Cranial nerves XIII and XIV: nerves in the shadows

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Abstract: It has been known for over a century that these cranial nerves exist, and that they are not typographical errors nor a sensational event reported in the medical literature. A number of scientific articles on anatomy highlight how textbooks on descriptive anatomy do not always consider variables such as differences related to the geographical areas where people live, and these differences do exist. This is an important concept not only for surgeons, but also for all medical professionals who use manual techniques when treating their patients, ie, osteopaths, chiropractors, physiotherapists, and other manual therapists. This paper highlights the latest developments regarding these cranial nerves, offering at the same time some ideas for further reflection when looking at clinical scenarios that appear to bear little relationship to each other. Inclusion of these concepts in everyday anamnesis is encouraged.

Keywords: cranial nerve, facial nerve, trigeminal system, Tolosa-Hunt syndrome

Introduction

Cranial nerve XIII is also known as the “zero nerve” or “nerve N”. First discovered in 1870 in sharks and other types of fish, it was initially referred to as the nerve of pinkus. In 1905, it became classified as terminal because, in other animal species, it was observed to extend into the region of the lamina terminalis (delineating the anterior wall of the third ventricle and, in its lower section, before the optic chiasma, forming the optic recess).1 From an osteopathic perspective, this area can develop into a dysfunction point with regard to cranial kinetics, with different clinical presentations.2 This gives us an insight into the importance and complexity of this nerve. Recent textbooks, published after 1987, have classified cranial nerve XIII as the zero nerve, given that it is more rostral than the other cranial nerves. Although first formally mentioned in a paper on human anatomy in 1945,1 it has rarely been mentioned in the medical literature since. The probable reason for this is that, during anatomical dissection, when taking off the dura mater, this nerve is torn off and cannot be found on subsequent examination. However, as long as the pia mater remains intact, the nerve can be located between the olfactory stalk and the rostral section of the optic chiasma.1 The nerve is attached to the pia mater by connective tissue,1 and this concept is useful for osteopaths in order to link dysfunction of the nerve to craniosacral movement.

Cranial nerve XIV was first identified in 1563, but it was not until 1777 that it was mentioned in a textbook as the nerve of Wrisberg.3 In modern textbooks, it is referred to as the nervus intermedius or “intermediary nerve”.1 Its name is consistent with its intermediary location between the facial nerve (cranial nerve VII) and the superior section of the vestibulocochlear nerve (cranial nerve VIII).4 It is well known to cause...
otalgia and related surgical operations. This confirms that this nerve is an independent entity and not a mere ramification of cranial nerve VII.

**Nerve N**

Anatomy is still a field rich in findings, and as regards the data collected so far, nothing can be taken for granted. There are many reference texts on anatomical dissection that demonstrate numerous variations in the human body. In humans, nerve N is composed of a small tract of nerves with nonmyelinated fibers. Unlike the other cranial nerves, nerve N lies in the most rostral area where the brainstem and olfactory bulb are located. Nerve N passes through the medial region of the stria olfactoria (ie, medial in relation to the anterior perforated space) and, via the rectus gyrus of the orbital surface of the frontal lobe, reaches the crista galli laterally. Finally, it passes through the cribiform plate of the ethmoid bone more medially and deeply than do the tracts of the olfactory nerve.

Nerve N is very thin in adults but is easily identifiable in the fetal stage. It has a small ganglion, with other as yet unidentified afferents, indicating that it might have some residual function to warrant consideration in diagnosis and treatment. As demonstrated by research in other species, the nerve is linked with the olfactory nerve in terms of its location and probably also its function. From an embryological perspective, it seems to have developed from the area of the olfactory cells, although some have hypothesized that it may originate from the neural crest.

Generally speaking, stimulation of a hormonal response in men can be linked with the nasal epithelium (through chemosensitive receptors or unidentified bonds with pheromones). Research in humans shows that nerve N is directly connected to the limbic system, at least in the fetus, and consequently to the olfactory system. Further studies in the human model suggest that this nerve has the ability to store small polypeptides, including luteinizing hormone-releasing hormone, acetylcholine, and neuropeptide Y. The latter can also be found in the amygdala, with a number of functions in influencing behavior, including nociception.

The current view is that nerve N has the ability to affect sexual behavior, but the extent to which it does this is unknown as yet. It is believed to act by stimulating luteinizing hormone-releasing hormone on contact with pheromones, improving olfaction and identifying odors, and this series of events will affect sexual behavior.

In several animal species, nerve N secretes or stimulates the secretion of luteinizing hormone-releasing hormone, also known as gonadotropin-releasing hormone. This hormone coordinates the hypothalamus-pituitary-gonadal axis. In animals, this nerve can be located without difficulty, and its functions are more evident, ie, it is presumed to control sexual behavior and reproductive activity. Stimulation of nerve N is believed to trigger a series of hormonal cascades from the nasal region to the reproductive system. Therefore, one hypothesis is that nerve N might trigger an endocrine response, directly or jointly with other nerve structures, such as the kisspeptin neural network. In humans, particularly in females, this group of neurons can be identified as being mainly in the preoptic area and in the infundibular nucleus of the mediobasal hypothalamus.

Could this mean that women may be more affected by nerve N? We still do not know. Undoubtedly, a woman’s sense of smell is stronger during ovulation. Moreover, in the human model, the area in question is a source of interest regarding sexual behavior, although there are no decisive writings on this topic.

Undeniably, these are interesting hypotheses, but await verification. Finally, the trigeminal system can be stimulated by odors. If a connection between the cervical sympathetic ganglia, the trigeminal system, and the aforementioned nerve was confirmed, we might better understand and explain some cervical disorders, such as those related to the menstrual cycle or chronic rhinitis. In addition, we could link certain sexual behavioral disorders to trigeminal pain.
Many other hypotheses might be formulated, but what is certain is that the human body is a sole entity. In order for the body to maintain homeostasis, each and every part needs to be in harmonious and tensegritive collaboration, with regard to not only fascial but also immunological, neuroendocrine, and psychological functioning. Recent research on the olfactory system theorizes a correlation between olfactory changes and certain psychiatric and behavioral disorders, and other neurological pathologies. \(^{33-35}\) Recently, some scientific articles have been published suggesting that the olfactory sense is a survival tool and essential for enabling attention in conscious individuals. \(^{36}\) A possible connection between the olfactory nerve and nerve N offers further clinical scenarios. However, we are again awaiting further research on this relationship.

**Intermediary nerve**

The intermediary nerve has always been considered a ramification of the facial nerve, which is complex\(^ {37,38}\) and all its connections have not been clearly revealed as yet. However, the intermediary nerve is actually a separate and independent nervous entity.\(^ {39,40}\) One study has demonstrated at autopsy that this nerve has four different anatomical origins: the brainstem, where it is evident as a tract of nerve fibers; the top of cranial nerve VII; the top of cranial nerve VIII; and finally, multiple sites in the greater area of the brainstem.\(^ {39}\) According to this study, its ramifications are between 2 and 5, not 4 as we have always thought.\(^ {39}\) It comprises visceral efferents comprised of preganglionic and parasympathetic fibers, which involve the submandibular and sublingual lacrimal glands as well as minor salivary glands. It is a short nerve, measuring no more than 21 mm in length.\(^ {40,41}\) These efferents contact the pterygopalatine ganglion and pass through the base of the pterygoid process of the sphenoid bone, then extend into the pterygoid (vidian) canal, finally reaching the pterygopalatine fossa and developing synaptic connections with the homonymous ganglion.\(^ {40}\) Another portion of these preganglionic and parasympathetic efferents reaches the geniculate ganglion in the middle ear through a pathway that runs in parallel with the motor component of cranial nerve VII, and finally arrives at the tympanic section and the chorda tympani nerve, which develops from the facial nerve.\(^ {40,42}\)

In contrast, the afferents of the intermediary nerve derive from receptors known as the concha auriculae located in a small area behind the ear, from the external layer of the membrana tympani, and from part of the wall of the external auditory meatus.\(^ {40}\) Moreover, some afferents develop from the taste receptors on the anterior two thirds of the tongue, from the floor of the mouth and palate, from the nasal mucosa and nasopharynx (presumably because of the connection with the lingual nerve, a branch of the mandibular division of the trigeminal nerve); and finally, from the chorda tympani nerve that branches from the facial nerve.\(^ {3,40}\) Most textbooks report that this nerve has a sensory function, yet motor efferents of this nerve have also been identified, and electrophysiological studies carried out during surgery on patients have revealed that stimulation of this nerve can result in contraction of the orbicularis oris muscle, which has four segments. This has further confirmed its separation from the facial nerve.\(^ {3}\)

From an anatomical perspective, which involves taking into consideration its direct connections with the geniculate nucleus, the pterygopalatine ganglion, and the brainstem, and from an electrophysiological point of view, the intermediary nerve is now regarded as a nervous structure in its own right.\(^ {3}\) This prompts some interesting considerations. First, the intermediary nerve is connected indirectly to the trigeminal system.\(^ {40}\) Moreover, when it comes into contact with the chorda tympani nerve, the two nerves become one and the same, and then, passing through the petrotympanic fissure, the intermediary nerve becomes an extra cranial nerve.\(^ {4,40}\) Further research suggests that if the temporomandibular joint does not move properly for whatever reason, the pterygopalatine ganglion will be affected, and may alter the functions of the aforementioned intermediary nerve. Finally, looking more closely at the intermediary nerve, there are minute anastomoses with nerves VII and VIII at the level of their origin.\(^ {3}\) A logical deduction is that disorders in any of these nerves may affect the physiology of the intermediary nerve and vice versa, which may alter their function.

The recent medical literature cites the intermediary nerve in terms of the problems it may cause, although these do not occur frequently. For example, paroxysmal otalgia (lasting seconds or minutes), found in combination with alterations in lacrimation, salivation, and taste perception, is known as geniculate neuralgia, neuralgia of the intermediary nerve, or Tolosa-Hunt syndrome.\(^ {4,40,41}\) Its causes may be different, ranging from malignancy to vascular compression, but are mainly idiopathic.\(^ {3,41,43,44}\) Finally, it is important to consider the intermediary vein,\(^ {39}\) which is adjacent to the intermediary nerve, which may alter the electrical function of this nerve in the event of anomalous venous drainage. As osteopaths, we know that anomalous drainage of the cranial venous system may cause dysfunction in the cranial nerves.\(^ {2}\) These data can be useful to us in order to understand and treat the symptoms mentioned in this paper.
Conclusion
In the light of the information currently available in the medical literature, the nomenclature for the cranial nerves should be revised. Further studies are needed to improve our knowledge of the clinical scenarios that may arise from physiological alteration of their function, as well as a more detailed anatomical description of medical strategies which may be helpful in treating related problems.

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The authors report no conflicts of interest in this work.

References