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## Sphenopalatine ganglion block with botulinum toxin type A using a three-dimensional injection guide for facial pain of cancerous origin: study of a clinical case.

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## Abstract

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**Objective:** Sphenopalatine ganglion block remains a noninvasive therapeutic option to treat diverse facial pain syndromes. The aim of this case report was to verify the effectiveness and safety of three-dimensional (3D) injection guide for sphenopalatine ganglion block in a tongue cancer patient.

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**Case report:** A 53-year-old man presented with a large squamous cell carcinoma of the right base of the tongue, classified cT4N2M0 and exhibiting excruciating facial pain. A botulinum toxin injection was performed following Dr. Yoshida's method, using a 3D injection guide.

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**Conclusions:** Sphenopalatine ganglion block with 3D injection guide can be an effective treatment against cancer pain. In this case report a complete withdrawal of painkillers was obtained less than 24 hours after botulinum toxin injection.

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**Keywords:** Sphenopalatine ganglion block, tongue cancer, pain, botulinum toxin, 3D guide

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## Introduction

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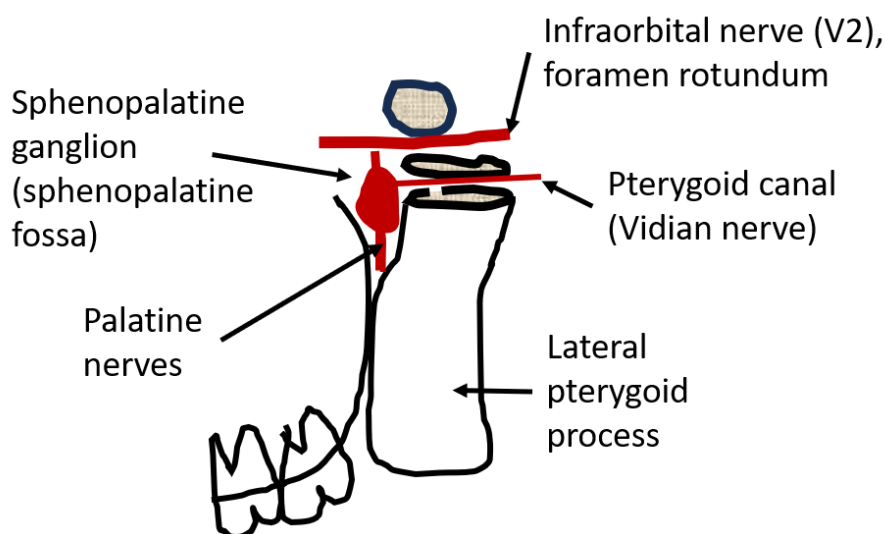
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The sphenopalatine ganglion block is used in the management of cluster headache, trigeminal neuralgia, migraine, postoperative pain of head and face, postherpetic neuralgia, musculoskeletal pain, in other pain syndromes of head and face, and for head and neck cancer pain [1]. Prasanna et al., used sphenopalatine ganglion block for advanced cancer in tongue and floor of mouth with immediate pain relief for the patients [1, 2]. Prasanna et al., used local anesthetic with nasal sinuscope approach without any additional imaging to improve finding the area of sphenopalatine ganglion [2]. However, there is still controversy over both the ideal method and the pharmacological agent used [1].

The ideal technique should be efficient, minimally invasive, and likely to cause the fewest side effects. In addition, faced with a real demand from the population, it should be possible to perform it in private practice in order to unclog hospitals and in particular interventional radiology departments.

The sphenopalatine ganglion is a parasympathetic ganglion located in the sphenopalatine fossa, opposite the sphenopalatine foramen, posterior to the middle turbinate and maxillary sinus, anterior to the pterygoid canal and round foramen and inferior to the sphenoid sinus and to the maxillary branch of the trigeminal nerve (Figure 1).



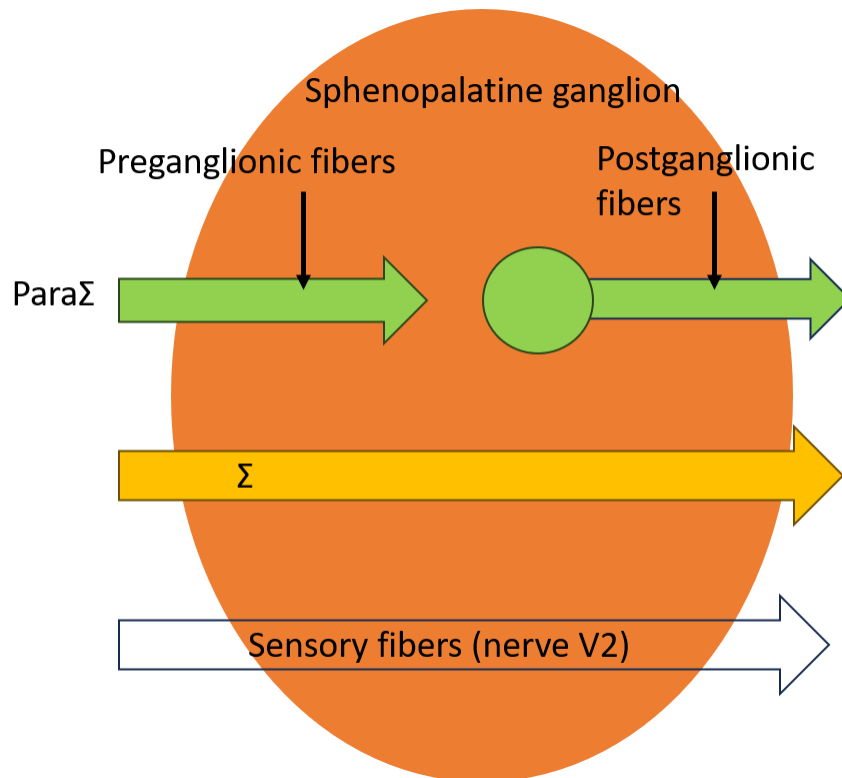
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Fig. 1. Transverse section at the level of the sphenopalatine ganglion.

62 The nerve fibers then distribute to the nasal cavity, palate, upper pharynx, lacrimal  
63 gland and meningeal vessels. Although traversed by orthosympathetic and sensory  
64 fibers, its designation as a parasympathetic ganglion results from the fact that only  
65 this system establishes a synaptic connection within the ganglion (Figure 2).  
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67 **Fig. 2. Nerve afferents and outflows from the sphenopalatine ganglion.**  
68 ParaΣ: parasympathetic nervous system; Σ: sympathetic nervous system.  
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71 This essential characteristic makes this ganglion a privileged target for the action of  
72 botulinum toxin type A (TBA), a protein complex produced by the anaerobic gram-  
73 positive bacterium *Clostridium botulinum*, by inhibiting the release of acetylcholine  
74 and therefore the neurotransmission between pre- and post-ganglionic  
75 parasympathetic fibers for up to a year [3].  
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77 The intraoral technique presented in this article has already been performed  
78 successfully on patients with trigeminal neuralgia by Yoshida, without any adverse  
79 effects described [4]. The idea here is to use the same methodology and operative  
80 technique but this time to evaluate its effectiveness on pain of oncological origin.  
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## Case report

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A 53-year-old patient presented to a maxillofacial surgery consultation because of very debilitating right hemifacial pain that had been present for 5 months. Several dentists and doctors were consulted without success and tooth 48 was extracted. Palpation of the base of the tongue in consultation revealed the presence of a hardened mass, however without any surface mucosal lesion. The requested MRI detected the presence of a large mass of 42 mm long axis invading the extrinsic and intrinsic musculature of the posterior part of the free border and the base of the tongue on the right side, with suspicion of extension beyond from the midline. During the assessment also including a PET-CT and a biopsy, the diagnosis of basic squamous cell carcinoma of the tongue classified cT4N2M0 was made (Figure 3).



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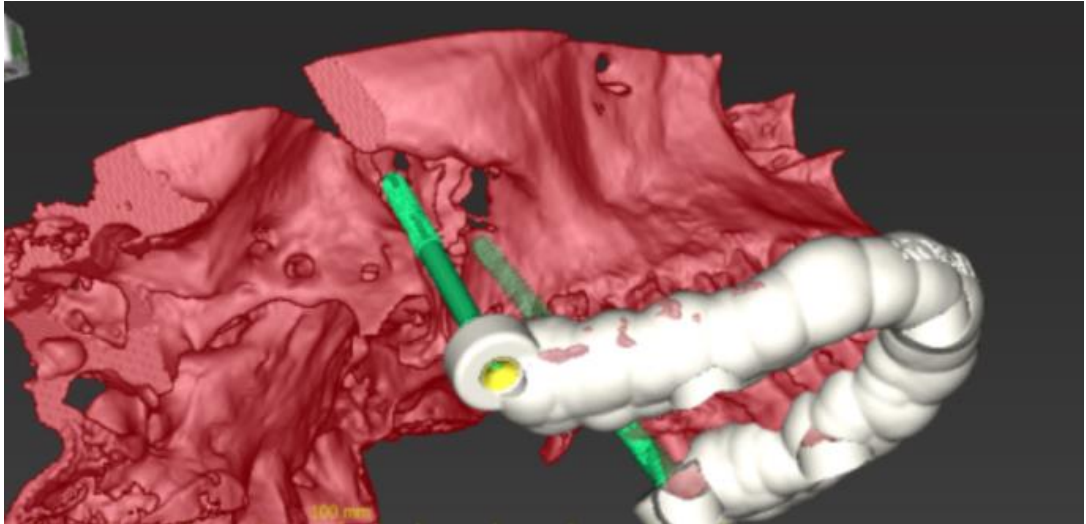
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**Fig. 3. MRI (T2) of tongue base squamous cell carcinoma (star) classified cT4N2M0.**

The tumour, deemed unresectable, was treated with radiotherapy, chemotherapy and immunotherapy. On a numerical scale from 0 to 10, the patient judged his pain at 6/10 with analgesic treatment and at 9/10 in the event of therapeutic abstention. His analgesic treatment was: Lyrica 75mg 3x in the morning; Lyrica 150mg 3x in the evening, Dafalgan 1g 4 tablets/day; Oxyxontin 40mg 2x/day; Oxynorm 10mg 2x/day; Tegretol 200mg 2x/day; Dexamethasone 6mg 1x/day. After written consent from the patient, the intraoral injection guide was performed using the technique mentioned by Yoshida.

A cone beam computed tomography (CBCT) examination of the upper jaw including the ceiling of the sphenopalatine fossa was performed, and then merged with an intraoral scan recording the dentition of the upper jaw. After topographic validation of the sphenopalatine ganglion by a radiologist, the needle insertion path

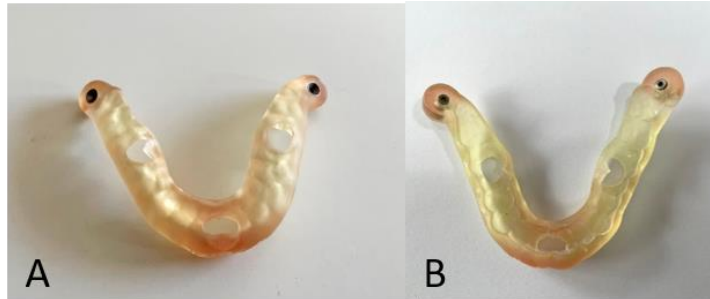
110 was simulated using a 40mm zygomatic implant pointing towards the ganglion  
111 (CoDiagnostiX (Straumann)) (Figure 4).  
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114 **Fig. 4. 3D simulation of the needle insertion path, injection guide in**  
115 **place, within the sphenopalatine fossa.** In this specific case, a 40 mm  
116 zygomatic implant makes it possible to point the ceiling of the pit with the  
117 surface of the guide as a reference. Simulation performed using the  
118 CoDiagnostiX software (Straumann).  
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120 A custom-made tray fitted with two metallic guides located posterior to the wisdom  
121 teeth was made and 50 units of botulinum toxin type A diluted in 1cm<sup>3</sup> of  
122 physiological fluid was injected to the sphenopalatine fossa using a 50mm X 21G  
123 needle (Microlance) at a depth of 40mm after local anesthesia with 0.85 ml of  
124 mepivacaine hydrochloride (Scandonest) in the area of the injection site (Figures 5,  
125 6).  
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127 The pain was rated 1/10 with the visual analogue scale 24 hours after the injection.  
128 A month later, the pain remained perfectly controlled and unchanged. Our patient  
129 relapsed in pain 2 months later and died soon after without having received a second  
130 injection.  
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**Fig. 5. A. Extrados of the tooth-supported injection 3D guide.  
B. Intrados of the tooth-supported injection 3D guide.**



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**Fig. 6. The injection time.**

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## Discussion

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Botulinum toxin type A has been shown to be effective in the treatment of chronic migraines as well as cluster headaches [5, 6]. However, the transnasal and percutaneous routes seem to cause more undesirable effects such as epistaxis, pain, swelling at the injection site, diplopia, dysphagia, asymmetry of the nasolabial groove, dry eye. The transnasal approach can be made difficult in case of nasal obstruction [7, 8].

152 Approaching the sphenopalatine ganglion intraorally via the greater palatine  
153 foramen is generally considered painful and technically difficult [9].  
154 On the other hand, access to the sphenopalatine fossa by an approach lateral rather  
155 than medial to the wisdom tooth seems to be a more promising technique [4]. The  
156 needle is oriented by the custom-made intraoral guide and the vestibular approach  
157 seems easier than an approach via the greater palatine canal. Local anesthesia before  
158 injection thus seems optional [4].  
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160 However, the depth of the insertion of the needle was not controlled by the 3D  
161 guide. Yoshida used a needle of 60mm (Yoshida). We used a needle of 50mm but  
162 inserted 40mm for more safety. Yoshida measured the target point on CT scan of  
163 pterygopalatine fossa, however, not explained in this article principles how the depth  
164 was controlled to arrive to the selected depth [4]. We may use in the future  
165 endodontic stops inserted on the needle to control the depth of insertion more  
166 accurately and closely to the 3D simulation. Also, we did not control the time of  
167 injection.  
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169 The 3D guide is reusable, which represents a clear advantage in terms of cost. The  
170 action is fast, from the first injection and the injection can be repeated if necessary.  
171 Yoshida recommends in case of recurrence, a reinjection at least 3 months after [4].  
172 For our patient, the effectiveness of botulinum toxin decreased 2 months after the  
173 injection.  
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175 The analgesic mechanism of botulinum toxin does not seem to be fully elucidated.  
176 The inhibition of the parasympathetic system by inhibition of the exocytosis of  
177 acetylcholine is the most widespread theory but the action on the peripheral sensory  
178 fibers and on the central nervous system is under study [6].  
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180 Finally, it is surprising to note that the infiltration of the sphenopalatine ganglion, in  
181 close connection with the maxillary nerve, relieves pain originating from a territory  
182 innervated by the third trigeminal branch. The presence of dendrites coming from  
183 the ganglion of Gasser within the sphenopalatine ganglion could explain this  
184 phenomenon [10].  
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186 The intra-oral method of injection of botulinum toxin using a 3D guide seems to be  
187 particularly safe and likely to provide a real analgesic effect in patients for whom  
188 conventional treatments have proven to be ineffective. The treatment proposed in  
189 this clinical case allowed a total withdrawal of painkillers for a period of 2 months  
190 in an oncological patient.  
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- **Ethical approval:** there was no need for the ethical approval for this case report

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- **Informed consent:** there was no need for the informed consent for this ca se

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report as all the images were anonymized and no private data were provided

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allowing the patient's identification.

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**Authors contribution:**

Author	Contributor role
Bourgeois Thomas	Conceptualization, Investigation, Methodology, Data curation, Resources, Validation, Writing original draft preparation, Writing review and editing
Gilon Yves	Writing original draft preparation, Writing review and editing

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