A hemangioma in the masseter muscle: a case report

Daehwan Park¹, Sulki Park¹, Bongsoo Baik¹, Sunyoung Kim²
Departments of ¹Plastic and Reconstructive Surgery and ²Pathology, Dongkang Medical Center, Ulsan, Korea

INTRODUCTION

Intramuscular hemangiomas (IMHs) constitute less than 1% of all hemangiomas [1], and are usually located in the skeletal muscles of the trunk or limbs. However, 13.8% of IMHs are located in the head and neck region, where the masseter muscle is the most common site [2]. Due to the rarity of these tumors, their deep intramuscular location and the presence of the overlying parotid gland, accurate preoperative diagnosis occurs in less than 8% of cases [3].

Magnetic resonance imaging (MRI) has shown superiority in the exquisite delineation and contrast of the lesion from its surrounding tissue [4]. Computed tomography (CT) following injection of the contrast agent iopamidol is also helpful in diagnosing IMHs. A major disadvantage of the CT scan is that it is unable to clearly differentiate between the anatomic extent of the IMHs and the muscle [5]. Ultrasonography is able to differentiate between muscle, vascular tissues and surrounding tissues. Color Doppler can differentiate between arterial, venous, and lymphatic lesions. Both modalities are economical, rapid and free of radiation. They can be used as adjunctive tools for a preoperative diagnosis and are best used for follow-up studies [6].

In the present case, the diagnosis of IMH was made with Doppler ultrasonography. Surgical removal of IMHs of the masseter muscle has commonly been done through a preauricular incision [4], but transcervical approach is also a convenient way to expose the masseter muscle without causing injury to the facial nerve. A collar incision in the submandibular region has been used for the transcervical approach. The authors modified the location of the collar incision from the submandibular region, to the lateral cervical region near the lateral mandibular border and called it a lateral cervical incision. The incision started at the inferior portion of the earlobe and proceeded to within 1 cm of the posterior border of the mandibular ramus, with a slight anterior curve at the mandibular angle.

CASE REPORT

A 66-year-old woman was referred to our hospital with a 2-year history of painless swelling in her left cheek during mastication.

Intramuscular hemangiomas of the masseter muscle are uncommon tumors and therefore can be difficult to accurately diagnose preoperatively, due to the unfamiliar presentation and deep location in the lateral face. A case of intramuscular hemangioma of the masseter muscle in a 66-year-old woman is presented. Doppler ultrasonography showed a 34 × 15 mm hypoechoic and hyper-vascular soft tissue mass in the left masseter muscle, suggesting hemangioma. The mass was excised via a lateral cervical incision near the posterior border of the mandibular ramus. The surgical wound healed well without complications.

Abbreviations: CT, computed tomography; IMH, intramuscular hemangioma; MRI, magnetic resonance imaging.

Keywords: Hemangioma / Masseter muscle / Ultrasonography

Correspondence: Bongsoo Baik
Department of Plastic and Reconstructive Surgery, Dongkang Medical Center, 239 Taehwa-ro, Jung-gu, Ulsan 44455, Korea
E-mail: bsbaiksy@naver.com
Received July 7, 2021 / Revised August 10, 2021 / Accepted August 20, 2021
The patient recalled no clear history of trauma to that region. The swelling was soft and unnoticeable in the resting state but became a round, smooth, protruding mass with a diameter of 3 cm when the patient clenched her teeth (Fig. 1). The mass was slightly compressible and movable on the anterior surface of the lateral mandibular body. No pulsation or bruits was noted. Ultrasonography showed a hypoechoic intramuscular mass with tubular extension and internal hypervascularity on Doppler study (Fig. 2). IMH of the masseter muscle was suggested.

Under a provisional diagnosis of masseteric IMH, an incision line was drawn on the left lateral neck about 10mm from the posterior border of the mandibular ramus. After sedation and local anesthesia (0.5% lidocaine and 1:200,000 epinephrine), an incision of less than 5 cm in length was made on the design from the inferior part of the earlobe to the mandibular angle; the incision was slightly curved anteriorly to follow the mandibular angle. Taking care not to disrupt the marginal mandibular branch of the facial nerve, the anterior skin flap was deeply undermined to expose the platysma muscle. The platysma muscle was cut along the mandibular border and the fibrous capsule of the mass was exposed on the anterior surface of the ramus. The fibrous capsule was separated from the covering tissue to isolate the mass from the muscle. During the separation procedure, a short incision was made on the capsule, which revealed a bulging of dark purple, soft tissue, but no bleeding from the mass. The extruded tissue was not pure hemangioma but looked like an unusual mixture of hematoma and multiple small fibers. The extruded mass with its fibrous capsule was completely removed from the muscle tissue after the pedicle was ligated. The soft mass was about 15 mm in diameter (Fig. 3).

After saline irrigation of the wound, the cut ends of the soft tissue were resutured to cover the masseter muscle and the incisional wound was closed by layers without a drain. The postoperative course was uneventful, without surgical complications such as infection or facial palsy. There was no swelling when the patient clenched her teeth and the cosmetic results were excellent (Fig. 4). Histological examination showed mixed vessel-type IMH (Fig. 5).

**DISCUSSION**

Accurate preoperative diagnosis of IMHs is very difficult due to the rarity of these tumors, which account for only 0.8% of all benign vascular tumors [1,7], and a lack of awareness of the characteristic clinical and radiological presentations. A proper radiological evaluation is required before choosing the surgical...
approach, and complete resection is the treatment of choice [8]. MRI is the most helpful and accurate tool for the diagnosis of IMHs that show low signal intensity on T1-weighted images and high signal intensity on T2-weighted images [4]. Plain X-ray and fine needle aspiration cytology may not be sufficiently specific for diagnostic purposes [4]. CT with injection of contrast agent is also helpful for diagnosis of IMHs [5]. Ultrasonography is the first-line approach to assess soft tissue masses [9] and can be used to differentiate between muscular and vascular structures, and Doppler ultrasonography is useful to demonstrate the presence of vascular structures in and around muscles and to evaluate pathological changes like fibrosis and calcification [10]. Angiography is not indicated unless there is a high suspicion of large vascular connections to the tumor with pulsation, bruits and thrills [11].

There is some confusion regarding the use of terms “hemangioma” and “vascular malformation.” Mulliken and Glowacki [12] classified vascular lesions into hemangiomas and vascular malformations. Hemangiomas were defined as present at birth as small red marks with a biphasic growth phase consisting of proliferation and spontaneous involution. Vascular malformations were usually noted at birth, grew proportionately with the child, did not regress, and consisted of capillary, venous, arterial and lymphatic vascular elements.

Allen and Enzinger [13] subdivided hemangiomas into three groups; small-vessels, large-vessels and mixed group with large and small vessels. Vessels with a diameter less than 140 μm were regarded as small vessels while those with a diameter greater than 140 μm were classified as large vessels. In practice, the small-vessel structure would be regarded as capillary hemangiomas, while the large-vessel group might be called cavernous or venous hemangioma.

The treatment of choice for IMHs of the masseter muscle is surgical excision of the lesion. This is preferred over other treatment modalities such as cryotherapy, steroid and sclerosing agent injection, embolization, and radiation therapy [2]. Local recurrence after surgical excision occurs in approximately 18% of cases due to incomplete excision [2]. The surgical approach for IMHs of the masseter muscle should allow wide exposure of the tumor with preservation of the facial nerve and normal regional structures. The preauricular approach may be necessary for complete excision, with superficial parotidectomy to expose IMHs of the masseteric muscle [4]. In the intraoral approach, a fine tube is inserted into Stensen’s duct for positional marking and a mucosal incision is created anterior to Stensen’s duct to approach the tumor [5]. The main advantages of the intraoral approach are easy access to the tumor and an invisible postoperative scar. The disadvantages are that this approach is less likely to provide adequate exposure for optimal tumor resection and identification of adjacent nerves [2].

Fig. 4. Postoperative photographs. (A) State of surgical site and (B) anteroposterior view on clenching her teeth 1 week postoperatively.

Fig. 5. Histopathologic examination. Photomicrograph shows intramuscular hemangioma composed of mixed vessel types including small vessel type (green arrows) and cavernous-like vascular spaces (blue arrows) (H&E, ×40).
In the case described here, the lateral cervical incision within 1 cm of the posterior border of the mandibular ramus is a convenient way to expose the masseter muscle without causing injury to the facial nerve. It leaves a scar on the area that is not easily visible and is associated with a shorter operative time and better preservation of the facial nerves and parotid gland.

In conclusion, Doppler ultrasonography is a preferred initial imaging modality for superficial masses because of its low cost, general availability and high resolution. This study provides a useful reference for the diagnosis of IMH of the masseter muscle. Doppler ultrasonography is also the best choice for follow-up imaging of vascular tumor sites. A lateral cervical incision near the lateral mandibular margin provides a simple and easy approach that can clearly expose an IMH of the masseter muscle without causing injury to the facial nerves.

NOTES

Conflict of interest
No potential conflict of interest relevant to this article was reported.

Ethical approval
The study was approved by the Institutional Review Board of Dongkang Medical Center (IRB No. 2021-05-03) and performed in accordance with the principles of the Declaration of Helsinki. Written informed consent was obtained.

Patient consent
The patient provided written informed consent for the publication and the use of her images.

ORCID
Daehwan Park https://orcid.org/0000-0001-9671-6510
Sulki Park https://orcid.org/0000-0003-0243-2790
Bongsoo Baik https://orcid.org/0000-0002-8019-3310
Sunyoung Kim https://orcid.org/0000-0001-9479-0549

Author contribution
Project administration: SP. Writing - original draft: SP. Writing - review & editing: DP, BB. Resources: SK. Supervision: DP.

REFERENCES

1. Barnes L. Tumors and tumor like lesions of the soft tissue. In: Barnes L, editor. Pathology of the head and neck. 2nd ed. New York: Marcel Dekker; 2001. p. 900.
2. Wolf GT, Daniel F, Krause CJ, Kaufman RS. Intramuscular hemangioma of the head and neck. Laryngoscope 1985;95:210-3.
3. Avci G, Yim S, Misirlioglu A, Akoz T, Kartal LK. Intramuscular hemangioma. Plast Reconstr Surg 2002;109:1748-50.
4. Narayanan CD, Prakash P, Dhanasekaran CK. Intramuscular hemangioma of the masseter muscle: a case report. Cases J 2009;2:7459.
5. Cheng YT, Lai CC. Transcervical excision of intramuscular cavernous hemangioma: a case report. Oncol Lett 2016;11:1657-60.
6. Rahmani G, McCarthy P, Bergin D. The diagnostic accuracy of ultrasonography for soft tissue lipomas: a systematic review. Acta Radiol Open 2017;6:2058460117716704.
7. Park H, Kim JS, Park H, Kim JY, Huh S, Lee JM, et al. Venous malformations of the head and neck: a retrospective review of 82 cases. Arch Plast Surg 2019;46:23-33.
8. Wee SJ, Park MC, Chung CM, Tak SW. Intramuscular hemangioma in the zygomaticus minor muscle: a case report and literature review. Arch Craniofac Surg 2021;22:115-8.
9. Kim JH, Ahn CH, Kim KH, Oh SH. Multifocal intraosseous calvarial hemangioma misdiagnosed as subgaleal lipoma. Arch Craniofac Surg 2019;20:181-5.
10. Lee JJW, Cho WS, Kim SB, Lee DH. Intramuscular cavernous hemangioma of the masseter muscle in child and adolescent. Korean J Head Neck Oncol 2015;31:27-30.
11. Odabasi AO, Metin KK, Mutlu C, Basak S, Erpek G. Intramuscular hemangioma of the masseter muscle. Eur Arch Otorhinolaryngol 1999;256:366-9.
12. Mulliken JB, Glowacki J. Hemangiomas and vascular malformations in infants and children: a classification based on endothelial characteristics. Plast Reconstr Surg 1982;69:412-22.
13. Allen PW, Enzinger FM. Hemangioma of skeletal muscle: an analysis of 89 cases. Cancer 1972;29:8-22.