ORIGINAL ARTICLE

Treatment of orofacial pain in patients with stylomandibular ligament syndrome (Ernest syndrome)☆

D. Peñarrocha-Oltraa, J. Ata-Alia,b,*, F. Ata-Alia, M.A. Peñarrocha-Diagoa, M. Peñarrochaa

a Facultad de Medicina y Odontología, Universidad de Valencia, Valencia, Spain
b Equipo de Atención Primaria, Agencia Valenciana de Salud, Valencia, Spain

Received 8 February 2012; accepted 30 June 2012
Available online 5 July 2013

Abstract

Introduction: Ernest syndrome involves the stylomandibular ligament. It is characterised by pain in the preauricular area and mandibular angle, radiating to the neck, shoulder, and eye on the same side, and associated with pain during palpation of that ligament. The purpose of this study is to describe the clinical characteristics, treatment, and course of the disease in a series of patients with Ernest syndrome.

Methods: Retrospective observational study covering the period from 1998 to 2008. We recorded patients’ age, sex, duration of the disorder, and pain characteristics. All patients were injected with 40 mg triamcinolone acetonide at the mandibular insertion of the stylomandibular ligament.

Results: The study included a total of 6 patients. Mean age was 40.3 years (range, 35–51). All the subjects were women. Four patients had undergone lengthy dental treatments in the month prior to onset of the pain. The mean time between pain onset and first consultation was 23 months. The syndrome resolved completely in all cases after treatment, with a minimum follow-up period of 12 months.

Conclusions: We analysed the clinical characteristics, treatment, and course of disease in 6 patients with Ernest syndrome. Correct diagnosis is the key to being able to provide proper treatment. This disorder is sometimes confused with other types of orofacial pain, and may therefore be more prevalent than the literature would indicate.

© 2012 Sociedad Española de Neurología. Published by Elsevier España, S.L. All rights reserved.

Keywords

Ernest syndrome; Myofascial pain; Stylomandibular; Eagle syndrome; Orofacial pain

PALABRAS CLAVE

Síndrome de Ernest; Dolor miofascial; Estilomandibular; Síndrome de Eagle; Dolor orofacial

Tratamiento del dolor orofacial en pacientes con síndrome del ligamento estilomandibular (síndrome de Ernest)

Resumen

Introducción: El síndrome de Ernest se define como una alteración del ligamento estilomandibular, caracterizado por la presencia de dolor en la región preauricular y en el ángulo mandibular, irradiado al cuello, el hombro y el ojo del mismo lado, asociado a dolor durante la


* Corresponding author.
E-mail address: javiataali@hotmail.com (J. Ata-Ali).

2173-5808/$ - see front matter © 2012 Sociedad Española de Neurología. Published by Elsevier España, S.L. All rights reserved.
Introduction

Ernest syndrome was described by Dr Ernest in 1981 and a published article appeared one year later; Shankland et al. were the first to publish a case series in the PubMed database. Ernest syndrome affects the stylomandibular ligament. It is characterised by the presence of pain in the preauricular region and angle of the mandible radiating to the neck, shoulder, and eye on the same side, and it is associated with pain upon palpation of that ligament. Since 1987 when Shankland et al. described 68 patients with Ernest syndrome, no new case series have been published in the medical literature.

A good medical history, manual examination of the insertion of the stylomandibular ligament in the gonial angle, and immediate relief of symptoms after administration of local anaesthesia to this region will confirm this diagnosis.

The current article presents 6 additional cases with masticatory orofacial pain that were diagnosed with the Ernest syndrome and discusses the symptoms, treatment, and course of the disease.

Patients and methods

This retrospective observational study conducted between 1998 and 2008 examined patients who visited the oral surgery unit of the dental clinic pertaining to the University of Valencia’s School of Medicine and Dentistry due to orofacial pain. The following inclusion criteria were used to identify cases of Ernest syndrome: (a) presence of pain in the preauricular region and angle of the mandible; (b) pain upon palpation of the mandibular insertion of the stylomandibular ligament; (c) pain relief after injection of mepivacaine 3% (Normon Laboratory, Madrid, Spain) in the mandibular insertion of the painful stylomandibular ligament, and (d) no significant changes in the extraoral panoramic radiography.

We recorded data for age, sex, progression timeline, and pain characteristics. We also searched for any traumatic episodes or prolonged use of a dental treatment that may have proceeded onset of pain. Patients were considered to have a positive history when they had undergone treatment in the month prior to the onset of pain. We examined each patient’s oral cavity, muscles of mastication, stylomandibular ligament (Fig. 1), and temporomandibular joint. We also took extraoral panoramic radiographs. Pain was scored on following scale: 0: no pain; 1: mild pain (pain is present, but it does not cause discomfort); 2 moderate (pain causes discomfort, but is tolerable), and 3: intense (pain causes significant discomfort and it is difficult to tolerate).

Once a diagnosis of Ernest syndrome had been established, all patients were injected with 40 mg of triamcinolone acetonide (Trigon depot®, Bristol-Myers Squibb, Madrid, Spain) in the mandibular insertion of the stylomandibular ligament (Fig. 2). Patients whose symptoms persisted received an additional injection 15 days after the first one. In all cases, occlusion was corrected with a prosthesis. Patients were told to avoid forcing the jaw open and...
The patients’ clinical characteristics are listed in Table 1. The series included 6 female patients aged between 35 and 51 years, with a mean age of 40.3. None of the patients reported prior history of traumatic episodes; however, 4 of the 6 women experienced pain following dental treatment (in 2 patients after endodontic procedures and in the other 2 patients after complex extractions) and they reported prolonged mouth opening during the procedures.

The pain timeline prior to the patient’s first visit ranged from 6 months to 4 years, with a mean of 23 months. Pain was mild in 3 patients and moderate in the other 3; all subjects reported that pain appeared slowly with no clear trigger event and that it was dull, continuous, and poorly defined. All patients presented symptoms unilaterally, with the left side being dominant (66.6%).

Intraoral examination revealed missing teeth in 5 patients. During palpation of the muscles of mastication, 2 patients experienced pain in the ipsilateral masseter muscles and 2 in the ipsilateral internal masseter and pterygoid muscles. Palpation of the temporomandibular joint was painless in all cases.

Four patients achieved complete symptom remission with treatment (Table 2). The 2 remaining patients needed additional injections 15 days after the first one since residual pain persisted. Occlusion was corrected in all 5 patients with missing teeth by means of removable partial dental prostheses in 4 patients (cases 2 to 5) and a fixed dental prosthesis in the other case (case 1) (Table 2). All patients recovered completely after treatment and remained stable at least during those months in which they were monitored (minimum follow-up period of 12 months).

Discussion

The present study presents 6 clinical cases of orofacial pain that were diagnosed as Ernest syndrome. Other similar entities do exist, such as Eagle syndrome, which presents as a painful alteration of the stylohyoid process in patients who have undergone a tonsillectomy. This syndrome was described by Dr Eagle in 1937.4 The aetiology and pathogenesis of Eagle syndrome are both controversial.5 Symptoms include mineralisation, lengthening, or even calcification of the stylohyoid ligament. Unlike Ernest syndrome, which only affects the stylohyoid ligament, especially at the mandibular insertion, Eagle syndrome is usually bilateral.2 The clinical, radiological, and therapeutic characteristics of both syndromes are listed in Table 3.

No new studies of Ernest syndrome have been published in the medical literature since 1987, the year in which the largest series in the literature appeared.2 This study6 analysed 68 patients with a mean age of 37.7 years; 82% of the patients in the sample were female. These results are similar to those from our study, in which the mean age was 40.3 and 100% of the subjects were women. A retrospective study7 of 850 patients with temporomandibular joint disorder showed that men were affected less than women by a ratio of 1:6.2. Several studies8 have tried to explain this discrepancy between the sexes by evaluating psychological, endocrine, physical, and behavioural factors, but researchers have been unable to draw any conclusions.

Performing a literature search of the PubMed database using the key words ‘Ernest Syndrome’ and ‘stylohyoid ligament’ yields 7 hits; as 2 are duplicated, only 5 hits are related to this syndrome. Of these articles, 4 are reviews9,12–14 and 1 is the case series by Shankland et al.2 The scarcity of literature may be due to the fact that Ernest syndrome is a little-known syndrome that may be underdiagnosed. This would not have to be the case, however, if the syndrome were included in the International Classification of Headache Disorders.15

The stylohyoid ligament is a thickened band of cervical fascia that extends from the styloid process to the

**Table 1** Patient characteristics: age, sex, prior dental treatments, and type of pain.

<table>
<thead>
<tr>
<th>Age (years)</th>
<th>Sex</th>
<th>Prior dental treatments</th>
<th>Progression (months)</th>
<th>Intensity</th>
<th>Radiation</th>
<th>Side</th>
</tr>
</thead>
<tbody>
<tr>
<td>Case 1</td>
<td>42</td>
<td>Female</td>
<td>Yes</td>
<td>12</td>
<td>Moderate</td>
<td>Arm</td>
</tr>
<tr>
<td>Case 2</td>
<td>51</td>
<td>Female</td>
<td>Yes</td>
<td>48</td>
<td>Mild</td>
<td>Neck</td>
</tr>
<tr>
<td>Case 3</td>
<td>35</td>
<td>Female</td>
<td>No</td>
<td>6</td>
<td>Mild</td>
<td>Shoulder</td>
</tr>
<tr>
<td>Case 4</td>
<td>38</td>
<td>Female</td>
<td>No</td>
<td>12</td>
<td>Moderate</td>
<td>Neck</td>
</tr>
<tr>
<td>Case 5</td>
<td>35</td>
<td>Female</td>
<td>Yes</td>
<td>36</td>
<td>Mild</td>
<td>No</td>
</tr>
<tr>
<td>Case 6</td>
<td>41</td>
<td>Female</td>
<td>Yes</td>
<td>24</td>
<td>Moderate</td>
<td>Eye</td>
</tr>
</tbody>
</table>
Table 2  Data referring to clinical examination, treatment, and follow-up time.

<table>
<thead>
<tr>
<th>Missing teeth</th>
<th>Pain during the palpation of the muscles of mastication</th>
<th>Treatment</th>
<th>Follow-up (months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Case 1</td>
<td>Inferior Yes</td>
<td>No. doses (triamcinolone 40 mg)</td>
<td>Prosthesis type</td>
</tr>
<tr>
<td>Case 2.</td>
<td>Sup. and inf. Yes</td>
<td>1</td>
<td>Fixed bridge 14</td>
</tr>
<tr>
<td>Case 3</td>
<td>Sup. and inf. No</td>
<td>2</td>
<td>RPP 17</td>
</tr>
<tr>
<td>Case 4.</td>
<td>No</td>
<td>1</td>
<td>RPP 12</td>
</tr>
<tr>
<td>Case 5.</td>
<td>Sup. and inf. No</td>
<td>2</td>
<td>RPP 14</td>
</tr>
<tr>
<td>Case 6.</td>
<td>Sup. and inf. Yes</td>
<td>1</td>
<td>RPP 15</td>
</tr>
</tbody>
</table>

Fixed P: fixed prosthesis; RPP: removable partial prosthesis; Sup. and inf.: superior and inferior.

The posterior border of the ramus of the mandible, medially to the angle of the mandible. The stylomandibular ligament is responsible for limiting the opening and protrusion movements of the jaw. The pathogenesis of Ernest syndrome is not completely clear; it could be related to macrotrauma caused by blows to the jaw, cranial trauma, or whiplash caused by traffic accidents.1617 All of these processes produce the same clinical symptoms: distension of the ligament connecting the stylohyoid process to the mandibular gonial angle.918 Researchers have indicated that when this ligament stretches more than 25 to 30 mm, it may cause cervical and pharyngeal pain as well as pain when swallowing, speaking, and opening the mouth, perception of a foreign body in the oropharynx, and pain radiating to the ear.519 Some have suggested that prolonged dental procedures may lengthen and strain the stylomandibular ligament, thereby causing painful symptoms.2 This hypothesis is supported by the fact that 4 out of our patients experienced pain during the month following lengthy dental procedures.

The pain due to Ernest syndrome can be confused with that resulting from masticatory muscle disorders, especially in cases of myofascial pain, and other entities such as Eagle syndrome, neuralgia of the third branch of the trigeminal nerve, glossopharyngeal neuralgia, persistent idiopathic facial pain, temporomandibular joint dysfunction syndrome, pain associated with periconoritis, parotid pain, and cervical pain. Trigeminal neuralgia is a disorder related to the peripheral or central fibres of the trigeminal nerve. As such, it causes pain limited to one or more branches of the trigeminal nerve. This neuralgia may be idiopathic or present as a result of the syndrome’s underlying aetiology (secondary or symptomatic). Table 4 presents the differential diagnosis for idiopathic and symptomatic trigeminal neuralgia.20

The study by Shankland et al.2 described 68 patients with Ernest syndrome; of this number, 77.9% were successfully treated non-surgically with injection of an anaesthetic in the insertion of the ligament, localised corticosteroid injections, and soft diet. Our patients underwent the same treatment.

While the sample size is small, 4 patients recovered completely after treatment. The other 2 patients required an additional injection 15 days after the first treatment due to residual pain. Based on these results, we believe that this treatment, which is easily provided, may alleviate pain and improve quality of life in patients. Here, we analyse the clinical characteristics, treatment, and progression of 6 patients with Ernest syndrome. We believe that the prevalence of this syndrome is higher than the literature may lead us to think, and that Ernest syndrome may be mistaken for other causes of orofacial pain.

Table 3  Diagnosis, symptoms, aetiology, and treatment of both syndromes.

<table>
<thead>
<tr>
<th>Syndrome</th>
<th>Ernest syndrome</th>
<th>Eagle syndrome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Symptoms</td>
<td>Initial phase: sensitivity of the infra-auricular and mandibular region near the mandibular angle.</td>
<td>Type 1 presentation: spastic and persistent pharynx pain in patients following tonsillectomy with styloids &gt; 3 cm or ossification of the stylohyoid ligament.</td>
</tr>
<tr>
<td></td>
<td>Later symptoms: pain in the TMJ, orbital and temporal areas, oropharynx, and coronoid apophysis; toothache in the postero-inferior teeth.16</td>
<td>Type 2 presentation: pharyngeal pain increasing upon moving the head, headache, and vertigo in patients with no prior surgical history.7</td>
</tr>
<tr>
<td></td>
<td>Radiographic imaging is not very useful, since the pathogenesis of the syndrome is related to the inflammation of the stylomandibular ligament.16</td>
<td>Simple radiography of the cervical spine shows lengthening of the styloids, which can be confirmed using a CT (length &gt; 3 cm).8</td>
</tr>
<tr>
<td>Treatment</td>
<td>Local injections with corticosteroids and local anaesthetics. Surgical treatment in severe cases9</td>
<td>Local injections with corticosteroids and local anaesthetics; surgical resection via the oropharynx if medical treatment fails410</td>
</tr>
</tbody>
</table>
Table 4  Differential diagnosis for idiopathic and symptomatic trigeminal neuralgia.

<table>
<thead>
<tr>
<th>Trigeminal neuralgia</th>
<th>Idiopathic</th>
<th>Symptomatic</th>
</tr>
</thead>
<tbody>
<tr>
<td>Clinical examination</td>
<td>Normal results and no organic disease responsible for the symptoms</td>
<td>Sensory or motor deficits of the cranial nerves and signs typical of origin (traumatic, infectious, tumoral, or associated with BTSN)</td>
</tr>
<tr>
<td>Complementary examination</td>
<td>No relevant changes</td>
<td>With changes</td>
</tr>
<tr>
<td>Age</td>
<td>Advanced stage</td>
<td>Any</td>
</tr>
<tr>
<td>Localisation</td>
<td>2nd and 3rd trigeminal branch</td>
<td>1 or more than 1</td>
</tr>
<tr>
<td>Side</td>
<td>Unilateral</td>
<td>Unilateral or bilateral</td>
</tr>
<tr>
<td>Clinical history</td>
<td>No</td>
<td>Yes</td>
</tr>
</tbody>
</table>

BTSN: benign trigeminal sensory neuropathy.

Conflicts of interest

The authors have no conflicts of interest to declare.

References