Dysregulation of the Descending Pain System in Temporomandibular Disorders Revealed by Low-Frequency Sensory Transcutaneous Electrical Nerve Stimulation: A Pupillometric Study

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Abstract

Using computerized pupillometry, our previous research established that the autonomic nervous system (ANS) is dysregulated in patients suffering from temporomandibular disorders (TMDs), suggesting a potential role for ANS dysfunction in pain modulation and the etiology of TMD. However, pain modulation hypotheses for TMD are still lacking. The periaqueductal gray (PAG) is involved in the descending modulation of defensive behavior and pain through μ, κ, and δ opioid receptors. Transcutaneous electrical nerve stimulation (TENS) has been extensively used for pain relief, as low-frequency stimulation can activate μ receptors. Our aim was to use pupillometry to evaluate the effect of low-frequency TENS stimulation of μ receptors on opioid descending pathways in TMD patients. In accordance with the Research Diagnostic Criteria for TMD, 18 females with myogenous TMD and 18 matched-controls were enrolled. All subjects underwent subsequent pupillometric evaluations under dark and light conditions before, soon after (end of stimulation) and long after (recovery period) sensorial TENS. The overall statistics derived from the darkness condition revealed no significant differences in pupil size between cases and controls; indeed, TENS stimulation significantly reduced pupil size in both groups. Controls, but not TMD patients, displayed significant differences in pupil size before compared with after TENS. Under light conditions, TMD patients presented a smaller pupil size compared with controls; the pupil size was reduced only in the controls. Pupil size differences were found before and during TENS and before and after TENS in the controls only. Pupillometry revealed that stimulating the descending opioid pathway with low-frequency sensory TENS of the fifth and seventh pairs of cranial nerves affects the peripheral target. The TMD patients exhibited a different pattern of response to TENS stimulation compared with the controls, suggesting that impaired modulation of the descending pain system may be involved in TMD.

Introduction

Recent literature has suggested that TMD patients may suffer from dysfunction in the brain network that supports sensory, pain, emotional, and cognitive processes [1–8]. Some authors have focused on the dysregulation of the autonomic nervous system (ANS) in TMD patients [9–13], suggesting that TMD could be the clinical manifestation of multisystem dysregulation [14].

Among the structures that are involved in the brain network regulating the sensory, pain, emotional, and cognitive systems, the periaqueductal gray (PAG) has a key role. It receives afferent connections from cortical areas associated with cognition and motivation related to sensory and pain perception [15–17], and it then projects the connections to the centers controlling the peripheral afferent inputs and couples autonomic reactions in a specific manner [18,19].

The role of the PAG in descending pain control via endogenous opioids is one of the most studied pathways [20–22], and several findings suggest that chronic pain is promoted by an abnormal modulation of the descending endogenous pain system [23,24].

Transcutaneous electric nerve stimulation (TENS) has been used for a long time to relieve pain [25–27]. Its main effect is believed to be achieved through the modulation of descending inputs from the ventral-lateral PAG to the rostroventral medial medulla (RVM) [28,29]. In particular, low-frequency TENS is suggested to activate μ opioid receptors [30–32].

Bilateral low-frequency TENS of the fifth and seventh cranial nerves has been suggested to treat TMD [33–36] on the basis of the resulting pain relief [37,38] monitored by surface electromyography (EMG) of masticatory muscles and kinesiography of jaw movements [39,40]. However, despite the clinical relevance of TENS in TMD, the above-cited studies focused on clinical pain or jaw muscle effects. Therefore, no information can be drawn regarding the hypothesis of autonomous or central dysfunction in patients with TMD.

Recent findings promote the use of pupillometry to study the effects of opioids in opioid-maintained patients and healthy subjects [41–43]. Moreover, the dysregulation of the ANS has been reported in TMD [44], obstructive sleep apnea syndrome [45–48], and other autonomous disorders [49–52] by pupillometry. For these reasons, pupillometry can be considered a non-invasive, sensitive and cost-effective tool for investigating, via pupil constriction (miosis), the ANS and opioid-related miosis [53–55].

According to the above-mentioned literature, it is possible that the effects of low-frequency TENS that are mediated by the supraspinal center and the endogenous opioid path can be demonstrated at the pupil level (miosis). The aim of this study was to evaluate pupillometric indexes using low-frequency TENS to test the hypothesis that the opioid descending pathway is dysregulated in patients with TMD.

Materials and Methods

Subjects

This study was conducted in accordance with the Declaration of Helsinki. The Committee on Ethics in Science of the University of L'Aquila, L'Aquila, Italy, approved the study, and written informed consent was obtained from each subject and electronically stored as suggested by our institutional guidelines.

Inclusion/Exclusion Criteria

Eighteen Caucasian patients (mean age 26.5±5.3 years) who fulfilled the following criteria were included in the study group: female gender; age less than 40 years; myogenous TMD; pain

duration longer than 3 months; presence of complete permanent dentition, with the possible exception of the third molars; normal occlusion. Patients were excluded from the study if they met one or more of the following criteria: presence of systemic or metabolic diseases; eye diseases or visual defects; history of local or general trauma; neurological or psychiatric disorders; muscular diseases; cervical pain; bruxism, as diagnosed by the presence of parafunctional facets and/or anamnesis of parafunctional tooth clenching and/or grinding; pregnancy; assumed use of anti-inflammatory, analgesic, anti-depressant, opioid, or myorelaxant drugs; smoking; fixed or removable prostheses; fixed restorations that affected the occlusal surfaces; and either previous or concurrent orthodontic or orthognathic treatment.

For comparison with previous literature [7,8,56,57], the diagnosis of myofascial-type TMD was provided after clinical examination by a trained clinician according to group 1a and 1b of the Research Diagnostic Criteria for TMD (RDC/TMD) [58], in a blinded manner (RC).

The control group consisted of 18 age- and gender-matched Caucasian subjects scheduled for a routine checkup at the University Clinic, without signs or symptoms of TMD, who fulfilled the inclusion and exclusion criteria (mean age 25.1±5.7 years).

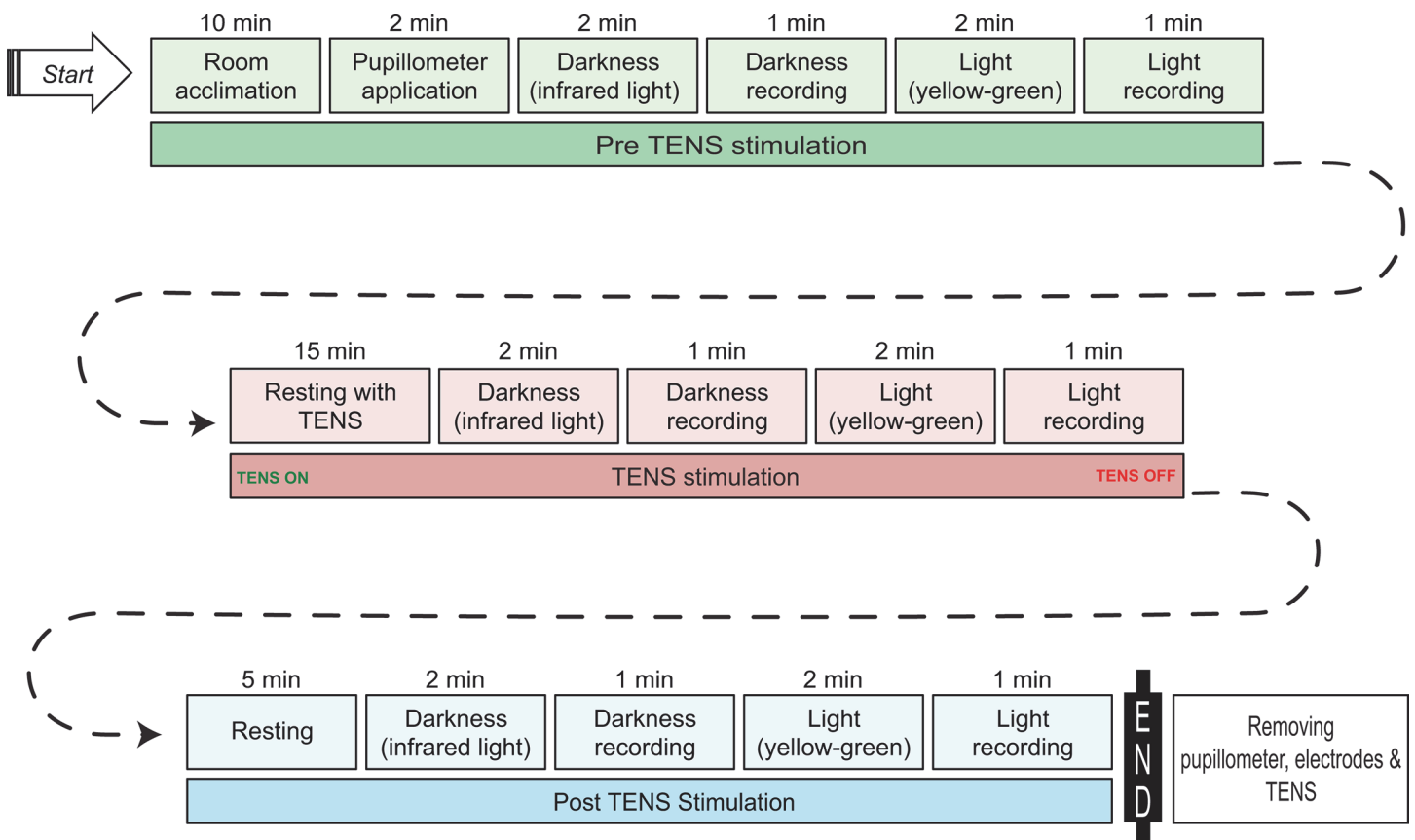Each enrolled subject underwent the experimental protocol described in Fig 1.

Fig 1. Diagram of the Experimental Protocol. Before beginning the experimental session, each enrolled subject was prepared with TENS electrodes and connected to a turned-off TENS device. Subsequently, they underwent three experimental phases: Pre TENS stimulation (green); During TENS stimulation (red); and Post TENS stimulation (blue). Pupillometer application and its removal occurred after the acclimation period and at the end of the experimental session, respectively (green). Based on the literature, central nervous system stimulation was obtained by sensory stimulation of cranial nerves V and VII with low-frequency TENS for 21 minutes (red) [28–32]. To avoid operator interference, the pupillometer, electrodes and TENS device were removed only at the end of the experimental session.

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Pupillometry

Pupillometry was performed with an infrared pupillometer (Oculus system, Inventis srl, Padova, Italy), which was composed of two infrared CCD cameras (resolution of 720x576 pixels, 256 gray levels) mounted on a light helmet (1.5 kg), with a sampling frequency of 25 frames/s. To stabilize accommodation, the subjects were asked to focus their eyes on the light point in the pupillometer [42].

The assessment of pupil size was performed under light conditions, with the eyes illuminated using a yellow-green LED with a 740-nanometer wavelength, as well as under dark conditions that were obtained using only three infrared diodes with a wavelength of 880 nanometers. Pupillometric recordings were acquired in digital form and processed using the Tarjan algorithm designed to evaluate strongly connected components [59] to obtain frame-by-frame measurements of the pupil area, expressed as the number of pixels covering it. A template was positioned on the computer screen to allow for the correction of the eye position to avoid errors in the alignment of the pupils, as previously described [44].

Pupillometry was performed with the subjects in a horizontal supine position on a bed. Room temperature (21°C) and relative humidity (50%) remained constant. Any external or internal noise sources were excluded.

Before the pupillometric recording sessions, patients were invited to lie on the bed for clinical examination with their eyes open for at least 10 minutes to adapt to the temperature and humidity of the room, as well to reduce their anxiety (Fig 1). Then, the pupillometer was applied and maintained until the end of the recording session.

Recording Procedure

All recording procedures are described in the protocol diagram (Fig 1). Briefly, after 2 minutes of darkness under an infrared light condition, 1 minute of recording was obtained, followed by the application of yellow-green light for 2 minutes and subsequent recording for 1 minute; this procedure was applied before, during and after sensory TENS stimulation, for a total of 6 pupillometric recordings of 60 seconds each (Fig 1). The recording sessions were performed by an expert operator (RC) in a blinded manner.

Based on previous literature [44], darkness and light adaptation was minimized, due to the short duration of infrared and yellow-green stimulation, and any potential, residual adaptive effect, if present, could be observed in both groups.

Stimulation Procedure

The method for sensory TENS was described previously [60]. Briefly, a J5 Myomonitor TENS Unit device (Myotronics-Noromed, Inc., Tukwila, WA, USA) with disposable electrodes (Myotrode SG Electrodes, Myotronics-Noromed, Inc., Tukwila, WA, USA) was used. This low-frequency neurostimulator generates a repetitive synchronous and bilateral stimulus delivered at 1.5-s intervals, with an adjustable amplitude of approximately 0–24 mA, a duration of 500 μs, and a frequency of 0.66 Hz. The two TENS electrodes were placed bilaterally over the cutaneous projection of the notch of the fifth pair of cranial nerves, which was located between the coronoid and condylar processes and was retrieved by manual palpation of the zone anterior to the tragus; a third grounding electrode was placed in the center of the back of the neck.

Based on the literature, central nervous system stimulation was obtained by sensory stimulation of cranial nerves V and VII with low-frequency TENS for 21 minutes [28–32]. The amplitude of TENS stimulation started at 0 mA, with the stimulator turned on and the rheostat, which controls the amplitude, positioned at 0. The amplitude of stimulation was progressively increased at a rate of 0.6 mA/s until the patients reported the sensation of pricking. Particular

attention was paid to avoid reaching the threshold of motor stimulation; if any movement of the investigated muscles was observed, the patient was excluded from the study. One patient with TMD and 1 control subject were excluded because the amplitude reached the motor threshold of the seventh cranial nerve, as shown by the contraction of the masseter and temporal muscles.

The same operator (RC) applied the pupillometer and delivered the TENS according to the manufacturer's guidelines.

Statistical Analysis

Statistical analysis was performed using STATA 10 (StataCorp LP, College Station, TX, USA) on average pupil sizes, computed on 60 seconds of recordings, as previously described [44]. Preliminary analysis of pupil size showed, as expected, a high correlation between the left and right pupil sizes, allowing the use of the left-right mean value to simplify the statistics. The ratio between pupil size in the presence of yellow-green light and that in darkness (herein referred to as the L/D ratio) was calculated. The Shapiro-Wilk test revealed a normal distribution of data. Within-group differences in the pupil size and L/D ratio were analyzed using a paired t test, while differences in pupil size between groups were analyzed using an unpaired t test. The level of significance was set at $p = 0.05$ for all tests. The results are expressed as the mean and standard deviation (SD).

Results

The size of the pupil in darkness (infrared light) was significantly reduced during sensory low-frequency TENS both in the control (overall mean of 7886.51 before TENS vs. 7434.33 during TENS; $p = 0.02$) and TMD (7599.88 vs. 7148.94; $p = 0.001$) groups (Table 1). After TENS, the reduction of pupil size remained significant in the control group (7886.51 vs. 7427.11; $p = 0.003$) but not in the TMD group (7599.88 vs. 7490.61; $p = 0.082$) because of the return of pupil size to that prior to TENS.

The unpaired t test did not reveal any significant difference in pupil size between the control and TMD groups before TENS (7886.51 vs. 7599.88; $p = 0.24$), during TENS (7434.33 vs. 7184.94; $p = 0.11$), or after TENS (7427.11 vs. 7490.61; $p = 0.47$). In the TENS condition, it is possible that the higher value of data dispersion in the TMD group (SD 1983.31) was responsible for the lack of significance.

Table 2 shows the statistics of pupil size under yellow-green light. The size of the pupils in the control group was significantly reduced during TENS compared with the size prior to

Table 1. Comparison between the Pupil Size in the Control and TMD Groups in Infrared Light.

	Infrared pupil size Control group	Infrared pupil size TMD group
Before TENS	7886.51 (1231.0)	7599.88 (1254.6)
During TENS	7434.33 (1464.0)*	7148.94 (1983.3)*
After TENS	7427.11 (1168.6)**	7490.61 (1477.6)

The data are expressed as the mean and standard deviation (in parentheses). Asterisks indicate statistically significant differences.

*paired t test within groups between the pre-TENS and TENS conditions: $p = 0.02$ in the control group and $p = 0.001$ in the TMD group.

**paired t test within groups between the pre-TENS and post-TENS conditions in the control group: $p = 0.003$.

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Table 2. Comparison between Pupil Size in the Control and TMD Groups in the Yellow-Green Light Condition.

	Yellow-Green pupil size Control group	Yellow-Green pupil size TMD group
Before TENS	3934.38 (821.35)	2911.51 (1041.97)°
TENS	3523.55 (867.38)*	2836.38 (1184.54)°°
After TENS	3294.16 (617.32)**	2973.83 (1978.37)

The data are expressed as means and standard deviations (in parentheses). Asterisks and circles show the statistically significant differences.

*paired t test within groups between the pre-TENS and TENS conditions in the control group: p = 0.003.

**paired t test within groups between the pre-TENS and post-TENS conditions in the control group: p = 0.0005.

°unpaired t test between the groups before TENS: p = 0.001.

°°unpaired t test between the groups during TENS: p = 0.02.

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TENS (3934.98 vs. 3523.55; p = 0.005), and the pupil size continued to decrease post-TENS (3934.98 vs. 3294.16; p = 0.005). The TMD group did not show significant differences in pupil size between the pre-TENS and TENS conditions (2911.51 vs. 2836.38: p = 0.21) or between the pre-TENS and post-TENS conditions (2911.51 vs. 2973.83: p = 0.34). The unpaired t test between groups showed that the pupil size in the TMD group was significantly lower than that in the controls in the pre-TENS (2911.51 vs. 3934.38: p = 0.001) and TENS (2836.38 vs. 3523.55: p = 0.02) conditions, but no significant differences between groups were found in the post-TENS condition (2973.83 vs. 3294.16: p = 0.14). It is possible that, as in the post-TENS condition in darkness, the higher dispersion of data in the TMD group was responsible for the lack of significance. Note that the dispersion of data in the control group was lower than that in the TMD group, and, in the post-TENS condition, it was approximately one-third of that in the TMD group (sd: 617.32 vs. 1978.37). The absolute value of size differed by 10% (control 3294.16 vs. TMD 2973.83), suggesting more homogeneous pupil behavior among the control subjects.

Results of the L/D ratio within and between groups are shown in Table 3 in the pre-TENS, TENS, and post-TENS conditions. The ratio decreased significantly in the control group

Table 3. Comparison of the Pupil Size and Yellow-Green/Infrared Light (L/D) Ratio between the Control and TMD Groups.

	L/D ratio Control group	L/D ratio TMD group
Before TENS	0.504 (0.111)	0.370 (0.091)°
TENS	0.478 (0.096)*	0.387 (0.123)°°
After TENS	0.444 (0.063)**	0.402 (0.112)

The data are expressed as means and standard deviations (in parentheses). Asterisks and circles show the statistically significant differences.

*paired t test within groups between the pre-TENS and TENS conditions in the control group: p = 0.03.

**paired t test within groups between the pre-TENS and post-TENS conditions in the control group: p = 0.01.

°unpaired t test between the groups before TENS: p = 0.0002.

°°unpaired t test between the groups during TENS: p = 0.01.

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pre-TENS compared with TENS (0.504 vs. 0.478: p = 0.03) and pre-TENS compared with post-TENS (0.504 vs. 0.444: p = 0.01). The TMD group did not show statistically significant differences in this ratio during the within-group comparisons.

The L/D ratio was significantly higher in the control group compared with the TMD group, both pre-TENS (0.504 vs. 0.370: p = 0.0002) and during TENS (0.478 vs. 0.387: p = 0.01). Post-TENS, the L/D ratio remained higher in the control group (0.444 vs. 0.402), but the difference between the two groups was not significant (p = 0.08). Note that the ratio in the TMD group trended toward the value of the control group, with a value less than that in the TENS and pre-TENS conditions.

Discussion

The data in our study can be summarized as follows:

1. No significant difference was found in pupil size in darkness between the control and TMD groups.

2. Significant within-group differences in pupil size in the darkness condition before TENS compared with during TENS in both the control and TMD groups were found; specifically, the pupil size was reduced during TENS.

3. Significant differences in pupil size in the darkness condition were found pre- compared with post-TENS in the control group but not in the TMD group. In the control group, the pupil size was reduced post-TENS, with the same reduction obtained during TENS; in the TMD group, the size of the pupil post-TENS returned close to the pre-TENS value.

4. Significant differences in pupil size in the light condition were found between the control and TMD groups; specifically, the TMD group showed a significantly smaller pupil size.

5. Significant within-group differences in pupil size in the light condition were found pre-TENS compared with during TENS and pre-TENS compared with post-TENS in the control group but not in the TMD group; pupil size in the light condition was reduced only in the control group.

Pupil size is a peripheral effect of a central network that controls the balance between sympathetic and parasympathetic outflow. The ambient light and the physiological and pathological causes of sympathetic excitation and/or parasympathetic inhibition evoke pupil dilation, and, vice versa, causes inducing sympathetic inhibition and/or parasympathetic excitation evoke pupil constriction. For this reason, pupil dynamics have been studied in physiological conditions that are suspected to impair autonomic balance and enhance arousal and sympathetic drive [61,62], enabling the assessment of the relationship between pupil dilation and noradrenergic activity [63,64]. Some authors have focused on the locus coeruleus-noradrenergic neuromodulatory system [65,66], which is considered to be one of the main centers regulating arousal, vigilance, alertness [67–70], pain, and sensory afferents, and it is likely involved in a variety of disorders, such as chronic pain and TMD, characterized by dysfunction of the noradrenergic arousal system [71–73].

The locus coeruleus and the noradrenergic arousal/vigilance system receive opposing regulation by corticotropin-releasing factor and opioids [74,75]. Opioids reduce the activity of the locus coeruleus [76,77] and the diameter of the pupil [41–43,55]. In contrast, a wakefulness-promoting drug, likely acts on the locus coeruleus-noradrenergic system and induces pupil dilation [78,79] (Fig 2).
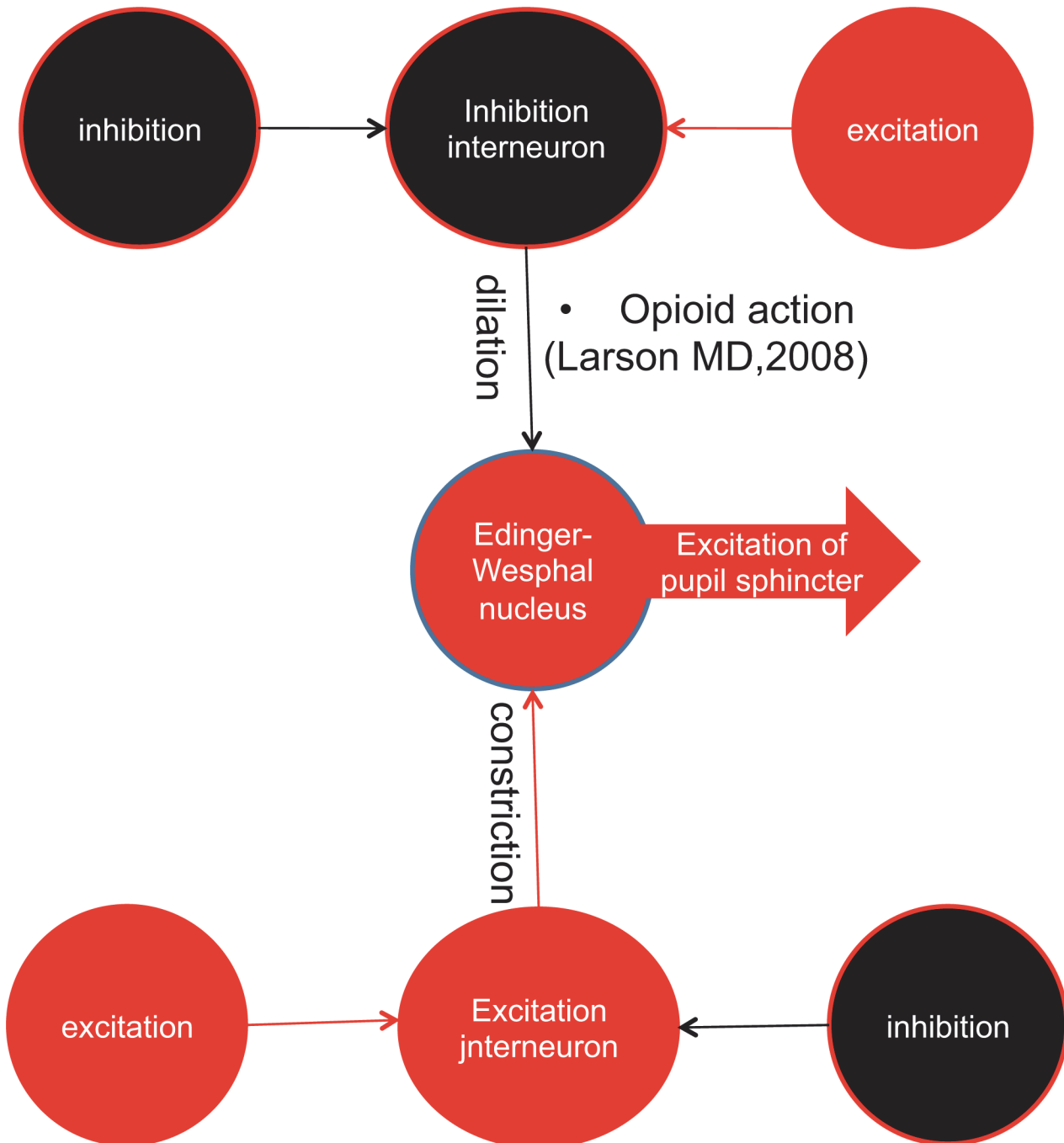
Fig 2. The Edinger-Westphal Nucleus (EWN) Exerts a Tonic Activity on the Pupil Sphincter. Larson [55] proposed that in anesthetized patients, opioid action can be exerted, blocking the efferents of inhibitory neurons to the EWN. Translating this hypothesis to non-anesthetized patients, it could be suggested that opioids reduce the inhibition of interneurons, increasing the effects of excitatory afferents of the EWN. The final effect was opioid miosis. In the figure, excitatory components of the model are shown in red, while inhibitory components are shown in black.

Low-frequency TENS induces a release of endorphins from the ventral-lateral PAG and the activation of the opioid descending path to the RVM. Simultaneously, PAG fibers extend to the pericoeruleus region, in which opioids are released in the dendritic system from the locus coeruleus neurons [80]. For this reason, it is possible that low-frequency TENS induces a reduction

in the firing frequency in the locus coeruleus, simultaneously reducing the state of arousal, the perception of pain, and the diameter (size) of the pupil.

According to the above data, in our study, the pupil size decreased with low-frequency sensory TENS in darkness, both in the control and TMD groups. Five minutes after the end of stimulation, the effect was still present in the control group, but not in the TMD group.

The dilation of the pupil in darkness is mediated by the adrenergic sympathetic branch of the ANS and is supplied by a nerve originating from Budge's cilio-spinal center that excites the dilator muscles of the pupil. The action of the dilator muscles is counterbalanced by the cholinergic parasympathetic branch, which originates from the Edinger-Westphal nucleus and inhibits the dilation. Moreover, pupil dilation obtained by sphincter inhibition can be equal to 1/3 of the maximum physiological dilation [81], and during mental or physical efforts, the pupil dilation may occur via central inhibition of the parasympathetic center [82].

It is possible that in the darkness condition in TMD patients, the endorphin effect due to TENS is sufficient to stimulate the parasympathetic inhibition of the dilator muscle; however, after the stimulation, if the baseline condition is characterized by impaired parasympathetic control, the stimulation provided by TENS rapidly completes its action on the pupillary muscles.

In the light condition, the parasympathetic branch responds to light that activates the sphincter muscle of the iris, causing miosis. At the same time, the sphincter receives beta-adrenergic innervation that is sufficient to reduce and/or counterbalance the contraction [83,84].

In our study, the subjects in the control and TMD groups responded to light as evidenced by a reduction in pupil size, demonstrating the proper activation of the central parasympathetic pathway that controls this basic function. Compared with the control group, the TMD patients showed a significantly higher reduction of pupil size, as indicated by both the absolute values and the ratio between the pupil size in light and in darkness. This finding might reflect impaired counterbalance of adrenergic inhibition or hyperexcitation of parasympathetic effectors in the TMD group. The result that TENS in the light condition reduces the size of the pupil in the control group but has no effect in the TMD group could suggest the impairment of the adrenergic counterbalance system, given that the opioid path works as an inhibitor of the parasympathetic branch of pupil control [55].

Taken together, the data from the present study show that TMD patients suffer from a type of impairment of the descending pathway that controls the sensory, pain, and autonomic output. It is possible that one of the mechanisms involved in this impairment can be addressed in the opioid descending pathway, considering that TMD patients have a weaker response to low-frequency sensory TENS than controls, with respect to modulating the inhibition of physiological parasympathetic excitation. This study focused on adrenergic dysfunction in TMD and confirmed previous studies in which TMD patients showed impairment in the recruitment of adrenergic activity in a weakly stressful task of muscle activation, such as forced voluntarily clenching [44] and other cognitive or emotional tasks [8].

It is probable that the impairments could be due to changes in the connectivity among the structures that belong to the distributed network of pain processing, such as the PAG, which is a node between cortical descending control and midbrain/spinal effectors [56,57,85,86] (Figs 2 and 3).

Confirmation of the involvement of the central descending pain pathway in future studies would explain the observation that therapies such as occlusal splint therapy, orthodontics, and physiotherapy yield short-term pain management success in only 75–80% of cases [87]. However, simple instruments are needed to test the impairment of the descending pathway. Konchak et al. [88] demonstrated that low-frequency TENS of the fifth and seventh pairs of cranial nerves "relaxed" only 75–80% of TMD subjects who received the stimulation. The
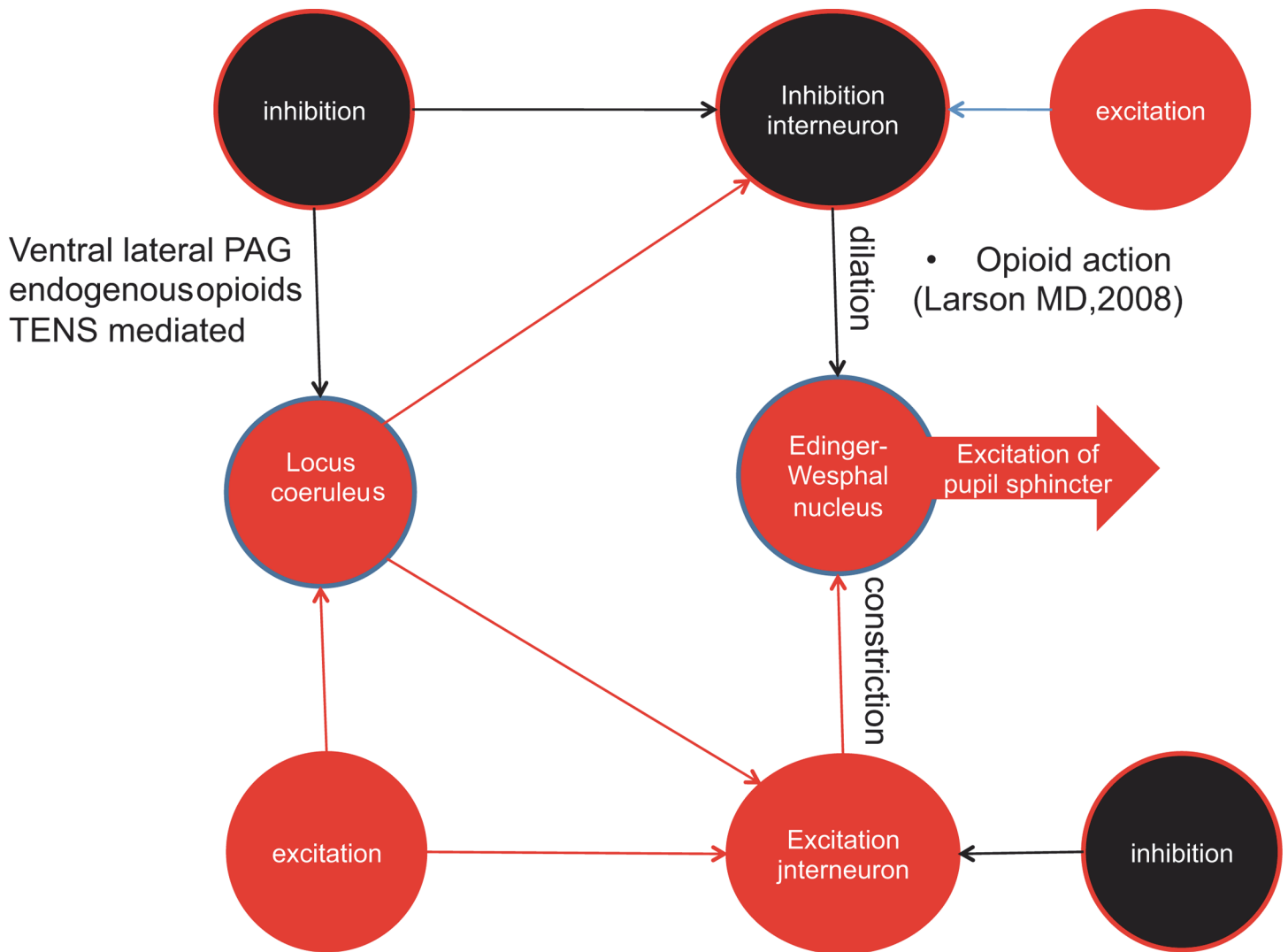
Fig 3. Mechanism of Endogenous Opioid Action Induced by LF TENS in Non-Anesthetized Subjects with Pupil Miosis. The locus coeruleus-noradrenergic system (LC) has been suggested to be a key structure in arousal, and its firing frequency tone has been correlated with wakefulness, vigilance, and sensory and pain perception. The size of the pupil correlates with the activity of LC, and light increases the frequency of firing. The action of the LC excites the excitation pathway that directly induces pupil constriction and excites inhibitory interneurons in the inhibitory pathway, exciting the inhibitor of the inhibitor; the net result is a reduction of inhibition and, indirectly, an agonist effect of the excitatory pathway of pupil constriction. Opioids act on the inhibitory pathway, and TENS has been suggested to be activated via the ventral lateral PAG and the opioid pathway. According to Valentino [74] and Curtis [75], opioids reduce the activation of the LC, and corticotropin-releasing hormone (CRH) increases the firing rate. It is possible that low-frequency sensory TENS inhibits LC inhibitory interneurons, increasing the tone of the pupillary constrictor of the LC. TMD patients could suffer from an impairment of the inhibitory pathway stemming from the ventral lateral PAG. Consequently, pupil constriction in response to light is not counterbalanced by inhibition (smaller pupil size under yellow-green light), and low-frequency sensory TENS is unable to further induce pupil constriction. Red, excitation pathway; black, inhibitory pathway.

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remaining 20–25% showed signs of irritation after TENS, namely, an increase in EMG values of the jaw elevator muscles at rest and a lack of interocclusal freeway space. This result disagrees with the activation of the opioid descending system and directs attention to the activation of the sympathetic pathway, perhaps due to the stimulation of the dorsolateral PAG root.

As in a previous study [88], we used low-frequency TENS of the fifth and seventh pairs of cranial nerves to stimulate the descending control system and to examine the effect on the peripheral target. Beyond the absolute values of pupil size and the L/D ratio, it is noteworthy to mention that the data from TMD subjects are characterized by a higher standard deviation, an

index of the dispersion of the values, compared with the controls. This higher dispersion of TMD data was particularly seen during and after TENS, while the dispersion was reduced in the control subjects. We speculate that low-frequency sensory TENS could aid in differentiating TMD patients with a higher impairment of the descending pain modulation pathway from those with a less impaired system. Unfortunately, we did not assess the response to TENS for the prediction of therapy. Thus, we were unable to determine whether different degrees of impairment, such as a lower light-darkness ratio or greater TENS reduction of the pupil size in the light condition, could predict the therapy outcome. Another limitation of our study is that we did not assess whether TENS induced a reduction in pain perception in the examined subjects; therefore, we could not directly assess the efficacy of our stimulation protocol on experimental pain. However, our goal did not concern the symptom "pain" and the efficacy of low-frequency TENS used in dentistry for its relief, and thus we did not take into account a specific study design to address the response to experimental pain. These topics are beyond the scope of the present work and are suggested for future studies.

Compared with the previous work [44], TMD patients in this study exhibited a smaller pupil size in darkness. The average patient age in the present study was 5.32 years older than that in the previous investigation. It is possible that this age discrepancy is responsible for this difference; in fact, it is known that pupil size decreases with age [49,50]. Moreover, it is possible that the duration of the disorder influences the degree of dysregulation, which shifts from a high sympathetic tone (greater pupil size in darkness) in the early stage of TMD to the failure of sympathetic counterbalance (lower pupil size in darkness) with the progression of the disorder. We did not assess the TMD duration in our sample; however, to assess the possible effect of this variable, the duration of the disorder must be investigated in the future through designed studies.

Conclusion

The stimulation of the descending opioid pathway with low-frequency sensory TENS of the fifth and seventh pairs of cranial nerves and the evaluation of the effect on the peripheral target with a simple pupillometric parameter (pupil size) lead to the conclusion that TMD patients suffer from an impairment in the modulation of the descending pain system. Further studies are necessary to confirm this hypothesis and to relate the impairment of the descending pain system to the therapeutic outcome.

Author Contributions

Conceived and designed the experiments: AM RC DP. Performed the experiments: EO DP RC. Analyzed the data: LM RC. Contributed reagents/materials/analysis tools: AM MG. Wrote the paper: AM RC DP LM.

References

1. Maixner W, Fillingim R, Booker D, Sigurdsson A. Sensitivity of patients with painful temporomandibular disorders to experimentally evoked pain. Pain. 1995; 63: 341–351. PMID: 8719535

2. Maixner W, Fillingim R, Sigurdsson A, Kincaid S, Silva S. Sensitivity of patients with painful temporomandibular disorders to experimentally evoked pain: evidence for altered temporal summation of pain. Pain. 1998; 76: 71–81. PMID: 9696460

3. Svensson P, List T, Hector G. Analysis of stimulus-evoked pain in patients with myofascial temporomandibular pain disorders. Pain. 2001; 92: 399–409. PMID: 11376913

4. Eliav E, Teich S, Nitzan D, El Raziq DA, Nahlieli O, Tal M, et al. Facial arthralgia and myalgia: can they be differentiated by trigeminal sensory assessment? Pain. 2003; 104: 481–490. PMID: 12927620

5. Ayesh EE, Jensen TS, Svensson P. Somatosensory function following painful repetitive electrical stim-ulation of the human temporomandibular joint and skin. Exp Brain Res. 2007; 179: 415–425. PMID: 17146645

6. Schmidt JE, Carlson CR. A controlled comparison of emotional reactivity and physiological response in masticatory muscle pain patients. J Orofac Pain. 2009; 23: 230–242. PMID: 19639103

7. Nebel MB, Folger S, Tommerdahl M, Hollins M, McGlone F, Essick G. Temporomandibular disorder modifies cortical response to tactile stimulation. J Pain. 2010; 11: 1083–1094. doi: 10.1016/j.jpain.2010.02.021 PMID: 20462805

8. Weissman-Fogel I, Moayedi M, Tenenbaum HC, Goldberg MB, Freeman BV, Davis KD. Abnormal cor-tical activity in patients with temporomandibular disorder evoked by cognitive and emotional tasks. Pain. 2011; 152: 384–396. doi: 10.1016/j.pain.2010.10.046 PMID: 21167644

9. Korszun A, Young EA, Singer K, Carlson NE, Brown MB, Crofford L. Basal circadian cortisol secretion in women with temporomandibular disorders. J Dent Res. 2002; 81: 279–283. PMID: 12097314

10. Nilsson AM, Dahlström L. Perceived symptoms of psychological distress and salivary cortisol levels in young women with muscular or disk-related temporomandibular disorders. Acta Odontol Scand. 2010; 68: 284–288. doi: 10.3109/00016357.2010.494620 PMID: 20500119

11. Maixner W, Greenspan JD, Dubner R, Bair E, Mulkey F, Miller V, et al. Potential autonomic risk factors for chronic TMD: descriptive data and empirically identified domains from OPPERA case-contrl study. J Pain. 2011: 12(11 Suppl): T75–T91. doi: 10.1016/j.jpain.2011.09.002 PMID: 22074754

12. Light KC, Bragdon EE, Grewen KM, Brownley KA, Girdler SS, Maixner W. Adrenergic dysregulation and pain with and without acute beta-blockade in women with fibromyalgia and temporomandibular dis-order. J Pain. 2009; 10: 542–552. doi: 10.1016/j.jpain.2008.12.006 PMID: 19411061

13. Eze-Nliam CM, Quartana PJ, Quain AM, Smith MT. Nocturnal heart rate variability is lower in temporo-mandibular disorder patients than in healthy, pain-free individuals. J Orofac Pain. 2011; 25: 232–239. PMID: 21837290

14. Chen H, Nackley A, Miller V, Diatchenko L, Maixner W. Multisystem dysregulation in painful temporo-mandibular disorders. J Pain. 2013; 14: 983–996. doi: 10.1016/j.jpain.2013.03.011 PMID: 23721875

15. Lumb BM. Inescapable and escapable pain is represented in distinct hypothalamic-midbrain circuits: specific roles for Adelta- and C-nociceptors. Exp Physiol. 2002; 87: 281–286. PMID: 11856975

16. Ossipov MH. The perception and endogenous modulation of pain. Scientifica. 2012; 2012: 561761. doi: 10.6064/2012/561761 PMID: 24278716

17. Kobayashi S. Organization of neural systems for aversive information processing: pain, error, and pun-ishment. Front Neurosci. 2012; 6: 136. doi: 10.3389/fnins.2012.00136 PMID: 23049496

18. Bandler R, Keay KA, Floyd N, Price J. Central circuits mediating patterned autonomic activity during ac-tive vs. passive emotional coping. Brain Res Bull. 2000; 53: 95–104. PMID: 11033213

19. Keay KA, Clement CI, Depaulis A, Bandler R. Different representations of inescapable noxious stimuli in the periaqueductal gray and upper cervical spinal cord of freely moving rats. Neurosci Lett. 2001; 313: 17–20. PMID: 11684329

20. Heinricher MM, Tavares I, Leith JL, Lumb BM. Descending control of nociception: specificity, recruit-ment and plasticity. Brain Res Rev. 2009; 60: 214–225. doi: 10.1016/j.brainresrev.2008.12.009 PMID: 19146877

21. Bruehl S, Burns JW, Chung OY, Chont M. Pain-related effects of trait anger expression: neural sub-strates and the role of endogenous opioid mechanisms. Neurosci Biobehav Rev. 2009; 33: 475–491. doi: 10.1016/j.neubiorev.2008.12.003 PMID: 19146872

22. Graeff FG. New perspective on the pathophysiology of panic: merging serotonin and opioids in the peri-aqueductal gray. Braz J Med Biol Res. 2012; 45: 366–375. PMID: 22437485

23. Staud R. Abnormal endogenous pain modulation is a shared characteristic of many chronic pain condi-tions. Expert Rev Neurother. 2012; 12: 577–585. doi: 10.1586/ern.12.41 PMID: 22550986

24. Linnman C, Moulton EA, Barmettler G, Becerra L, Borsook D. Neuroimaging of the periaqueductal gray: state of the field. Neuroimage. 2012; 60: 505–522. doi: 10.1016/j.neuroimage.2011.11.095 PMID: 22197740

25. DeSantana JM, Walsh DM, Vance C, Rakel BA, Sluka KA. Effectiveness of transcutaneous electrical nerve stimulation for treatment of hyperalgesia and pain. Curr Rheumatol Rep. 2008; 10: 492–499. PMID: 19007541

26. Johnson M, Martinson M. Efficacy of electrical nerve stimulation for chronic musculoskeletal pain: a meta-analysis of randomized controlled trials. Pain. 2007; 130: 157–165. PMID: 17383095

27. Sluka KA, Walsh D. Transcutaneous electrical nerve stimulation: basic science mechanisms and clini-cal effectiveness. J Pain. 2003 April 4: 109–121. PMID: 14622708

28. Ainsworth L, Budelier K, Clinesmith M, Fiedler A, Landstrom R, Leeper BJ, et al. Transcutaneous electrical nerve stimulation (TENS) reduces chronic hyperalgesia induced by muscle inflammation. Pain. 2006; 120: 182–187. PMID: 16360266

29. DeSantana JM, Da Silva LF, De Resende MA, Sluka KA. Transcutaneous electrical nerve stimulation at both high and low frequencies activates ventrolateral periaqueductal grey to decrease mechanical hyperalgesia in arthritic rats. Neuroscience. 2009; 163: 1233–1241. doi: 10.1016/j.neuroscience.2009.06.056 PMID: 19576962

30. Sluka KA, Deacon M, Stibal A, Strissel S, Terpstra A. Spinal blockade of opioid receptors prevents the analgesia produced by TENS in arthritic rats. J Pharmacol Exp Ther. 1999; 289: 840–846. PMID: 10215661

31. Kalra A, Urban MO, Sluka KA. Blockade of opioid receptors in rostral ventral medulla prevents antihyperalgesia produced by transcutaneous electrical nerve stimulation (TENS). J Pharmacol Exp Ther. 2001; 298: 257–263. PMID: 11408550

32. Radhakrishnan R, King EW, Dickman JK, Herold CA, Johnston NF, Spurgin ML, et al. Spinal 5-HT(2) and 5-HT(3) receptors mediate low, but not high, frequency TENS-induced antihyperalgesia in rats. Pain. 2003; 105: 205–213. PMID: 14499437

33. Jankelson B. Electronic control of muscle contraction—a new clinical era in occlusion and prosthodontics. Sci Educ Bull. 1969; 2: 29–31. PMID: 4949886

34. Katch EM. Application of transcutaneous electrical nerve stimulation in dentistry. Anesth Prog. 1986; 33: 156–160. PMID: 3488697

35. Cooper BC. International College of Cranio-Mandibular Orthopedics (ICCMO). Temporomandibular disorders: A position paper of the International College of Cranio-Mandibular Orthopedics (ICCMO). Cranio. 2011; 29: 237–244. PMID: 22586834

36. Chipaila N, Sgolastra F, Spadaro A, Pietropaoli D, Masci C, Cattaneo R, et al. The effects of ULF-TENS stimulation on gnathology: the state of the art. Cranio. 2014; 32: 118–130. PMID: 24839723

37. Rodrigues D, Siriani AO, Bérzin F. Effect of conventional TENS on pain and electromyographic activity of masticatory muscles in TMD patients. Braz Oral Res. 2004; 18: 290–295. PMID: 16089258

38. Cooper BC, Kleinberg I. Establishment of a temporomandibular physiological state with neuromuscular orthosis treatment affects reduction of TMD symptoms in 313 patients. Cranio. 2008; 26: 104–117. PMID: 18468270

39. Monaco A, Sgolastra F, Ciarrocchi I, Cattaneo R. Effects of transcutaneous electrical nervous stimulation on electromyographic and kinesiographic activity of patients with temporomandibular disorders: a placebo-controlled study. J Electromyogr Kinesiol. 2012; 22: 463–468. doi: 10.1016/j.jelekin.2011.12.008 PMID: 22245620

40. Kamyszek G, Ketcham R, Garcia R, Radke J. Electromyographic evidence of reduced muscle activity when ULF-TENS is applied to the Vth and VIIth cranial nerves. Cranio. 2001; 19: 162–168. PMID: 11482827

41. Murillo R, Crucilla C, Schmittner J, Hotchkiss E, Pickworth WB. Pupillometry in the detection of concomitant drug use in opioid-maintained patients. Methods Find Exp Clin Pharmacol. 2004; 26: 271–275. PMID: 15319805

42. Fliegert F, Kurth B, Göhler K. The effects of tramadol on static and dynamic pupillometry in healthy subjects—the relationship between pharmacodynamics, pharmacokinetics and CYP2D6 metaboliser status. Eur J Clin Pharmacol. 2005; 61: 257–266. PMID: 15906019

43. Matouskova O, Slanar O, Chytil L, Perlik F. Pupillometry in healthy volunteers as a biomarker of tramadol efficacy. J Clin Pharm Ther. 2011; 36: 513–517. doi: 10.1111/j.1365-2710.2010.01203.x PMID: 21729116

44. Monaco A, Cattaneo R, Mesin L, Ciarrocchi I, Sgolastra F, Pietropaoli D. Dysregulation of the autonomous nervous system in patients with temporomandibular disorder: a pupillometric study. PLOS ONE. 2012; 7: e45424. doi: 10.1371/journal.pone.0045424 PMID: 23028999

45. Monaco A, Cattaneo R, Mesin L, Fiorucci E, Pietropaoli D. Evaluation of autonomic nervous system in sleep apnea patients using pupillometry under occlusal stress: a pilot study. Cranio. 2014; 32: 139–147. PMID: 24839725

46. Wilhelm B, Giedke H, Lüdtke H, Bittner E, Hofmann A, Wilhelm H. Daytime variations in central nervous system activation measured by a pupillographic sleepiness test. J Sleep Res. 2001; 10: 1–7. PMID: 11285049

47. Wilhelm B, Wilhelm H, Widmaier D, Ludtke H, Ruhle K. Therapy control in sleep apnea patients by means of a pupillographic sleepiness test. Sleep. 1998; 21: 266.

48. Yamamoto K, Kobayashi F, Hori R, Arita A, Sasanabe R, Shiomi T. Association between pupillometric sleepiness measures and sleep latency derived by MSLT in clinically sleepy patients. Environ Health Prev Med. 2013; 18: 361–367. doi: 10.1007/s12199-013-0331-0 PMID: 23420264

49. Bär KJ, Schulz S, Koschke M, Harzendorf C, Gayde S, Berg W, et al. Correlations between the autonomic modulation of heart rate, blood pressure and the pupillary light reflex in healthy subjects. J Neurol Sci. 2009; 279: 9–13. doi: 10.1016/j.jns.2009.01.010 PMID: 19195664

50. Keivanidou A, Fotiou D, Arnaoutoglou C, Arnaoutoglou M, Fotiou F, Karlovasitou A. Evaluation of autonomic imbalance in patients with heart failure: a preliminary study of pupillomotor function. Cardiol J. 2010; 17: 65–72. PMID: 20104459

51. Jain S, Siegle GJ, Gu C, Moore CG, Ivanco LS, Jennings JR, et al. Autonomic insufficiency in pupillary and cardiovascular systems in Parkinson's disease. Parkinsonism Relat Disord. 2011; 17: 119–122. doi: 10.1016/j.parkreldis.2010.11.005 PMID: 21115264

52. Muppidi S, Adams-Huet B, Tajzoy E, Scribner M, Blazek P, Spaeth EB, et al. Dynamic pupillometry as an autonomic testing tool. Clin Auton Res. 2013; 23: 297–303. doi: 10.1007/s10286-013-0209-7 PMID: 23880969

53. Gutstein HB, Akil H. Opioid analgesics. In: Brunton LL, Lazo JS, Parker KL, editors. Goodman and Gilman's the pharmacological basis of therapeutics. 11th ed. New York: McGraw-Hill; 2006. pp. 547–590.

54. Verster JC, Veldhuijzen DS, Volkerts ER. Effects of an opioid (oxycodone/paracetamol) and an NSAID (bromfenac) on driving ability, memory functioning, psychomotor performance, pupil size, and mood. Clin J Pain. 2006; 22: 499–504. PMID: 16772806

55. Larson MD. Mechanism of opioid-induced pupillary effects. Clin Neurophysiol. 2008; 119: 1358–1364. doi: 10.1016/j.clinph.2008.01.106 PMID: 18397839

56. Ichesco E, Quintero A, Clauw DJ, Peltier S, Sundgren PM, Gerstner GE, et al. Altered functional connectivity between the insula and the cingulate cortex in patients with temporomandibular disorder: a pilot study. Headache. 2012; 52: 441–454. doi: 10.1111/j.1526-4610.2011.01998.x PMID: 21929661

57. Younger JW, Shen YF, Goddard G, Mackey SC. Chronic myofascial temporomandibular pain is associated with neural abnormalities in the trigeminal and limbic systems. Pain. 2010; 149: 222–228. doi: 10.1016/j.pain.2010.01.006 PMID: 20236763

58. Dworkin SF, LeResche L. Research diagnostic criteria for temporomandibular disorders: review, criteria, examinations and specifications, critique. J Craniomandib Disord. 1992; 6: 301–355. PMID: 1298767

59. Tarjan R. Depth-first search and linear graph algorithms. SIAM J Comput. 1972; 1: 146–160.

60. Monaco A, Sgolastra F, Pietropaoli D, Giannoni M, Cattaneo R. Comparison between sensory and motor transcutaneous electrical nervous stimulation on electromyographic and kinesiographic activity of patients with temporomandibular disorder: a controlled clinical trial. BMC Musculoskelet Disord. 2013; 14: 168. doi: 10.1186/1471-2474-14-168 PMID: 23672400

61. Hayashi N, Someya N, Fukuba Y. Effect of intensity of dynamic exercise on pupil diameter in humans. J Physiol Anthropol. 2010; 29: 119–122. PMID: 20558970

62. Ishigaki H, Miyao M, Ishihara S. Change of pupil size as a function of exercise. J Hum Ergol. 1991; 20: 61–66. PMID: 1820381

63. Preuschoff K, 't Hart BM, Einhäuser W. Pupil dilation signals surprise: evidence for noradrenaline's role in decision making. Front Neurosci. 2011; 5: 115. doi: 10.3389/fnins.2011.00115 PMID: 21994487

64. Gabay S, Pertzov Y, Henik A. Orienting of attention, pupil size, and the norepinephrine system. Atten Percept Psychophys. 2011; 73: 123–129. doi: 10.3758/s13414-010-0015-4 PMID: 21258914

65. Murphy PR, O'Connell RG, O'Sullivan M, Robertson IH, Balsters JH. Pupil diameter covaries with BOLD activity in human locus coeruleus. Hum Brain Mapp. 2014; 35: 4140–4154. doi: 10.1002/hbm.22466 PMID: 24510607

66. Alnaes D, Sneve MH, Espeseth T, Endestad T, van de Pavert SHP, Laeng B. Pupil size signals mental effort deployed during multiple object tracking and predicts brain activity in the dorsal attention network and the locus coeruleus which represents one of the most important center of arousal. J. Vision. 2014; 14: 1–20.

67. Aston-Jones G. Brain structures and receptors involved in alertness. Sleep Med. 2005; 6(Suppl 1): S3–S7. PMID: 16140243

68. Aston-Jones G, Cohen JD. Adaptive gain and the role of the locus coeruleus-norepinephrine system in optimal performance. J Comp Neurol. 2005; 493: 99–110. PMID: 16254995

69. Stone EA, Lin Y, Sarfraz Y, Quartermain D. The role of the central noradrenergic system in behavioral inhibition. Brain Res Rev. 2011; 67: 193–208. doi: 10.1016/j.brainresrev.2011.02.002 PMID: 21315760

70. Berridge CW, Schmeichel BE, España RA. Noradrenergic modulation of wakefulness/arousal. Sleep Med Rev. 2012; 162: 187–197.

71. Berridge CW, Waterhouse BD. The locus coeruleus-noradrenergic system: modulation of behavioral state and state-dependent cognitive processes. Brain Res Brain Res Rev. 2003; 42: 33–84. PMID: 12668290

72. Szabady E. Modulation of physiological reflexes by pain: role of the locus coeruleus. Front Inegr Neurosci. 2012; 6: 94.

73. Brightwell JJ, Taylor BK. Noradrenergic neurons in the locus coeruleus contribute to neuropathic pain. Neuroscience. 2009; 160: 174–185. doi: 10.1016/j.neuroscience.2009.02.023 PMID: 19223010

74. Valentino RJ, Van Bockstaele E. Opposing regulation of the locus coeruleus by corticotropin-releasing factor and opioids. Potential for reciprocal interactions between stress and opioid sensitivity. Psychopharmacology. 2001; 158: 331–342. PMID: 11797054

75. Curtis AL, Bello NT, Valentino RJ. Evidence for functional release of endogenous opioids in the locus ceruleus during stress termination. J Neurosci. 2001; 21: RC152. PMID: 11406637

76. Mazei-Robinson MS, Nestler EJ. Opiate-induced molecular and cellular plasticity of ventral tegmental area and locus coeruleus catecholamine neurons. Cold Spring Harb Perspect Med. 2012; 2: a012070. doi: 10.1101/cshperspect.a012070 PMID: 22762025

77. Valentino RJ, Van Bockstaele E. Convergent regulation of locus coeruleus activity as an adaptive response to stress. Eur J Pharmacol. 2008; 583: 194–203. doi: 10.1016/j.ejphar.2007.11.062 PMID: 18255055

78. Hou RH, Freeman C, Langley RW, Szabadi E, Bradshaw CM. Does modafinil activate the locus coeruleus in man? Comparison of modafinil and clonidine on arousal and autonomic functions in human volunteers. Psychopharmacology. 2005; 181: 537–549. PMID: 15983798

79. Nikolaou A, Schiza SE, Giakoumaki SG, Roussos P, Siafakas N, Bitsios P. The 5-min pupillary alertness test is sensitive to modafinil: a placebo controlled study in patients with sleep apnea. Psychopharmacology. 2008; 196: 167–175. PMID: 17899016

80. Ishimatsu M, Williams JT. Synchronous activity in locus coeruleus results from dendritic interactions in pericoerulear regions. J Neurosci. 1996; 16: 5196–5204. PMID: 8756448

81. Tachado SD, Akhtar RA, Abdel-Latif AA. Activation of beta-adrenergic receptors causes stimulation of cyclic AMP, inhibition of inositol trisphosphate, and relaxation of bovine iris sphincter smooth muscle. Biochemical and functional interactions between the cyclic AMP and calcium signalling systems. Invest Ophthalmol Vis Sci. 1989; 30: 2232–2239. PMID: 2551839

82. Steinhauer SR, Siegle GJ, Condray R, Pless M. Sympathetic and parasympathetic innervation of pupillary dilation during sustained processing. Int J Psychophysiol. 2004; 52: 77–86. PMID: 15003374

83. Topalkara A, Karadas B, Toker MI, Kaya T, Durmus N, Turgut B. Relaxant effects of beta-adrenoceptor agonist formoterol and BRL 37344 on bovine iris sphincter and ciliary muscle. Eur J Pharmacol. 2006; 548: 144–149. PMID: 16973159

84. Patil PN, Weber PA. In vivo functional implications of isoproterenol-mediated relaxation of isolated human iris sphincter. J Ocul Pharmacol. 1991; 7: 297–300. PMID: 1809789

85. Ploner M, Lee MC, Wiech K, Bingel U, Tracey I. Flexible cerebral connectivity patterns subserve contextual modulations of pain. Cereb Cortex. 2011; 21: 719–726. doi: 10.1093/cercor/bhq146 PMID: 20713505

86. Ploner M, Lee MC, Wiech K, Bingel U, Tracey I. Prestimulus functional connectivity determines pain perception in humans. Proc Natl Acad Sci U S A. 2010; 107: 355–360. doi: 10.1073/pnas.0906186106 PMID: 19948949

87. Suvinen TI, Reade PC, Kemppainen P, Könönen M, Dworkin SF. Review of aetiological concepts of temporomandibular pain disorders: towards a biopsychosocial model for integration of physical disorder factors with psychological and psychosocial illness impact factors. Eur J Pain. 2005; 9: 613–633. PMID: 15978854

88. Konchak PA, Thomas NR, Lanigan DT, Devon RM. Free way space measurement using mandibular kinesiograph and EMG before and after TENS. Angle Orthod. 1988; 58: 343–350. PMID: 3264667