




## Comparative effect of aerobic physical activity and occlusal stabilization appliances on musculoskeletal orofacial pain in individuals with temporomandibular disorders

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### Abstract

This study evaluated the efficacy of aerobic physical activity (APA) in managing musculoskeletal orofacial pain in individuals with temporomandibular disorders (TMD) compared to treatment with an occlusal stabilization appliance (OSA). Volunteers with musculoskeletal pain aged between 18 and 40 and with bimaxillary teeth involved. The analog verbal scale was used to evaluate the pain variable. Each treatment group included seven individuals evaluated for eight (APA) and six (OSA) weeks, respectively. APA showed a significant weekly decrease of 0.155 points in pain measurement. As for the OSA, the trend was even stronger than in the APA treatment—the difference was substantial.

This study demonstrated that both the OSA and APA contributed to reducing musculoskeletal orofacial pain in volunteers suffering from TMD.

**Keywords:** temporomandibular disorders, musculoskeletal pain, aerobic exercise, occlusal devices.

## Introduction

Temporomandibular disorders (TMD) are a heterogeneous group of pathologies that affect the temporomandibular joints (TMJ), the masticatory muscles, or both<sup>(1)</sup>. The glossary of prosthodontic terms describes them as conditions that produce an abnormal, incomplete, or altered function of the TMJ and the masticatory muscles<sup>(2)</sup>. Temporomandibular pain is relatively common, affecting approximately 10% of the population over the age of 18. This disorder mainly affects young adults, middle-aged people, and it is nearly twice as common in women as in men<sup>(3)</sup>.

TMDs are highly prevalent across the general population. The prevalence of signs in a study conducted in a European population is 34.5%<sup>(4)</sup>. Another study carried out in Australia, in a cohort of dental students, showed a prevalence of TMD symptoms of 77.2%<sup>(5)</sup>. In Uruguay, 55% of the population has at least one symptom, and 44% of the

population has at least one sign<sup>(6)</sup>. These studies also show a higher prevalence in women.

It is crucial to identify the main factors leading to TMD to select the most appropriate and effective treatment for each patient. Reversible therapies most commonly used for the pain symptom include therapy with an occlusal stabilization appliance (OSA), pharmacotherapy, physiotherapy, psychotherapy (behavioral strategies), among others. These can be indicated separately or in combination<sup>(7)</sup>.

OSAs have proved to be efficient in treating myalgia and arthralgia of the masticatory system<sup>(8)</sup>. However, their action mechanism is unclear and still controversial.

Different methodological problems relating to research designs have made it difficult to understand the real effectiveness and action mechanisms of OSAs. For example, among the issues detected in previous studies are a small sample size, inadequate blinding, and short follow-up times<sup>(9)</sup>.

Aerobic physical activity (APA) and manual therapy are the therapeutic measures that have proved to be most effective in changing attitudes to pain and stress are<sup>(10)</sup>.

Past studies have shown that APA interventions have positive results for pain reduction in fibromyalgia treatment<sup>(11-13)</sup>.

To date, no studies have been conducted to evaluate whether APA is effective in reducing pain in patients with TMD muscle pain.

Beta-endorphin (BE) is a hormone, an endogenous opioid, secreted by the anterior pituitary gland. Among the most relevant effects are analgesia, increased lactate tolerance, and decreased muscle discomfort<sup>(14)</sup>. Its bloodstream levels increase during prolonged APA, thus reducing pain. It shows greater resistance to enzymatic degradation compared to enkephalins<sup>(15)</sup>.

Valim, quoting other authors, argues that APA influences the serotonergic system, increases sympathetic activity, improves sleep, and promotes psychological well-being<sup>(11)</sup>.

This study assessed APA's efficacy in managing musculoskeletal orofacial pain in people with TMD, compared to treatment with an OSA.

## **Materials and methods**

### **Materials**

This was a comparative, prospective, longitudinal, controlled, and randomized clinical study.

The sample was selected among students of the School of Dentistry, Universidad de la República. Uruguay. It included 14 female volunteers. The recruitment method was consecutive, and a public call was made in 2016, seeking volunteers suffering from orofacial pain.

Ethical considerations: The project was approved by the Ethics Committee of the School of Dentistry of Universidad de la República (File 249/15).

Volunteers signed an informed consent before their participation.

Inclusion criteria: volunteers with musculoskeletal orofacial pain due to TMD, aged between 18 and 40 years, with bimaxillary teeth and who presented proof of physical fitness issued by a competent institution.

Exclusion criteria: volunteers with pain resulting from reasons other than musculoskeletal due to TMD, pregnant women who regularly do aerobic physical activity, OSA carriers at the time of the intervention, chronic and/or acute respiratory conditions, severe psychiatric disorders, unstable or untreated heart diseases and with a history of a heart attack in the past six months.

Elimination criteria: those who did not complete at least 80% of the APA sessions, as well as patients who did not meet the OSA indications for use and controls.

## **Method**

A general, regional, and local clinical examination of the patients was undertaken to check the inclusion and exclusion criteria.

Volunteers with musculoskeletal orofacial pain-related TMD were separated from those of other origin, following the DC-TMD criteria<sup>(16)</sup> and the calibration protocol of the Riva et al. study of May 2011<sup>(6)</sup>. Palpation of masticatory muscles (masseter and temporal) and external pole of the TMJ was performed.

### **Measuring instruments and variables**

The verbal analog scale (VAS) was used to evaluate the pain and emotional stress variables. For data collection, we used a survey that includes the aforementioned measuring instrument (Fig. 1). The survey was used on several occasions, from the beginning to the end of the interventions, to evaluate the progression of the symptom reported by the patient. Each of the treatment groups consisted of seven people, who were evaluated for eight (APA) and six (OSA) weeks, plus an initial evaluation to determine mean values at baseline. In each assessment, self-reports of pain and stress were recorded at three times of the day: morning, afternoon, and evening. For the initial evaluation, they were provided with a pain journal two weeks before the intervention to determine an average baseline pain. This considers pain symptoms and fluctuations.

#### **Fig. 1:**

GROUP: APA/ OSA	VOLUNTEER N°				
Date	PAIN (0 TO 10)	STRESS (0 TO 10)	MEDICATION	OTHER TREATMENTS	AGGRAVATING EVENTS
MORNING					
AFTERNOON					
EVENING					

## Interventions and Groups

The sample was randomly divided into two groups of seven volunteers each:

Group A (GA): people who did APA exclusively and Group B (GB) people who were fitted with an OSA.

The OSAs were made at the School of Dentistry clinic using a printed splint technique and made by a specialist considering ideal occlusion criteria.

The APA was supervised by advanced students and teachers from the Higher Institute of Physical Education (ISEF) of Universidad de la República. The type of APA was 60-minute aerobic fitness sessions. Each session started with a 5 to 10 min warm-up, followed by moderate aerobic exercises, which did not exceed 10 to 30% of the  $VO_2$  max<sup>14</sup>; the central part with planned exercises and the final stretch, in which aerobic activity is maintained, at a low to moderate intensity (5 to 10 minutes), with an exercise progression approach<sup>(17)</sup>.

Due to the study's limitations, to ensure BE release, we used a heart rate meter (*Advanced 3 Müller Pulse Oximeter*) and Karvonen's formula; by entering the data, we obtained the oxygen consumption level. To be considered BE release-related, it had to be between 60% and 85%<sup>(18-19)</sup>.

## Statistical methodology

Pain and stress levels were recorded multiple times (weekly and three times a day). The reduction in both variables (pain and stress) was compared using mixed-effect regression models [1], where inferences were made with a 5% significance level. The statistical software used was R[2].

The evolution of each group was compared using a regression model that compared the trends for each treatment at different times of the day. It should be noted that the control variable for this model was the presence of aggravating factors since this could skew each person's measurements at the different points in the study. This article presents the results of the pain variable.

## Results

## Pain variable

### Descriptive analysis

Table 1 shows the average evolution of both treatments. The APA group had a longer follow-up period where a downward trend in pain values seems to be observed.

**Table 1:** Weekly evolution of pain based on the type of treatment

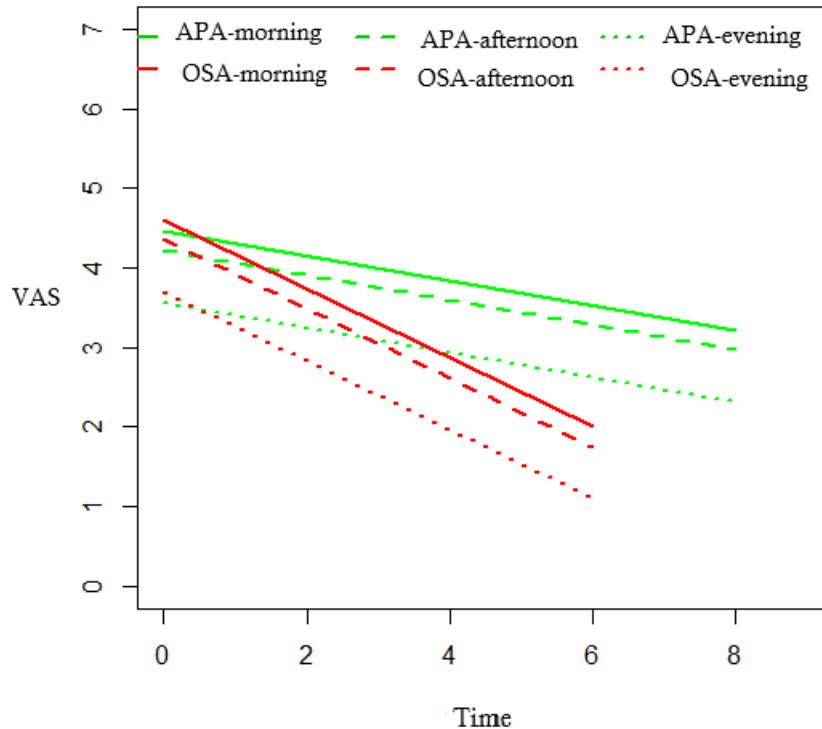
	Week									
	0	1	2	3	4	5	6	7	8	
<b>APA</b>	<b>4.10</b>	<b>3.70</b>	<b>4.00</b>	<b>3.57</b>	<b>4.05</b>	<b>2.05</b>	<b>3.90</b>	<b>3.52</b>	<b>2.52</b>	
Morning	4.43	5.00	4.29	3.14	3.71	2.43	4.14	4.14	3.43	
Afternoon	3.41	2.43	3.43	3.43	4.14	2.00	3.57	2.43	1.71	
Evening	4.47	3.86	4.29	4.14	4.29	1.71	4.00	4.00	2.43	
<b>OSA</b>	<b>4.51</b>	<b>3.89</b>	<b>3.17</b>	<b>2.71</b>	<b>3.35</b>	<b>2.00</b>	<b>1.86</b>			
Morning	4.51	4.67	3.83	3.14	3.67	2.57	2.57			
Afternoon	4.51	2.83	2.67	2.29	2.57	1.29	1.29			
Evening	4.51	4.17	3.00	2.71	3.86	2.14	1.71			
<b>Tot</b>	<b>4.31</b>	<b>3.79</b>	<b>3.62</b>	<b>3.14</b>	<b>3.71</b>	<b>2.02</b>	<b>2.88</b>	<b>3.52</b>	<b>2.52</b>	

**Table 2:** Mixed-effect regression model

	estimation	p-value
APA (start)	4.463	0.000
OSA (start) compared to APA	0.141	0.889
APA reduction per week	-0.155	0.004
Evening-morning	-0.908	0.000
Afternoon-morning	-0.247	0.330
aggravating event	0.794	0.057
OSA difference with APA per week	-0.279	0.004

Figure 1 shows the average evolution based on the results of the previously estimated regression model.

**Figure 1:** Pain evolution based on the type of treatment



The graph shows pain reduction is more significant with an OSA (red lines) than APA interventions (green lines). The three lines, corresponding to each treatment, show the evolution in the three moments of the day. Note that pain values at night are lower than those recorded in the morning and afternoon.

## Discussion

The first two lines of Table 2 indicate that the average pain value of the two groups was not significantly different, and the average was 4.463 points. The third line shows the first relevant result, which indicates that APA treatment resulted in a significant weekly reduction of 0.155 points in pain measurement. Building on the above statement, the last row shows that the trend was even more pronounced with OSA treatment than with APA treatment, and the difference is significant. By combining the two estimates, we see that pain reduction among the OSA group was almost half a point (0.434) per week.

In addition, the pain perceived by the participants was less intense (almost one VAS point) at night compared to the morning, and that the aggravating factors recorded were significant, so much so that when faced with these factors, participants' perception of pain was on average 0.794 points higher than usual.

This study found significant differences between the two groups following interventions considering the pain variable. Although APA significantly reduced pain, the trend was higher with OSA, and the difference in reduction between the two groups was significant. The evaluation period of the OSA group was six weeks. This study shows its effectiveness in reducing pain. This result is in line with studies such as that of Dao and Lavigne 1994, which shows pain reduction across a ten-week evaluation period, but without explaining

the mechanism for pain reduction<sup>(8)</sup>. In a subsequent literature review, Kreiner concluded that with the data available so far, the use of OSA for muscle and joint pain had sufficient scientific support<sup>(9)</sup>. In 2011, a randomized clinical study compared the efficacy of OSA and physical therapy (Tens) for TDM pain management across a four-month period for OSA and a four-week period for physical therapy. The study found a significant difference in pain reduction between the two, and OSA is the most effective<sup>(20)</sup>. The latter is in line with other studies on the efficacy of OSA for pain reduction in five-week<sup>(21)</sup> and ten-week periods, respectively<sup>(22)</sup>.

Our work is consistent with the studies reviewed, and despite controversy regarding the action mode or mechanism, OSAs do reduce musculoskeletal pain in patients with TMD, and despite varying evaluation times, their effectiveness is recognized.

APA showed a slower reduction than OSA; this could be because volunteers were sedentary and the intensity of the APA was gradually increased as the days of the intervention progressed. APA intensity is required to ensure BE release<sup>(14)</sup>. Some studies have shown that BE levels did not vary with low-intensity exercise, with oxygen volumes ranging from 25% to 60%<sup>(23-25)</sup>.

Another study of 12 untrained men found that pedaling at 60% oxygen volume did not significantly change BE levels. However, at 70 and 80% intensities, increases were significant, 2 to 5 times the baseline values, respectively<sup>(26)</sup>. Years later, the same author didn't find a gender difference in BE levels at 60% and 80% intensity, respectively<sup>(27)</sup>.

One of the most difficult parameters to estimate is the intensity of physical activity. There are different methods for measuring it: measuring heart rate, the person's perception of the activity, and perceived intensity using a standardized classification system<sup>(28)</sup>. Oxygen volume was not measured directly in our study; it was done using Karvonen's formula<sup>(19)</sup>. The study did not aim to measure BE.

The 2006 review article by Nishishiya assessed the efficacy of APA in fibromyalgia patients. It confirmed that APA is the most studied exercises and concluded that there is moderate evidence that APA helps improve pain in these patients. Likewise, there is no evidence that APA worsens the clinical manifestations of people with fibromyalgia<sup>(29)</sup>.

More recently, a review article published in 2017 states that the quality of the evidence to assess physical activity and exercise for chronic pain is low. This is primarily due to small sample sizes and potentially weak studies. Favorable effects were detected in reducing the severity of pain and improving physical function, although these were mainly of low to moderate impact and were not consistent in all reviews<sup>(30)</sup>. In this sense, our study would have the same effects as those reported in that article in terms of pain.

Available evidence suggests that physical activity and exercise have few adverse effects and can improve pain intensity, physical function, and the resulting quality of life. However, further research is needed and should focus on increasing the number of participants, including participants with a broader VAS pain level range, and extend both the intervention itself and the follow-up period<sup>(30)</sup>.

## Conclusions

This study showed that both OSA and APA helped reduce musculoskeletal orofacial pain in TMD volunteers.

The comparative study between these treatments showed that pain reduction was most effective in the OSA group.

Future studies should review and analyze the possibility of implementing other methodological designs for more accurate results: longer evaluation periods, larger samples, more diverse APA interventions.

It would be interesting to work with designs where both groups are fitted with an OSA, and one of the groups also undertakes APA. It would also be interesting to work with three groups: one with OSA, another with APA, and the third one with OSA and APA.

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#### **Authors' contribution note:**

1. Conception and design of study
  2. Acquisition of data
  3. Data analysis
  4. Discussion of results
  5. Drafting of the manuscript
  6. Approval of the final version of the manuscript.
- LARF has contributed in 1, 2, 3, 4, 5 y 6.  
LS has contributed in 2, 3 y 4.  
RRB has contributed in 1, 3, 4 y 6

#### **Editor's opinion:**

This article has been accepted by the Odontostomatología's editor Dra. Vanessa Pereira-Prado

