<0,05			
LIIMd	11,7±2,5 mm	4,8±1,1 mm		p<0,05			
CBCT data	CGIII						
	Before	After	Treatment efficiency	p-value			
	treatment	treatment					
BAMxW	55,6 ±1,3 mm	$62,7 \pm 1,5 \text{ mm}$		p<0,05			
AAMxW	$53,2 \pm 2,5$ mm	$58.9 \pm 2.3 \text{ mm}$	66,7±1,6%	p<0,05			

 $58,7 \pm 0,6 \text{ mm}$

 $57,5 \pm 2,9 \text{ mm}$

 $6,4 \pm 1,5 \text{ mm}$

2.9±0.5 mm

Table 2. CBCT data values in clinical groups by suggested protocol.

Evaluating the results of treatment using a standard protocol, we found that in patients with neutralgrowth skeletal effect of maxillary expansion was $2,7\pm0,6$ mm, alveolar effect of expansion was $2,4\pm0,8$ mm, the

 $54,8 \pm 1,5 \text{ mm}$

 $55,2 \pm 2,1 \text{ mm}$

18,1±3,5 mm

 $13,9\pm 2,4 \text{ mm}$

skeletal effect of mandibular expansion was1,8±0,5mm, while at the alveolar level mandibular expansion was3,9 \pm 0,8mm; at the same time, we managed to reduce the Little's Irregularity Index value of upper teeth by 5,8 \pm 1,5 mm, at the

p<0,05

p<0,05

p<0,05

p<0,05

BAMdW

AAMdW

LIIMx

LIIMd



same time, we managed to reduce the Little's Irregularity Index value of lower teethby $4,3\pm0,5$ mm, , i.e., we were able to transfer the severity degree of crowdingonly from severe to moderate on both maxilla and mandible. The effectiveness of the treatment was $25,6\pm1,7\%$.

Patients with horizontalgrowthhadlesser skeletal effect of maxillaryexpansion-2,5±0,8 mm, alveolareffectofexpansionwas3,6±1,3mm., the skeletal effect of mandibularexpansion was2,1±0,5mm, while at the alveolar levelmandibular expansion was 2.8 ± 0.7 mm; at the same time, we managed to reduce the Little's Irregularity Index value of upper teeth by 6,2 \pm 1,4mm, we managed to reduce the Little's Irregularity Index value of lower teeth by 5,1±0,7 mm, were able to transfer the severity degree of crowdingfrom severe to moderate on both maxilla and mandible. The effectiveness of the treatment was22.3±2.1%.

Patients with verticalgrowthhadbetter skeletal effect of maxillaryexpansion-3,4±0,9 mm, alveolareffectofexpansionwas2,9±0,5mm., the skeletal effect of mandibularexpansion was 3.3±0.4mm. at the alveolar levelmandibularexpansionwas3,8 ± 1,3mm;at the same time, we managed to reduce the Little's Irregularity Index value of upper teeth by $4,2 \pm$ 1,3mm, at the same time, we managed to reduce the Little's Irregularity Index value of lower teeth by $3,4\pm0,3$ mm, were able to transfer the severity degree of crowdingfrom severe to moderate on both maxilla and mandible. The effectiveness of the treatment was29.5±2.4%.

III. DISCUSSION.

Comparing the obtained results of the effectiveness of the treatment with the results of the effectiveness of the treatment of tooth crowding in variable bite according to traditional methods[10, 11] a significant difference in quantitative data was observed. Thus, the effectiveness of treatment of patients of CG III reached 66.7±1.6% (for the results obtained from patients who were treated according to the traditional algorithm, this indicator was 29.5±2.4%); CG II - up to 58.1±1.3% (for the results obtained in patients who were treated according to the traditional algorithm, this indicator was $22.3\pm2.1\%$); CG I - $52.3\pm0.9\%$ (for the results obtained in patients who were treated according to the traditional algorithm, this indicator was 25.6±1.7%).

Conclusions.The results of our conducted statistical analysis of the proposed protocol application efficiency of dental crowding treatment

allowed to improve treatment quality of this pathology in children. The results obtained after 16 months in patients with a horizontal type of growth indicate that the effectiveness of the treatment of CG II patients reached 58.1±1.3%; after 17 months, in patients with a vertical and neutral type of growth, the effectiveness of the treatment of CG III reached 66.7±1.6%, CG I - up to 52.3±0.9%.Our proposed algorithm is more effective in patients with a neutral type of growth by $26.9 \pm 1.2\%$, in patients with a vertical type of growth by $37.3 \pm$ 0.7% and allows to shorten their total treatment period by 3 months: more effective in patients with a horizontal type of growth by $35.8 \pm 0.9\%$ and allows to shorten their total treatment period by 4 months.

Source of funding. This article received no financial support from a government, public or commercial organization.

Conflict of interest.The authors declare that they have no conflict of interest that could be perceived as prejudicing the impartiality of the article.

Consent to publish. All authors read and approved the final version of the manuscript. All authors agreed to publish this manuscript.

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A - Conception and design of the work, B - Data collection and analysis, C - Responsibility for the statistical analysis, D - Writing the article, E - Critical review, F - Final approval of the article

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