



The use of the Front Plateau anterior partial interocclusal device in the management of temporomandibular disorders: a pilot study

O uso do dispositivo interoclusal parcial anterior Front Plateau no manejo das disfunções temporomandibulares: um estudo piloto

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ABSTRACT

BACKGROUND AND OBJECTIVES: The partial occlusal splints, such as the Front Plateau, a device that involves the anterior teeth (from canine to canine), have been used by some professionals as an alternative to the full-coverage rigid occlusal splints due to the ease and time involved in their manufacturing. However, there is little evidence of its mechanism of action. Therefore, the aim of this study was to evaluate the effectiveness of the *Front Plateau* associated with the counseling therapy in controlling chronic myalgia associated with Temporomandibular Disorders (TMDs).

METHODS: This pilot study conducted with 10 patients with chronic myalgia related to TMDs. Two groups were assessed (G1: Front Plateau with the counseling therapy; G2: the counseling therapy only) with a follow-up of 3 months. Pain levels, quality of life and stress were assessed. A $p < 0.05$ was considered statistically significant.

RESULTS: Among the 10 patients assessed, 80% (n=8) were female. None of the variables evaluated showed statistically significant results ($p < 0.05$). However, the numerical pain assessment scale of the group that received the Front Plateau with the counseling therapy (G1) showed relevant clinical results, going from 8.00 ± 1.22 (Mean \pm SD) at baseline to 4.2 ± 1.79 (Mean \pm SD) after 3 months.

CONCLUSION: The Front Plateau with the counseling therapy seems to be a clinically viable alternative for controlling chronic myalgias related to TMDs. However, future clinical trials are needed to assess its applicability by professionals in the field.

KEYWORDS: Myalgia, Occlusal splints, Temporomandibular joint disorders, Conservative treatment.

RESUMO

JUSTIFICATIVA E OBJETIVOS: As placas oclusais parciais, como o *Front Plateau*, dispositivo que envolve os dentes anteriores (de canino a canino), têm sido utilizadas por alguns profissionais como alternativa às placas oclusais rígidas de cobertura total, devido à facilidade e ao tempo envolvido na sua confecção. Apesar disso, existem poucas evidências sobre o seu mecanismo de ação. Assim, o objetivo desse estudo foi avaliar a eficácia do *Front Plateau* associado com orientações no controle de mialgias crônicas associadas às Disfunções Temporomandibulares (DTMs).

MÉTODOS: Estudo piloto realizado com 10 pacientes com mialgias crônicas relacionadas às DTMs. Foram avaliados dois grupos (G1: *Front Plateau* associado às orientações; G2: apenas orientações) por um período de 3 meses. Foram avaliados os níveis de dor, a qualidade de vida e o stress. Um valor de $p < 0,05$ foi considerado estatisticamente significativo.

RESULTADOS: Dos 10 pacientes avaliados, 80% (n=8) eram do sexo feminino. Nenhuma das variáveis avaliadas apresentou resultados estatisticamente significativos ($p < 0,05$). No entanto, a escala numérica de avaliação da dor do grupo que recebeu o *Front Plateau* associado às orientações (G1) apresentou resultados clínicos relevantes, passando de $8,00 \pm 1,22$ (Média \pm DP) no *Baseline* para $4,2 \pm 1,79$ (Média \pm DP) após 3 meses.

CONCLUSÃO: O *Front Plateau* associado às orientações parece ser uma alternativa clinicamente viável para o controle das mialgias crônicas relacionadas com as DTMs. No entanto, são necessários futuros ensaios clínicos para avaliar a sua aplicabilidade pelos profissionais da área.

DESCRIPTORIOS: Mialgia, Placas oclusais, Transtornos da articulação temporomandibular, Tratamento conservador.

HIGHLIGHTS

- Minimally invasive techniques for controlling TMDs, such as the Front Plateau associated with the counseling therapy, have shown good results
- Since it is easy to fabricate, the Front Plateau reduces clinical time and can be made in a single session
- This study highlights the importance of evaluating clinical research beyond the p-value

INTRODUCTION

Temporomandibular Disorders (TMDs) represent a set of muscular and/or articular conditions that can result in orofacial pain, involving the masticatory muscles, the Temporomandibular Joint (TMJ), bone tissues and adjacent soft tissues^{1,2}. The etiology of TMDs is complex and multifactorial, involving physical (axis I) and/or psychological (axis II) factors that can modulate the intensity and perception of pain by the patient, making its diagnosis challenging³⁻⁵.

These alterations affect approximately 5% to 12% of the world's population, mainly adult women between the ages of 30 and 40 and are considered a public health problem⁶⁻⁸. The main signs and symptoms include pain on palpation, joint noises, headaches, otalgia, facial pain, muscle fatigue, spasms and mandibular deviations⁹. The treatment of TMDs is generally aimed at identifying and intervening in the main causal factors. Therapeutic approaches include total or partial occlusal splints, acupuncture, laser therapy, patient counseling, physiotherapy, pharmacotherapy and arthrocentesis, which can be combined according to clinical need^{10,11}. Priority is given to non-invasive and reversible interventions and, if they are not effective, more invasive procedures are used, with the aim of achieving the patient's functional, emotional and orthopedic balance¹².

The full-coverage rigid occlusal splint is a non-invasive and effective intervention in the management of TMDs, providing both short- and long-term muscle relief. Its benefits go beyond the placebo effect, providing pain relief, psychological well-being and patient satisfaction¹³. However, their fabrication can involve multiple clinical steps and often requires a laboratory phase, which can increase the cost for the patient^{11,13,14}. In this context, partial occlusal devices, such as the Nociceptive Trigeminal Inhibition Tension Suppression System (NTI-tss), have been developed as an alternative to the full-coverage occlusal splints^{15,16}. The NTI-tss is a prefabricated material that covers only the incisors and offers disocclusion of the posterior teeth, providing muscle relaxation. However, due to their small size, there have been reports in the literature of these devices being aspirated and swallowed, posing a risk to patients' lives¹⁷.

Some professionals suggest the use of a partial interocclusal device that covers the 6 anterior teeth, called a Front Plateau, made from self-curing colorless acrylic resin, presenting a low-cost alternative. The device can be made in a single session, directly in the patient's arch, helping to control orofacial pain and the differential diagnosis of TMDs^{18,19}. Another widely used approach, associated with occlusal splints or not, is Counseling Therapy (CT), based on personalized oral and written guidance to prevent and eliminate parafunctional habits related to TMDs, promoting patients' well-being through an adaptable and daily approach²⁰⁻²².

Although several scientific studies have evaluated the effectiveness of various treatment modalities for TMDs, research into interocclusal devices, such as the Front Plateau, is still limited. In this sense, well-designed studies are essential to identify effective interventions that can be adopted by various professionals^{11,17,23}. Given this, the aim of the present study was

to evaluate the effectiveness of the Front Plateau associated with CT, on the level of pain, quality of life and stress, with a follow-up of 3 months, in patients with chronic myalgias related to TMDs. A control group that only received CT was used as a comparative.

METHODS

This study was conducted in accordance with the Consolidated Standards of Reporting Trials (CONSORT)²⁴ and was approved by the Research Ethics Committee (REC), number CAAE 70486823.0.0000 in full compliance with the Declaration of Helsinki, revised in 2013. After explaining the research, the individuals who agreed to take part signed the Free Informed Consent Term (FICT).

Study design

This research was conducted from January to July 2024 with patients diagnosed with chronic myalgias related to TMDs. As this was a pilot study, it was not based on any hypothesis, ruling out the feasibility of a sample calculation based on statistical power²⁵. An n=5 per group was adopted as sufficient to assess the feasibility of a randomized clinical trial with the same interventions. A detailed description of the patients who completed the study can be found in Figure 1.

Eligibility criteria

The inclusion criteria were patients aged 18 or over, with TMDs-related myalgia for more than 6 months, located in the temporal region, accompanied or not by other facial myalgias. A numerical pain assessment scale from 0 to 10 was used to collect the patients' data, and those who reported pain equal to or greater than 7 were included. Patients who were undergoing any treatment for TMDs, including physiotherapy, follow-up with a speech therapist, laser therapy and botulinum toxin-A application (in the last 6 months) were excluded. Patients with other types of headaches that were not associated with the stomatognathic system, such as migraine, were also excluded. In addition, individuals with full dentures, undergoing orthodontic treatment, undergoing Selective Serotonin Reuptake Inhibitors (SSRIs) treatment or who had a cognitive impairment that made it difficult to participate in the study were not included in the sample.

Interventions

Front Plateau

It is a rigid partial plate that covers the anterior region, from canine to canine, respecting the movements of protrusion and laterality. The procedure was performed by a professional duly calibrated by a TMD specialist following a standardized protocol^{18,19}. The following materials were used to make the Front Plateau: self-

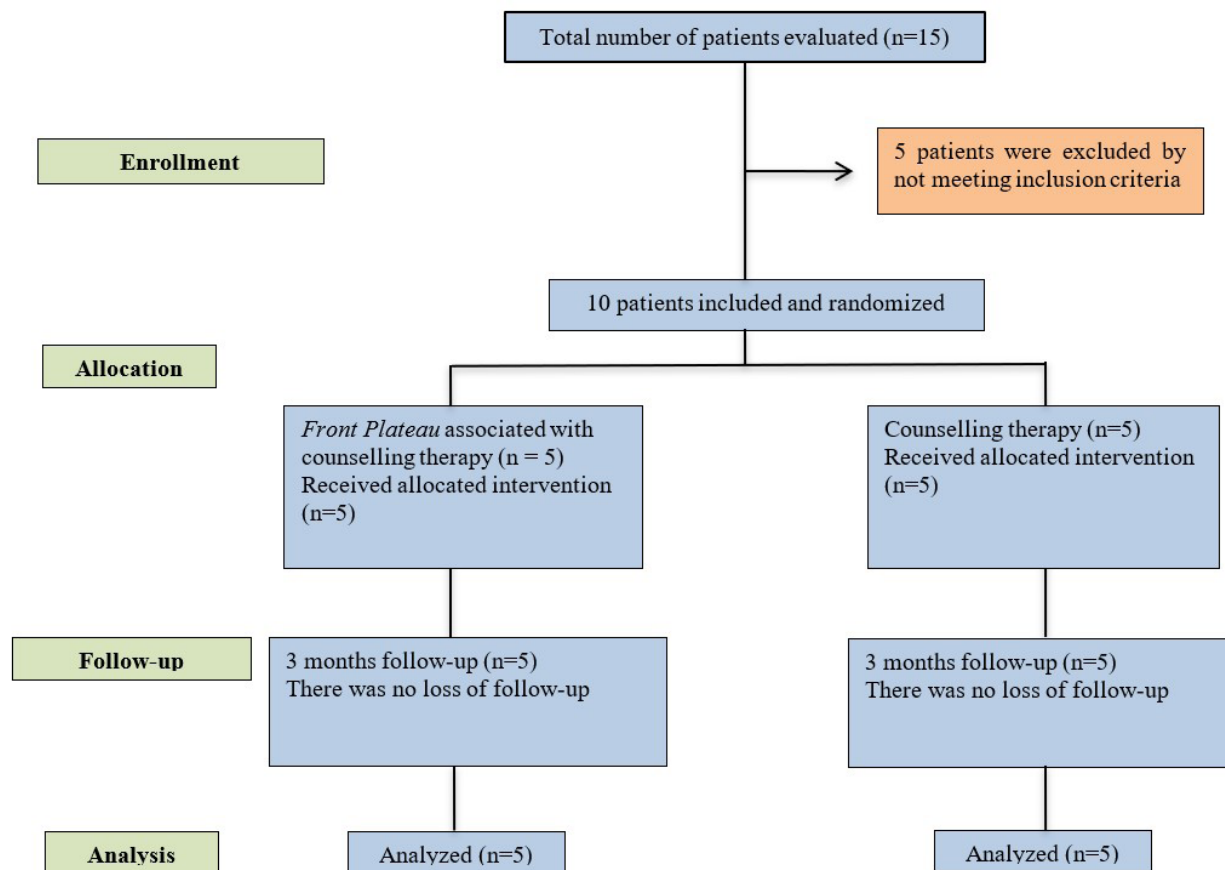


Figure 1. Flowchart of sample distribution.

curing colorless acrylic resin liquid and powder, dosing cup, kit of cutters and polishing tips, carbon paper, Muller tweezers, No. 31 spatula, brush, paladon pot, water tank and procedure gloves.

After organizing the materials, the acrylic resin powder was mixed with the liquid in the paladon pot for handling. Once a homogeneous mixture had been obtained, the paladon pot was closed and the acrylic resin was allowed to lose its shine and reach its plastic phase.

Next, the patient's teeth and the operator's fingers were insulated with liquid petroleum jelly to prevent the resin from sticking. The resin was manipulated with the operator's fingers until it took the shape of a stick, which was adapted to the area of the patient's upper front teeth. The patient was instructed to always bite down with the tongue on the palate to ensure that the Front Plateau acted in centric relation. During the setting time, due to the acrylic resin heating up, the Front Plateau was removed from the patient's mouth and placed to rest in a tub of water. The material was then returned to the patient's mouth. After the setting time, occlusal adjustments were made, ensuring that all anterior teeth were touching, and that protrusion and laterality movements were present.

At the end, the Front Plateau was finished and polished with a kit of cutters and abrasive tips for delivery to the patient. The patients were advised to wear the device at night, when they went to sleep, and to remove it when they woke up in the morning.

Counseling therapy

These include guidance on exercises for muscle relaxation, diet, sleep hygiene and ways to prevent and/or control the symptoms associated with TMDs. The instructions were given by a single professional, calibrated by a TMD specialist. During the 1- and 3-month meetings, patients were asked if they were following the CT and if they had any questions, they would be answered by the same calibrated professional.

Randomization, allocation concealment and masking

The randomization process was carried out using a block system by an independent researcher who was unaware of the patients, the intervention and the procedure for the control group. A total of 5 blocks were defined with 2 patients each. Each patient was randomly represented by a number from 1 to 10, the representation of which was unknown to the team directly involved in the research. Then, for each block, a number was placed in an opaque envelope along with a letter (A or B). These envelopes were sealed and only revealed on the day of the interventions. It was only at this point that the professionals responsible for each intervention found out which patient was going to receive which procedure. At the end of the session, the patients were instructed not to reveal the treatment they had received on the evaluation days after 1 month and 3 months, respectively.

Data collection

A clinical form adapted from the Diagnostic Criteria for Temporomandibular Disorders (DC/TMD) and a numerical pain assessment scale (1 to 10) were used for data collection at baseline and after 1 and 3 months. The Oral Health Impact Profile (OHIP-14) and Lipp's Inventory of Stress Symptoms for Adults (ISSL) were only applied at baseline and at the 3-month evaluation.

Statistical analysis

Were carried out using SPSS® for Windows® (Statistical Package for the Social Sciences Inc.) version 26. Exploratory data analysis provided frequencies, means and standard deviations. Normality was assessed using the Shapiro-Wilk test. Quantitative data was subjected to the Mann-Whitney test. Categorical data was analysed using the Chi-square test. Confidence intervals of 95% and a significance level of 5% were used.

RESULTS

Characterization of the sample

A total of 10 patients took part in the study, 5 of whom were allocated to G1 (Front Plateau and CT) with a mean age of 25.4 ± 7.77 (mean \pm SD) and 5 to G2 (CT) with a mean age of 26.2 ± 5.67 (mean \pm SD). Most patients were female, representing 80% (n=8) of the total sample. Among the diagnostic impressions, Myalgia and TMJ Disc Displacement with Reduction (DDwR) was present in all patients and hypermobility was identified in 60% of the sample (n=6). In addition, all the patients reported clenching their teeth, 30% (n=3) of which were accompanied by teeth grinding and 10% (n=1) onychophagia. No adverse effects were reported by patients.

Numerical pain assessment scale

The results of the numerical pain assessment scale used at baseline and after 1 and 3 months can be seen in Table 1. There were no statistically significant differences ($p < 0.05$), but it was possible to observe a clinical difference in G1 after 1 and 3 months.

Quality of life (Oral Health Impact Profile-14) and stress (ISSL)

The results for quality of life and stress after the treatments were summarized in Tables 2 and 3, respectively. For both variables, there were no statistically significant results.

Pain during active movement and pain on palpation

The assessment of pain during active movement considered the movements of maximum mandibular opening, protrusion and laterality on the right and left sides. On the other hand, pain on palpation was recorded in all the main masticatory muscles, especially the masseter and temporal muscles, as defined by the DC/TMD (Tables 4 and 5). None of these variables showed statistically significant results.

DISCUSSION

The effectiveness of occlusal devices in controlling symptoms associated with TMDs is still controversial in the literature²⁶. From the results of this pilot, the Front Plateau associated with CT, despite not presenting statistically significant data, obtained better clinical performance than the group that only received CT. These findings corroborate some studies in which the Front Plateau helped control pain, as well as identifying possible etiological factors^{18,19}. In clinical trials comparing a device like the Front Plateau, called NTI-tss with CT, both groups had a reduction in pain levels^{27,28}. The same result was found when compared with full-coverage rigid occlusal splints^{21,29,30}. Nevertheless, therapy with occlusal splints seems to promote an earlier improvement in symptoms, as found in the evaluation of the numerical pain assessment scale in this and other studies^{27,31}.

The variety of diagnostic criteria, types of devices and protocols used in randomized clinical trials in the field of TMDs make it difficult to assess the level of evidence found³². The use of occlusal splints is indicated by some professionals because they are more conservative approaches, presenting fewer risks and side effects, and are most often associated with exercises and CT for TMD patients³³. When used in isolation, in other words, without CT and compared to a group that only received CT for TMDs, occlusal splints performed less well than control group³⁴.

Table 1. Numerical pain assessment scale results.

	Front Plateau and CT		CT		p-value
	Mean	SD	Mean	SD	
Baseline	8.00	1.22	8.80	1.10	0.212
Assessment after 1 month	3.00	2.24	7.40	2.88	0.035
Assessment after 3 months	4.20	1.79	7.00	2.24	0.056

CT = Counselling Therapy; SD = standard deviation. A value of $p < 0.05$ was considered statistically significant.

Table 2. Quality of life assessment after 3 months (OHIP-14).

	Front Plateau and CT		CT		p-value
	Mean	SD	Mean	SD	
Baseline Total	23.80	8.56	19.20	8.64	0.458
Functional limitation	1.80	1.30	1.00	1.00	0.281
Physical pain	5.60	1.52	6.00	1.87	0.523
Psychological discomfort	4.80	1.92	5.40	1.95	0.655
Physical disability	2.00	1.58	1.40	1.67	0.522
Psychological disability	3.40	3.13	2.20	2.68	0.458
Social disability	3.20	1.79	1.60	1.67	0.178
Disadvantages	3.00	1.22	1.60	1.14	0.101
After 3 months Total	22.40	8.79	18.40	10.26	0.917
Functional limitation	1.80	1.48	1.00	1.73	0.277
Physical pain	4.40	2.19	5.00	2.00	0.655
Psychological discomfort	4.40	2.51	4.40	2.51	0.833
Physical disability	1.60	1.67	1.00	1.41	0.572
Psychological disability	3.40	3.58	2.00	1.58	0.671
Social disability	3.40	2.30	2.80	2.59	0.833
Disadvantages	3.40	1.52	2.20	2.59	0.522

CT = Counseling Therapy. The value of $p < 0.05$ was considered statistically significant.

Table 3. Stress assessment after 3 months (ISSL).

	Front Plateau and CT		CT		p
	n	%	n	%	
Baseline					
No stress	1	20.00	2	40.00	
Resistance	2	40.00	3	60.00	
Almost exhaustion	1	20.00	0	0.00	
Exhaustion	1	20.00	0	0.00	0.469
After 3 months					
No stress	0	0	1	20.00	
Resistance	4	80.00	2	40.00	
Almost exhaustion	1	20.00	1	20.00	
Exhaustion	0	0.00	1	20.00	0.446

CT = Counselling Therapy. $p < 0.05$ was considered statistically significant.

The fact that most patients were female reaffirms epidemiological findings in the field of TMDs, since these conditions are more common in women^{3,35}. Regarding parafunctional habits that can predispose to TMDs, dental clenching associated or not with teeth grinding and onychophagy were identified. These habits associated with psychological factors such as anxiety and stress, present in most of the patients in this study, can trigger or amplify TMDs^{36,37}.

In addition, conditions such as hypermobility were identified in some of the patients in the present study. According to a Randomized Clinical Trial (RCT) that evaluated the relationship between hypermobility and TMDs, it was possible to observe a significant association between the two conditions, especially in adult patients with disorders involving the articular disc³⁸. A considerable number of the patients assessed in this pilot had

Table 4. Temporal muscle palpation assessment.

	Front Plateau and CT		CT		p-value
	Mean	SD	Mean	SD	
Baseline - RPTM	1.20	0.84	1.00	1.00	0.740
1 month - RPTM	0.60	0.89	0.80	1.10	0.811
3 months - RPTM	0.00	0.00	0.40	0.89	0.317
Baseline - LPTM	1.00	0.71	1.00	1.00	0.999
1 month - LPTM	1.60	0.89	1.00	1.41	0.332
3 months - LPTM	1.40	0.55	0.40	0.89	0.060
Baseline - RMTM	1.40	1.14	1.00	1.41	0.514
1 month - RMTM	1.00	0.71	0.80	1.10	0.658
3 months - RMTM	0.80	0.45	0.80	0.84	0.905
Baseline - LMTM	1.80	0.84	1.20	1.30	0.389
1 month - LMTM	1.20	1.30	1.20	1.30	0.999
3 months - LMTM	1.20	0.84	1.00	1.22	0.584
Baseline - RATM	1.00	1.22	0.80	1.10	0.736
1 month - RATM	0.40	0.55	1.00	1.41	0.637
3 months - RATM	0.40	0.55	0.60	1.34	0.699
Baseline - LATM	1.40	1.14	0.80	1.10	0.391
1 month - LATM	0.40	0.55	0.40	0.55	0.999
3 months - LATM	1.00	0.00	0.80	1.30	0.232

CT = Counseling Therapy; RPTM = Right Posterior Temporal Muscle; LPTM = Left Posterior Temporal Muscle; RMTM = Right Middle Temporal Muscle; LMTM = Left Middle Temporal Muscle; RATM = Right Anterior Temporal Muscle; LATM = Left Anterior Temporal Muscle. The value of $p < 0.05$ was considered statistically significant.

Table 5. Masseter muscle palpation assessment.

	Front Plateau and CT		CT		p-value
	Mean	SD	Mean	SD	
Baseline - RSM	2.20	0.45	1.20	1.64	0.439
1 month - RSM	1.40	0.55	1.40	1.14	0.910
3 months - RSM	1.20	0.45	1.40	1.14	0.813
Baseline - LSM	2.20	0.45	1.20	1.64	0.439
1 month - LSM	1.60	0.89	1.60	1.34	0.910
3 months - LSM	1.60	0.55	1.60	1.34	0.913
Baseline - RMM	2.00	0.71	1.20	1.30	0.277
1 month - RMM	2.20	0.84	1.60	1.14	0.381
3 months - RMM	1.80	0.84	1.00	1.22	0.192
Baseline - LMM	2.60	0.55	1.40	1.52	0.217
1 month - LMM	2.00	1.00	2.00	1.22	0.913
3 months - LMM	2.20	0.45	1.80	1.10	0.606
Baseline - RIM	2.40	0.89	1.60	1.52	0.369
1 month - RIM	1.60	1.34	1.80	1.10	0.829
3 months - RIM	1.80	0.84	1.80	1.10	0.822
Baseline - LIM	2.60	0.55	2.00	1.22	0.419
1 month - LIM	1.60	1.34	1.80	1.30	0.827
3 months - LIM	2.00	0.71	1.40	1.52	0.518

CT = Counseling Therapy; RSM = Right Superior Masseter; LSM = Left Superior Masseter; RMM = Right Middle Masseter; LMM = Left Middle Masseter; RIM = Right Inferior Masseter; LIM = Left Inferior Masseter. The value of $p < 0.05$ was considered statistically significant.

hypermobility and DDwR. Despite this, some authors report that it cannot be said that there is a causal relationship between DDwR and hypermobility^{39,40}. This is because the presence of DdwR does not necessarily seem to depend on the existence of hypermobility, but rather on biomechanical factors that occur during adolescence^{41,42}. In the case of DDwR, it is only of concern and will require intervention if it interferes with TMJ function⁴².

When looking at the quality of life and stress assessment data, no statistically significant results were identified. However, it was possible to observe that in the group that received Front Plateau and CT there was a greater migration of patients to the resistance phase when stress was assessed after 3 months. In the group that only received CT, 2 patients moved from the stress-free and resistance phases to near-exhaustion and exhaustion, respectively. Significant results of improved quality of life and stress due to the use of occlusal splints have been found in studies with an evaluation time of at least 6 months^{11,43}. When comparing the results of studies with short evaluation times with studies that evaluated patients for up to 12 months, it was identified that a longer evaluation time provides more relevant and, consequently, statistically significant results⁴⁴.

As this is a pilot study, the results of the research should be evaluated with caution. It is worth noting that the primary aim of a pilot is to assess the feasibility of a clinical trial, allowing for adaptations of protocols, sample calculation and results to be evaluated⁴⁵. The absence of statistically significant results does not make the study unfeasible, as the p-value can mask relevant clinical changes not identified in the statistical analysis⁴⁶. In such cases, a critical interpretation of the results beyond the p-value is suggested, as in the case of recording the level of pain observed in the patients in this study. It is believed that an RCT with a larger sample and an evaluation time of more than 3 months could show more significant results^{27,31}.

CONCLUSION

The Front Plateau, when associated with CT for TMD patients, appears to be a clinically viable alternative for controlling chronic myalgia. Future clinical studies with greater methodological rigor, sample size and follow-up time are needed to assess the applicability of the device by professionals in the field.

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