Influence of Laser Therapy and Muscle Relaxant on the Masseter Muscle under Occlusal Wear - An Ultrastructural Study

In the context of summation, the purpose of this study was to investigate the influence of low-intensity laser therapy and muscle relaxant in the characteristic ultra structural masseter muscle occlusal wear. Animals and Methods: 40 male Wistar rats were randomly divided into four groups: the control group (GI), occlusal wear (G-II), laser occlusal wear (G-III), and the muscle relaxant occlusal wear (G-IV). Under general anesthesia given intraperitoneally, animals in groups II, III and IV had unilateral amputation of upper and lower molar cusps to simulate an occlusal wear situation. The masseter muscle G-III received laser therapy (830nm, 4J/cm², 40mW, f ~ 2mm) and the procedure was subsequently repeated every other day for 14/30 days. G-IV animals were treated with daily injection of dantrolene (2.5 mg / kg in 0.5 ml of H2O). From 24 hours after the elimination peak, the animals were euthanized with an overdose of anesthesia on days 14 and 30 after the removal of the cusps and the ipsilateral masseter muscle was excised and divided in two, one half was routinely processed for light microscopy and other for electron microscopy. There was no statistical difference between each experimental group and the control and between periods in each experimental group. However, the muscle fibers in the G-II showed the most pronounced changes. There is no causal relationship between muscles fibers injuries and occlusion and, despite signs of muscular tissue injury were more evident in the occlusal wear group. Results indicates a moderate action of laser therapy and muscle relaxants in skeletal muscle.

KEY WORDS: Lasertherapy; Ultrastructure; Dental occlusion; Masseter muscle relaxant.

INTRODUCTION

The etiology of temporomandibular disorder (TMD) involves craniofacial morphology, head posture, stress, psychological factors, trauma, joint hyper mobility and infections as risk or contributing factors (Egelmark et al., 2003; Liljeström et al., 2005; Niemi et al., 2006; Prasad et al., 2007). In addition, the occlusion plays a key role on the temporomandibular joint as it may influence the functional activity of masticatory muscles, resulting in fatigue and muscular pain (Gesh et al., 2004; Sonnesen & Svensson, 2008). Despite this, previous studies focusing on muscle activity in TMD patients have not provided convincing evidence that muscle hyperactivity occurs (Issa et al., 2006; Okano et al., 2007; Rodrigues & Ferreira, 2003; Suvinen & Kemppainen, 2007). Thus, the relationship between malocclusion and muscle morphofunctional aspects in the masticatory system is not yet fully understood (Bani et al., 1999).

Due to these aspects, several modalities of therapy have been approached for patients with temporomandibular joint (TMJ) dysfunction and chronic orofacial pain (Türp et al., 2007), in order to maximize their complementary actions. Among the therapies currently in use, muscle relaxants and low level laser therapy (LLL T) have been reported (Bani & Bergamini, 2001; Carrasco et al., 2008; Emshoff et al., 2008). Muscle relaxants are prescribed to reduce skeletal muscle tone and chronic orofacial pain, and alleviate muscle hyperactivity (Bani & Bergamini, 2011; Bani & Bergamini, 2002; Hersh et al., 2008; Krause et al., 2004).
LLLT is known as a new therapeutic approach used for TMD treatment. It stimulates the microcirculation, acts on tissue repair to reduce edema and pain (Mazzeto et al., 2007; Shinozaki et al., 2006). In addition, LLLT keeps the intensity of the force during muscle contraction and may delay its fatigue, and this may increase intracellular ATP levels sufficient to maintain muscle physical effort (Lopes-Martins et al., 2006). However, Gam et al. (1993) have not shown efficiency of LLLT on musculoskeletal disorders. Therefore, considering that the biological mechanisms resulting from action of LLLT on temporomandibular joint (TMJ), clinical trials remains unclear (Çetiner et al., 2006; Fikáková et al., 2007; Gorgey et al., 2008; Tullberg et al., 2003; Venancio et al., 2005).

The aim of the present study was to analyze the influence of laser therapy and muscle relaxant on the ultrastructural features of masseter muscle under occlusal wear.

MATERIAL AND METHOD

After approval by the Ethics Committee (Protocol 019/06), forty male adult Wistar rats weighing 300± 350g were included in the study and housed at the animal house of the School of Dentistry of the Federal University of Bahia. The animals were kept on 12:12 h light/dark cycle and were fed with standard pelleted diet and had water ad libitum. The animals were randomly divided into 4 groups with 10 animals each: control group (G-I), occlusal wear group (G-II), occlusal wear + LLLT (G-III), and occlusal wear + muscle relaxant (G-IV).

Under intraperitoneal general anesthesia (ketamine, 0.10mL/100g + xylazine, 0.25 mL/100mg), animals of groups II, III and IV had unilateral amputation of the left inferior and up molar cusps with a round diamond bur under saline irrigation in order to simulate an occlusal wear situation. Care was taken not to cause pulp exposure or damage.

The animals of G-III had the left masseter muscle region shaved and received LLLT (1830nm, 4J/cm², 40mW, f ~ 2mm spot, ‘Kondortech’ Laser Unit - São Paulo/Brazil) in two contact points, 1 cm apart, 24 h after the procedure and it was repeated at every other day during 14 days for half the animals and during 30 days on the other half. Animals of G-IV were treated with a daily intraperitoneal injection of Dantrolene® (Cristália, São Paulo/Brazil, 2.5 mg/Kg in 0.5 ml of H₂O) beginning 24 hours after occlusal wear simulation.

Half of the animals of each group were killed by overdose of general anesthetics on days 14 and 30 after cusps removal. The ipsilateral masseter muscle of the side of amputated cusps were excised and divided into two samples, being one half routinely processed for electron microscopy.

For transmission electron microscopy, the fresh samples were fixed in 4% cold glutaraldehyde, containing 0.1 M (0.1 mol/l, PBS, pH 7.2) sodium cacodylate buffer, at room temperature, and post-fixed in 1% osmium tetroxide containing 0.1 M (0.1 mol/l, pH 7.2) phosphate buffer during one hour, dehydrated in graded acetone, and embedded with Polybed resin. Ultra-thin sections (70nm) were cut and stained with uranyl and lead citrate for Zeiss-EM109 transmission electron microscope observation.

For the ultrastructural analysis, the samples were examined by a single previously trained observer using that microscope at magnifications of 4400x, 12000x and 20000x. The morphological features were established by the observation of these changes in three representative fields (each magnification) in the masseter muscle of each animal (adapted from Bani et al., 1998). The criteria used are reported in the Table I.

Data were reported as mean ± SEM. The difference between groups were tested using statistically Kruskall Wallis, Mann Whitney, and paired Wilcoxon test. All statistical calculations were performed using Minitab Program, 14.0 software (Minitab, Belo Horizonte, Minas Gerais, Brazil). The level of significance was set at 5% (P< 0.05).

Table I. Scoring method of tissue injury.

<table>
<thead>
<tr>
<th>SCORE</th>
<th>DEGREE OF MORPHOLOGICAL FEATURES</th>
</tr>
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<tbody>
<tr>
<td>0</td>
<td>Normal fiber</td>
</tr>
<tr>
<td></td>
<td>Vacuoles</td>
</tr>
<tr>
<td>1</td>
<td>Slight</td>
</tr>
<tr>
<td></td>
<td>mitochondrial swelling intracellular edema</td>
</tr>
<tr>
<td>2</td>
<td>Moderate</td>
</tr>
<tr>
<td></td>
<td>vacuoles</td>
</tr>
<tr>
<td></td>
<td>mitochondrial swelling</td>
</tr>
<tr>
<td></td>
<td>mitochondria with loss of cristae</td>
</tr>
<tr>
<td></td>
<td>intracellular edema</td>
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<tr>
<td></td>
<td>vacuoles</td>
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<tr>
<td></td>
<td>mitochondrial swelling</td>
</tr>
<tr>
<td></td>
<td>mitochondrial with loss of cristae</td>
</tr>
<tr>
<td></td>
<td>disarrangement of myofibrils</td>
</tr>
<tr>
<td>2</td>
<td>Severe</td>
</tr>
</tbody>
</table>

RESULTS

Group I (control) - This group showed muscle fibers with a normal aspect. The organelles exhibited conspicuous mitochondrias usually arranged duplets at Zine line. The sarcoplasmatic reticulum was well developed, and occasional cytoplasmatic vacuoles were also observed (Fig. 1).

Group II - This group showed many cytoplasmatic vacuoles exhibiting a small size. They were especially numerous in some muscle fibers, however, other fibers did not show to be affected (Fig. 2). The percentage of vacuolated fibers was significantly increased when compared to the control group. In addition, clusters of muscle fibers were also found with disarrangement of myofibrils, hypercontraction and swollen mitochondria with disrupted crista and cleared matrix (Fig. 3). The occlusion wear group (II) at 14 days presented severe injury in 20% of the fields, 60% with light or moderate injury, and 20% without injury. At 30 days, severe injury was only seen in 6.6% of the fields, with 53.4% presenting light or moderate injury. The muscle fibers in group II showed more pronounced alterations than those of rats without molar amputation in group I. Besides these characteristics, there was no significant difference between group I and II or between the periods in the groups (P>0.05).

Group III and Group IV - These groups showed few of the above alterations. Only sparse muscle fibers showed moderate mitochondrial swelling, whereas myofibril hypercontraction was rarely observed (Fig. 4 and 5). In these groups, moderate and severe scores were not as evident as in group II. There were also no statistical differences between these groups and between the experimental periods in each group.

Despite the particular characteristics in each group, the Kruskal-Wallis test demonstrated no interaction between groups I, II, III, IV (P= 0.251). Paired analysis with Wilcoxon test found no difference between 14 and 30 days on Group II (P=0.2718), Group III (P=1.000), Group IV (P=0.8241). The score for the electron microscopic examination of specimens are in Fig 6.
DISCUSSION

Little is known about the association between occlusion and TMD. This study attempted to analyze the morphological features of masseter muscle under occlusal wear and treated with LLLT or muscle relaxant therapy, as no therapy has shown itself capable of overcoming the others.

We used a model of occlusion wear to simulate an altered occlusion. This situation represented patients with lost or worn teeth. This model has been used in previous studies (Bani et al.; Bani & Bergamini, 2001; Nogueira-Filho et al., 2004) resulting in different degrees of morphologic and physiologic alterations on masticatory muscles, specially masseter. It is important to state that malocclusion may influence the functional performance of the masticatory muscles, resulting in muscular overwork and fatigue (Glaros et al., 2007; Pereira et al., 2009; Van Selms et al., 2008).

Previous studies using experimental animals have shown ultrastructural alterations in the masseter muscle related to occlusal wear (Bani & Bergamini, 2002; Santiwong et al., 2002) and pterygoid (Bazan et al., 2008; Iyomasa et al., 2008). Some authors as Iyomasa et al. and Bani et al. have shown that muscle fibers and capillaries are sensitive to different masticatory loads, indicating that this may contribute to understand the way in which the muscles adapt to different loads. Although, the group with occlusal wear showed ultra-structural changes at the two experimental periods in this study, they were not enough to demonstrate statistical difference between control group and occlusion wear group.

Fig. 3 Rats with molar amputation after 14 days: hypercontracted muscle fiber with disappearance of myofibril I bands, disorganization of myofibrils and disarrangement of contractile filaments (x20,000).

Fig. 4. In the laser group few alterations of muscle tissue could be found. Only sparse muscle fibers showed moderate mitochondrial swelling, although the myofibrils with thick Z lines; mitochondria located in pairs at the level of the Z lines and well-developed sarcoplasmatic reticulum. (x 4,400).
results it seemed to play positive effects on muscle tissue as important injuries were not observed at group III. Furthermore, although the findings of the present study were restricted to a specific set of parameters and that optimal treatment parameters (e.g., wavelength, dosage, number of treatment sessions) have not been, so far, agreed upon (Emshorff et al.; Venancio et al.), previous studies have been beneficial results using Infrared wavelengths (Çetiner et al.; Fikáková et al.; Kato et al., 2006; Kogawa et al., 2005; Leal Júnior et al., 2008).

The laser group (III) showed mild ultrastructural alterations, similar to control group. In this group, only 20% of fields had filled the criteria for severe and moderate injury. There were no statistical differences between the experimental periods in this group. These findings are similar to previous studies that demonstrated beneficial action of the laser on muscular TMD (Shinozaki et al.; Amaral & Salvini, 2001; Costardi et al., 2008; Farias, 2005; Frare & Nicolau, 2008). Despite the laser therapy role on muscle tissues, some authors have performed some studies. Lopes-Martins et al. investigated whether LLLT on visible spectra (655 nm) can reduce muscular fatigue during tetanic contractions in rats, and concluded that LLLT doses of 0.5 and 1.0 J/cm² can prevent development of muscular fatigue in rats during repeated tetanic contractions. Using infrared spectrum, Kato et al. and Kogawa et al. were successful in reducing pain in patients with TMD and increasing active range of motion on the masticatory muscles.

The group IV with dantrolen showed mild ultrastructural

Fig. 5. In the muscle relaxant group few alterations of muscle tissue could be found. The myofibrils with thick Z lines; mitochondria located in pairs at the level of the Z lines and well-developed sarcoplasmatic reticulum. Normal aspect (×20.000).

Fig. 6 Injury score for the electron microscopic examination

Therapies used on groups III and IV represented two ways for treatments for TMD. They are reversible therapies mentioned in previous reports as the initial treatment for TMD (Bani & Bergamini, 2001; Fikáková et al.; Naikmasur et al., 2008), and are effective in reducing pain and muscular tension. The laser protocol used in the present study was chosen in view to its ability to penetrate the muscle tissue, reducing pain and increasing active range of motion and according to our
alterations, also similar to control group. In this group, only 10% of fields had filled the criteria for moderate and severe injury and 40% of fields did not show signs of injury. Based on these features, it is possible to state that administration of skeletal muscle relaxant may prevent the occurrence of ultra-morphologic alterations. However, as there were no statistical differences between groups I, II, II and IV, other features especially those related to muscle biochemistry should be further investigated (Lisboa et al., 2010).

Bani et al. studied masseter muscle on a rodent model similar to ours and found vasoconstriction and damages on muscle fibers of animals corresponding to the occlusal wear group. Two years later, Bani & Bergamini (2001) carried out a study with the same animal model and used dantrolen (10 mg/kg b.w.) as therapy for muscle alterations. These authors demonstrated that the sub-cellular features were similar between control group and maloccluded rats given dantrolen group. In addition in their study, control group and maloccluded rats group given dantrolen showed results statistically different from those observed in maloccluded rat groups not given dantrolen. This medicine is not usually used for TMJ; thus, it has not been established dose for this purpose yet. The dose of 2.5mg/kg b.w used in our experiment represents that earlier dose used for malignant hyperthermia, and is four times smaller than that used by other authors (Bani & Bergamini, 2001). It may not be enough, as there were no significant differences in our study, however, the ultra-structural characteristics of masseter muscle observed here and in that study carried out by Bani & Bergamini (2001) showed normal appearance. This might suggest that this medicine can be a new tool focusing on the TMD. In conclusion, it is possible that LLLT and skeletal muscle relaxant play a positive influence on the masseter muscle in rats under occlusal wear, as morphological changes were more evident in occlusal wear group. Thus, our results did not show a cause effect relation between occlusion and TMD, although signs of muscle tissue injury were more evident in that group, indicating moderate action of LLLT and skeletal muscle relaxant as therapeutic agents.

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Correspondence to: Jean Nunes dos Santos
Av. Araujo Pinho, 62
Canela, Salvador, 40110-150
Bahia
BRAZIL.

Phone: +55 71 3283 9019
Fax : +55 71 3283 8962
E-mail: jeannunes@ufba.br

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