

# We are IntechOpen, the world's leading publisher of Open Access books Built by scientists, for scientists

5,300

Open access books available

130,000

International authors and editors

155M

Downloads

Our authors are among the

154

Countries delivered to

TOP 1%

most cited scientists

12.2%

Contributors from top 500 universities



WEB OF SCIENCE™

Selection of our books indexed in the Book Citation Index  
in Web of Science™ Core Collection (BKCI)

Interested in publishing with us?  
Contact [book.department@intechopen.com](mailto:book.department@intechopen.com)

Numbers displayed above are based on latest data collected.  
For more information visit [www.intechopen.com](http://www.intechopen.com)



# Proprioceptors in Cephalic Muscles

*Juan L. Cobo, Sonsoles Junquera, José Martín-Cruces,  
Antonio Solé-Magdalena, Olivia García-Suárez  
and Teresa Cobo*

## Abstract

The proprioception from the head is mainly mediated via the trigeminal nerve and originates from special sensitive receptors located within muscles called proprioceptors. Only muscles innervated by the trigeminal nerve, and rarely some muscles supplied by the facial nerve, contain typical proprioceptors, i.e. muscle spindles. In the other cephalic muscles (at the exception of the extrinsic muscles of the eye) the muscle spindles are replaced by sensory nerve formations (of different morphologies and in different densities) and isolated nerve fibers expressing mechanoproteins (especially PIEZO2) related to proprioception. This chapter examines the cephalic proprioceptors corresponding to the territories of the trigeminal, facial, glossopharyngeal and hypoglossal nerves.

**Keywords:** proprioception, muscle spindles, atypical proprioceptors, cephalic muscles, PIEZO2, mechanoproteins

## 1. Introduction

Proprioception is a quality of the somatosensory system that informs the central nervous system about the static and dynamics conditions of muscles and joints. This type of sensitivity has been studied in deep in the muscles depending on the spinal nerves and today the neurobiology of spinal proprioception is well known [1–4]. On the contrary, the neuroanatomy as well as the cellular and molecular bases of the proprioception in the cephalic muscles is not well known. Nevertheless, it is clear that cephalic muscles permanently develop fine adjustments of stretching and tone in facial movements, regulation of chewing force, oromotor reflex behaviors, verbal and nonverbal facial communication, swallowing, coughing, vomiting or breathing [5–7].

The skeletal muscles contain an intrinsic mechanosensory system, the proprioceptive system, which provides unconscious and conscious information to the central nervous system. The proprioceptive inputs originate in specialized sensory organs (proprioceptors) present in muscles (muscle spindles [8, 9]), tendons (Golgi's tendon organs [10]), joint capsules (Ruffini-like sensory corpuscles, Pacinian corpuscles and free nerve endings [11]), and presumably also the skin but their physiological properties suggest they are not the alternative to muscle spindles [2, 12–14]. The information encoded by the proprioceptors gives rise to unconscious

and conscious sensations, necessary for most basic motor functions [15]. For those interested in a recent review and in detail on both types of proprioceptors, we refer to the Banks [8] and Macefield and Knellwolf [16].

Some decades ago, Baumel [17] suggested that proprioceptive impulses from facial muscles are conveyed to the central nervous system via different branches of trigeminal nerve throughout multiple communications with the branches of the facial nerve. Actually, it is accepted that the proprioception of all cephalic muscles depends on the trigeminal nerve [6, 18].

Therefore, the first unresolved issue in cephalic proprioception is whether all cranial nerves that innervate striated muscles also collect their proprioceptive innervation. According to Lazarov [18] the proprioceptive innervation of all cephalic muscles depends exclusively on the trigeminal nerve. In other words: the sensory ganglia of cranial nerves lack of primary sensory neurons and the proprioceptors of the cephalic muscles are supplied by neurons from the trigeminal mesencephalic nucleus [19].

The second aspect pending clarification is: if the proprioception of the cephalic muscles depends exclusively, or mainly, on the trigeminal nerve, how do the fibers of this nerve reach the muscles of the territories of other nerves? This question can be answered because to extensive communications of the trigeminal nerve with other cranial nerves. The trigeminal nerve has numerous connections to the facial nerve [20–34] and the data collected from animal models indicate that the nerve fiber interchange is always from the trigeminal to the facial nerve and not on the contrary [35]. To serve facial proprioception additional connections between the facial and cervical spinal nerves exists [36, 37]. Apart from those communications no specific reference of communications between the trigeminal nerve with the glossopharyngeal, vagal and hypoglossal nerves were found. But presumably the trigeminal proprioceptive fibers pass from the trigeminal nerve to them directly on the target organs themselves (tongue, pharynx, palate) or through their connections with the facial nerve [28, 31, 32].

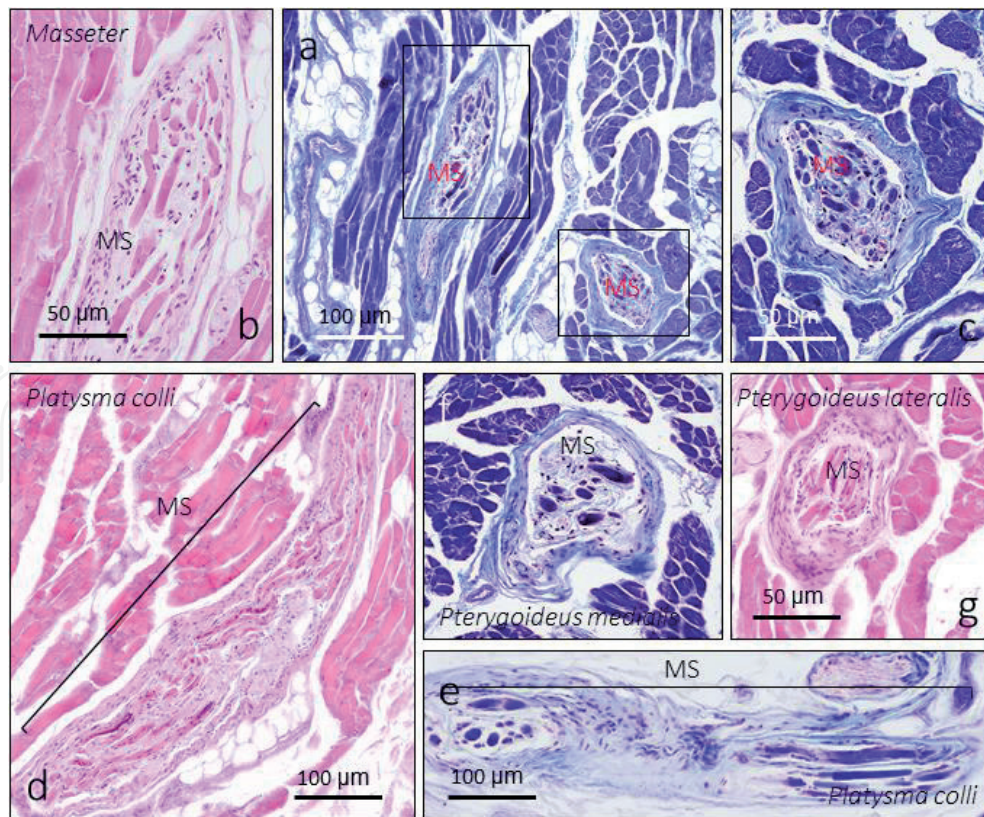
And the third main question of cephalic proprioception regards the identification and characterization of proprioceptors in the cephalic muscles. The skeletal muscles innervated by spinal nerves contain neuromuscular spindles and Golgi tendon organs, in addition to other types of corpuscles with less functional entity [8–11]. However, only the cephalic muscles supplied by the mandibular branch of the trigeminal nerve, and the *platysma colli* muscle contain neuromuscular spindles [38–40]. Therefore, cephalic proprioceptors, if any, have to be represented by other sensory nerve formations other than neuromuscular spindles. Recent studies, using immunohistochemistry techniques associated with specific markers related to mechanization, have shown that facial muscles [34, 41, 42] and some pharyngeal muscles [43] have differentiated sensory structures that presumably replace proprioceptors. However, it cannot be ruled out that sensitive nerve fibers reaching the muscles (especially nociceptive ones) can function as mechanoreceptors-proprioceptors (see [44]).

## 2. Distribution of typical proprioceptors in cephalic muscles

Typical proprioceptors of human cephalic muscles are represented by neuromuscular spindles as most of them lack Golgi tendon organs since they lack true tendons.

Muscle spindles have been found in muscles innervated by the trigeminal nerve while in the territory of the other cranial nerves, with very rare exceptions, are absent [6]. Recently, Junquera [45] determined the relative density of muscle





**Figure 1.**  
Longitudinal and transversal sections of muscle spindles from different cephalic muscles. MS: muscle spinde.

spindles in human jaw muscles (**Figure 1**; **Table 1**). The *M. temporalis*, *m. masseter*, *m. perygoideus medialis* and *m. pterygoideus lateralis* contained numerous muscle spindles whereas they were less abundant in the *digastricus* and *mylohyoideus* muscles [45, 46]. The absence [45, 47] or presence [48] of muscle spindles in the *tensor veli palatini* muscle, also innervated by the trigeminal nerve, has been reported. It should be noted that atypical proprioceptors were also found in these muscles (**Table 1**; see below).

In muscles where the density of muscle spindles is higher, they consist of thick capsule, a shallow intracapsular space filled with variable number of intrafusal muscle fibers (ranging from 4 to 12). In muscles where the density of neuromuscular spindles was low, in general, the size of the spindles was smaller, had fewer intrafusal fibers and the capsule was less developed [45].

In the territory of the facial nerve one muscle spindle was found in the muscle *orbicularis oculi* in one pediatric specimen [49] whereas abundant muscle spindles have been found in the *platysma colli* [40]. Junquera [45] in her doctoral dissertation also observed typical muscle spindles in the *plastysma colly* more numerous in the cervical segment of the muscle than in the suprahyoid one.

### 3. Atypical putative proprioceptors cephalic muscles

#### 3.1 Criteria to characterize atypical proprioceptors

The identification of putative sensory receptors in the cephalic muscles that may serve as proprioceptors was based on the following criteria: independence of the nerve trajectory, be placed in close relation to muscle fibers, show a morphologically differentiated aspect, and display immunoreactivity for any putative mechanoprotein [34].

Muscle	MS	Type I	Type II	Type III	INS*
<i>M. temporalis</i>	14	6	8	6	Yes
<i>M. masseter</i>	23	3	3	6	Yes
<i>M. pterygoideus lateralis</i>	18	3	14	7	Yes
<i>M. pterygoideus medialis</i>	21	5	10	3	Yes
Venter anterior m. digastricus	2	3	1	1	Yes
<i>M. mylohyoideus</i>	1	3	1	1	Yes
<i>M. tensor veli palatini</i>	0	2	2	1	Yes
<i>M. corrugator supercilii</i> + <i>M. depressor supercilii</i>		1	3	7	Yes
<i>M. orbicularis oculii</i> pars palpebralis	0	3	11	9	Yes
pars orbitalis	0	1	7	9	Yes
<i>M. orbicularis oris</i> pars marginalis	0	5	19	12	Yes
pars labialis	0	7	13	7	Yes
<i>M. zygomaticus maior</i>	0	1	4	4	Yes
<i>M. zygomaticus minor</i>	0	1	2	0	Yes
<i>M. buccinator</i>	0	19	28	10	Yes
<i>M. depressor labii inferioris</i> + mentalis	0	0	8	2	
<i>M. levator labii superioris</i>	0	1	1	3	Yes
Platysma colli**	12/8	11/7	4/7	6/8	Yes/ Yes
<i>M. genioglossus</i>	1	16	28	10	Yes
<i>M. palatoglossus</i>	0	0	5	3	Yes
<i>M. uvulae</i>	0	0	7	3	Yes
<i>M. constrictor pharyngis superior</i>	0	0	6	14	Yes
<i>M. constrictor pharyngis inferior</i>	0	0	5	9	Yes

\*Isolated nerve fibers displaying immunoreactivity for any of the mechanoproteins investigated.

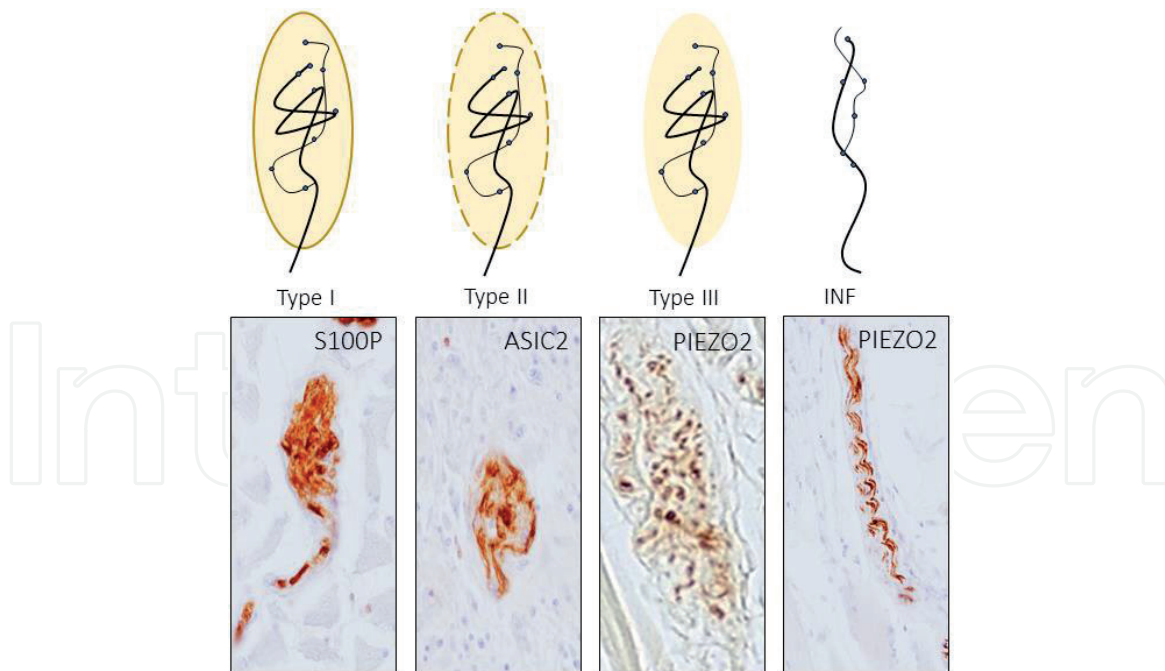
\*\*facial/cervical segments.

**Table 1.**

Distribution and density of muscle spindles (MS), atypical proprioceptors (types I to III) and isolated nerve fibers (INF) in muscles supplied by the trigeminal nerve (green), facial nerve (blue), hypoglossal nerve (white) and glossopharyngeal nerve (brown).

In agreement with the above premises, capsulated and non-capsulated corpuscle-like structures of variable size and shape containing numerous axon profiles complexly arranged, have been identified. Given the morphologic heterogeneity of the corpuscle-like structures that fulfill the preestablished criteria we attempt to classify them into three types: type I, capsulated by a thin capsule, the glial cells variably arranged and showing different morphologies; type II, partially capsulated (the capsule being continuous with the perimysium), with variable morphology and in most of the cases the direction of the long axis was parallel to the one of muscular fibers; type III, non-capsulated and both the axon and Schwann-like cells are variably arranged (**Figure 2**).

On the other hand, it is now well established that at the basis of mechanosensitivity are mechanically-gated ion channels [50]. At present acid-sensing ion



**Figure 2.**  
Types of putative proprioceptors in human cephalic muscles. INF: Isolated nerve fibers.

channel 2 (ASIC2) and Piezo2 have been detected in muscle spindles and are strong candidates to initiate the mechanotransduction in proprioceptors [50–56]. Also, the putative mechanoprotein transient-receptor potential vanilloid 4 (TRPV4) was detected in proprioceptors of the facial and pharyngeal muscles [42, 43].

### 3.2 Distribution in the territory of the facial nerve

No typical muscle spindles have been found in the human facial muscles [42, 57–61] with the exception of on the facial part of the muscle *platysma colli* [40, 45]. Conversely, they contain numerous atypical proprioceptors (**Table 1**) the type II of being the predominating and the greater density being observed in the *buccinator* and *orbicularis oris* muscles.

### 3.3 Distribution in the territory of the glossopharyngeal nerve

Most research have not found typical muscle spindles in the muscles innervated by the glossopharyngeal nerve although they are present in the human *palatoglossus* muscle [48].

Regarding the pharyngeal muscles, typical muscle spindles were never found with the exception of the *constrictor pharyngis inferior* of the crab-eating monkey (*Macaca irus*) [62]. Nevertheless, human pharyngeal muscles are richly innervated. In particular, the *constrictor pharyngis superior* and muscle *constrictor pharyngis inferior* (innervated by branches of the pharyngeal plexus, derived from the glossopharyngeal and vagal nerves, and a small contribution of facial nerve; [63]) contain type II and III putative proprioceptors and isolated nerve fibers that display immunoreactivity for mechanoproteins (**Table 1**) [43].

### 3.4 Distribution in the territory of the hypoglossal nerve

As far as we know no muscles spindles have been reported in tongue muscles. Junquera [45] observed one muscle spindle in the genioglossus muscle as well as numerous putative proprioceptors (**Table 1**).



Therefore, as a whole, the cephalic muscles have proprioceptive innervation, although only the muscles innervated by the trigeminal nerve and the *platysma colli* muscle innervated by the facial nerve contain neuromuscular spindles. The cephalic proprioceptors may be involved in the coordination of facial movements and non-verbal communication, in language, swallowing and some other reflexes [64–66].

IntechOpen

### Author details

Juan L. Cobo<sup>1,2\*</sup>, Sonsoles Junquera<sup>3</sup>, José Martín-Cruces<sup>1</sup>,  
Antonio Solé-Magdalena<sup>1</sup>, Olivia García-Suárez<sup>1</sup> and Teresa Cobo<sup>2,4</sup>

1 Departamento de Morfología y Biología Celular, Grupo SINPOS, Universidad de Oviedo, Oviedo, Spain

2 Instituto Asturiano de Odontología, Oviedo, Spain

3 Servicio de Radiología, Complejo Hospitalario Universitario de Santiago de Compostela, Santiago de Compostela, Spain

4 Departamento de Cirugía y Especialidades Médico-Quirúrgicas, Universidad de Oviedo, Oviedo, Spain

\*Address all correspondence to: juancobodiaz@gmail.com

### IntechOpen

© 2021 The Author(s). Licensee IntechOpen. This chapter is distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/3.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited. 

## References

- [1] Dijkerman HC, de Haan EH. Somatosensory processes subserving perception and action. *Behav Brain Sci.* 2007; 30: 189-201.
- [2] Proske U, Gandevia SC. The proprioceptive senses: their roles in signaling body shape, body position and movement, and muscle force. *Physiol Rev.* 2012; 92: 1651-1697.
- [3] Butler AA, Héroux ME, Gandevia SC. Body ownership and a new proprioceptive role for muscle spindles. *Acta Physiol. (Oxf)* 2016; 220: 19-27.
- [4] Blecher R, Heinemann-Yerushalmi L, Assaraf E, Konstantin N, Chapman JR, Cope TC, Bewick GS, Banks RW, Zelzer E. New functions for the proprioceptive system in skeletal biology. *Philos Trans R Soc Lond B Biol Sci.* 2018; 373 (1759). pii: 20170327.
- [5] Miller AJ. Oral and pharyngeal reflexes in the mammalian nervous system: their diverse range in complexity and the pivotal role of the tongue. *Crit Rev Oral Biol Med.* 2002; 13:409-425.
- [6] Cattaneo L, Pavesi G. The facial motor system. *Neurosci Biobehav Rev.* 2014; 38:135-159.
- [7] Frayne E, Coulson S, Adams R, Croxson G, Waddington G. Proprioceptive ability at the lips and jaw measured using the same psychophysical discrimination task. *Exp Brain Res.* 2016; 234:1679-1687.
- [8] Banks RW. The innervation of the muscle spindle: a personal history. *J Anat.* 2015; 227:115-135.
- [9] Bewick GS, Banks RW. Mechanotransduction in the muscle spindle. *Pflugers Arch.* 2015; 467:175-190.
- [10] Ackermann PW, Salo P, Hart DA. Tendon Innervation. *Adv Exp Med Biol.* 2016; 920:35-51.
- [11] Heppelmann B. Anatomy and histology of joint innervation. *J Peripher Nerv Syst.* 1997; 2: 5-16.
- [12] Collins DF, Refshauge KM, Todd G, Gandevia SC. Cutaneous receptors contribute to kinesthesia at the index finger, elbow, and knee. *J Neurophysiol.* 2005; 94: 1699-1706.
- [13] Macefield VG. Physiological characteristics of low-threshold mechanoreceptors in joints, muscle and skin in human subjects. *Clin Exp Pharmacol Physiol.* 2005; 32:135-144.
- [14] Andreatta RD, Barlow SM. Somatosensory gating is dependent on the rate of force recruitment in the human orofacial system. *J Speech Lang Hear Res* 2009; 52:1566-1578.
- [15] Akay T, Tourtellotte WG, Arber S, Jessell TM. Degradation of mouse locomotor pattern in the absence of proprioceptive sensory feedback. *PNAS USA.* 2014; 111:16877-16882.
- [16] Macefield VG, Knllwolf TP. Functional properties of human muscle spindles. *J Neurophysiol.* 2018; 120: 452-467.
- [17] Baumel J. Trigeminal-facial nerve communications. *Arch Otolaryngol* 1974; 99: 34-44.
- [18] Lazarov NE. Neurobiology of orofacial proprioception. *Brain Res Rew* 2007; 56: 362-383.
- [19] Yoshida A, Moritani M, Nagase Y, Bae YC. Projection and synaptic connectivity of trigeminal mesencephalic nucleus neurons controlling jaw reflexes. *J Oral Sci.* 2017; 59:177-182.



- [20] Bendella H, Spacca B, Rink S, Stoffels HJ, Nakamura M, Scaal M, Heinen H, Guntinas-Lichius O, Goldbrunner R, Grosheva M, Angelov DN. Anastomotic patterns of the facial parotid plexus (PP): A human cadaver study. *Ann Anat.* 2017; 213:52-61.
- [21] Shimada K, Moriyama H, Ikeda M, Tomita H, Shigihara S, Gasser RF. 1994. Peripheral communication of the facial nerve at the angle of the mouth. *Eur Arch Otorhinolaryngol.* S:110-112.
- [22] Hwang K, Han JY, Battuvshin D, Kim DJ, Chung IH. Communication of infraorbital nerve and facial nerve: Anatomic and histologic study. *J Craniofac Surg* 2004; 15:88-91.
- [23] Hwang K, Hwang JH, Cho HJ, Kim DJ, Chung IH. Horizontal branch of the supraorbital nerve and temporal branch of the facial nerve. *J Craniofac Surg.* 2005; 16: 647-649.
- [24] Hwang K, Jin S, Park JH, Kim DJ, Chung IH. Relation of mental nerve with mandibular branch of the facial nerve. *J Craniofac Surg* 2007. 18:165-168.
- [25] Hwang K, Yang SC, Song JS. Communications Between the Trigeminal Nerve and the Facial Nerve in the Face: A Systematic Review. *J Craniofac Surg.* 2015; 26: 1643-1646.
- [26] Tohma A, Mine K, Tamatsu Y, Shimada, K. Communication between the buccal nerve (V) and facial nerve (VII) in the human face. *Ann Anat* 2004;186: 173-178.
- [27] Hu KS, Kwak J, Koh KS, Abe S, Fontaine C, Kim HJ. Topographic distribution area of the infraorbital nerve. *Surg Radiol Anat.* 2007; 29:383-388.
- [28] Diamond M, Wartmann CT, Tubbs RS, Shoja MM, Cohen-Gadol AA, Loukas M. Peripheral facial nerve communications and their clinical implications. *Clin Anat.* 2011; 24:10-8.
- [29] Odobescu A, Williams HB, Gilardino MS. Description of a communication between the facial and zygomaticotemporal nerves. *J Plast Reconstr Aesthet Surg.* 2012; 65:1188-1192.
- [30] Yang HM, Won SY, Kim HJ, Hu KS. Sihler staining study of anastomosis between the facial and trigeminal nerves in the ocular area and its clinical implications. *Muscle & Nerve* 2013; 48:545-550.
- [31] Shoja MM, Oyesiku NM, Griessenauer CJ, Radcliff V, Loukas M, Chern JJ, Benninger B, Rozzelle CJ, Shokouhi G, Tubbs RS. Anastomoses between lower cranial and upper cervical nerves: a comprehensive review with potential significance during skull base and neck operations, part I: trigeminal, facial, and vestibulocochlear nerves. *Clin Anat.* 2014; 27:118-130.
- [32] Shoja MM, Oyesiku NM, Shokouhi G, Griessenauer CJ, Chern JJ, Rizk EB, Loukas M, Miller JH, Tubbs RS. A comprehensive review with potential significance during skull base and neck operations, Part II: glossopharyngeal, vagus, accessory, and hypoglossal nerves and cervical spinal nerves 1-4. *Clin Anat.* 2014b; 27: 131-144.
- [33] Tansatit T, Phanchart P, Chinnawong D, Apinuntrum P, Phetudom T, Sahraoui YM. A Cadaveric Study of the Communication Patterns Between the Buccal Trunks of the Facial Nerve and the Infraorbital Nerve in the Midface. *J Craniofac Surg.* 2016; 27:214-218.
- [34] Cobo JL, Solé-Magdalena A, Junquera S, Cobo T, Vega JA, Cobo J. The proprioception in the muscles supplied by the facial nerve. En: *Selected Topics in Facial Nerve Disorders*, I Al-Zwaini y M

Jalal Husseis (Eds). Intech, London, 2019; pp. 1-14.

[35] Bowden RE, Mahran ZY.

Experimental and histological studies of the extrapetrous portion of the facial nerve and its communications with the trigeminal nerve in the rabbit. *J Anat.* 1960; 94: 375-386.

[36] Yang HM, Kim HJ, Hu KS. Anatomic and histological study of great auricular nerve and its clinical implication. *J Plast Reconstr Aesthet Surg.* 2015; 68:230-236.

[37] Brennan PA, Elhamshary AS, Alam P, Anand R, Ammar M. Anastomosis between the transverse cervical nerve and marginal mandibular nerve: how often does it occur? *Br J Oral Maxillofac Surg.* 2017; 55:293-5.

[38] Scutter SD, Türker KS. The role of the muscle spindles in human masseter. *Hum Mov Sci.* 2001; 20: 489-497.

[39] Osterlund C, Liu JX, Thornell LE, Eriksson PO. Muscle spindle composition and distribution in human young masseter and biceps brachii muscles reveal early growth and maturation. *Anat Rec (Hoboken)* 2011; 294:683-693.

[40] May A, Bramke S, Funk RHW, May CA. The human platysma contains numerous muscle spindles. *J Anat.* 2018; 232: 146-51.

[41] Cobo JL. Neuroanatomía de la cara. Anatomía topográfica, quirúrgica, radiológica y microscópica de la inervación de la cara. Doctoral Thesis. Oviedo: Universidad de Oviedo; 2016.

[42] Cobo JL, Abbate F, de Vicente JC, Cobo J, Vega JA. Searching for proprioceptors in human facial muscles. *Neurosci Lett.* 2017; 640:1-5.

[43] de Carlos F, Cobo J, Macías E, Feito J, Cobo T, Calavia MG, García-Suárez O, Vega JA. The sensory

innervation of the human pharynx: searching for mechanoreceptors. *Anat Rec (Hoboken).* 2013; 296:1735-1746.

[44] Ackerley R, Watkins RH. Microneurography as a tool to study the function of individual C-fiber afferents in humans: responses from nociceptors, thermoreceptors, and mechanoreceptors. *J Neurophysiol.* 2018; 120: 2834-2846.

[45] Junquera S. Proprioceptores cefálicos: identificación basada en la expresión de PIEZO2. Doctoral Thesis. Oviedo: Universidad de Oviedo; 2020.

[46] Saverino D, De Santanna A, Simone R, Cervioni S, Cattrysse E, Testa M. Observational study on the occurrence of muscle spindles in human digastric and mylohyoideus muscles. *Biomed Res Int.* 2014; 2014: 294263.

[47] Kuehn DP, Templeton PJ, Maynard JA. Muscle spindles in the velopharyngeal musculature of humans. *J Speech Hear Res.* 1990; 33: 488-93.

[48] Liss JM. Muscle spindles in the human levator veli palatini and palatoglossus muscles. *J Speech Hear Res.* 1990; 33:736-746.

[49] Nelson CC, Blaivas M. Orbicularis oculi muscle in children. Histologic and histochemical characteristics. *Invest Ophthalmol Vis Sci.* 1991; 32:646-54.

[50] Kröger S. Proprioception 2.0: novel functions for muscle spindles. *Curr Opin Neurol.* 2018; 31:592-598.

[51] Simon A, Shenton F, Hunter I, Banks RW, Bewick GS. Amiloride-sensitive channels are a major contributor to mechanotransduction in mammalian muscle spindles. *J Physiol.* 2010; 588: 171-85.

[52] Chen CC, Wong CW. Neurosensory mechanotransduction through

acid-sensing ion channels. *J Cell Mol Med.* 2013; 17:337-349.

[53] Gautam M, Benson CJ. Acid-sensing ion channels (ASICs) in mouse skeletal muscle afferents are heteromers composed of ASIC1a, ASIC2, and ASIC3 subunits. *FASEB J.* 2013; 27:793-802.

[54] Woo SH, Lukacs V, de Nooij JC, Zaytseva D, Criddle CR, Francisco A, Jessell TM, Wilkinson KA, Patapoutian. Piezo2 is the principal mechano transduction channel for proprioception. *Nat Neurosci.* 2015; 18:1756-1762.

[55] Florez-Paz D, Bali KK, Kuner R, Gomis A. A critical role for Piezo2 channels in the mechanotransduction of mouse proprioceptive neurons. *Sci Rep.* 2016; 6:25923.

[56] Anderson EO, Schneider ER, Bagriantsev SN. Piezo2 in Cutaneous and Proprioceptive Mechano transduction in Vertebrates. *Curr Top Membr.* 2017; 79: 197-217.

[57] Stål P, Eriksson PO, Eriksson A, Thornell LE. Enzyme-histochemical differences in fibre-type between the human major and minor zygomatic and the first dorsal interosseus muscles, *Arch Oral Biol.* 1987; 32: 833-841.

[58] Stål P, Eriksson PO, Eriksson A, Thornell LE. Enzyme-histochemical and morphological characteristics of muscle fibre types in the human buccinator and orbicularis oris, *Arch Oral Biol.* 1990; 35: 449-458.

[59] Kamen G, De Luca CJ. Firing rate interactions among human orbicularis oris motor units. *Int J Neurosci.* 1992; 64: 167-75.

[60] Happak W, Burggasser G, Liu J, Gruber H, Freilinger G. Anatomy and histology of the mimic muscles and the supplying facial nerve, *Eur Arch Otorhinolaryngol.* 1994; 1994: S85-S86.

[61] Goodmurphy CW, Ovalle WK. Morphological study of two human facial muscles: orbicularis oculi and corrugator supercilii. *Clin Ant.* 1999; 12: 1-11.

[62] Sengupta BN, Sengupta S Muscle spindles in the inferior constrictor pharyngis muscle of the crab-eating monkey (*Macaca irus*). *Acta Anat (Basel)* 1978; 100:132-135.

[63] Shimokawa T, Yi SQ, Izumi A, Ru F, Akita K, Sato T, Tanaka S. An anatomical study of the levator veli palatini and superior constrictor with special reference to their nerve supply. *Surg Radiol Anat,* 2004; 26:100-105.

[64] Wild B, Erb M, Eyb M, Bartels M, Grodd W. Why are smiles contagious? An fMRI study of the interaction between perception of facial affect and facial movements. *Psychiatry Res.* 2003; 123: 17-36.

[65] van der Bilt A. Assessment of mastication with implications for oral rehabilitation: a review. *J Oral Rehabil.* 2011; 38, 754-780.

[66] Schötz S, Frid J, Löfqvist A 2013. Development of speech motor control: lip movement variability. *J Acoust Soc Am.* 2013; 133: 4210-4217.