CAROTID BODY **TUMOR**

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Carotid body tumor is an uncommon neoplasm arising from the carotid bifurcation, and complete surgical resection is the treatment of choice. Most of the tumors generally present a slow growth. Despite slow growth, these tumors enlarge to encase the carotid bifurcation and adjacent cranial nerves. A 57-year-old female was referred to our clinic with a mass in the neck. Neurological examinations revealed no abnormality includung the function of the cranial nerves. She underwent preoperative ultrasonography and computerized tomography. The tumor, which arising from carotid bifurcation, was removed. A temporary shunt was not used, because of the absence of noticeable changes in EEG on the carotid compression test. The patient's neurological status was stable after surgery. The results of the surgical treatment of carotid body tumors are successful with a recurrence rate after complete resection of about 5%, and a low incidence of metastatic disease of 1.7-2%. We think that almost every carotid body tumor can be removed with excellent long-term survival and with low incidence of stroke.

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TÜRKİYE

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arotid body tumor is an uncommon neoplasm arising from the carotid bifurcation, and complete surgical resection is the treatment of choice. The carotid body tumor can be defined as a nonfunctioning paraganglioma1. Bilateral tumors are described in about 4% of cases with a maximum of 25% in familial manifestation (autosomal dominant mechanism)^{2,3}. Most of the tumors generally present a slow growth. Despite slow growth, these tumors enlarge to encase the carotid bifurcation and adjacent cranial nerves4. In addition to the problems related to the local growth, they can show malignant changes from 2-30%, and can metastize4,5.

CASE REPORT

A 57-year-old female was referred to our clinic with a mass in the neck. Her sympoms were dizyness and syncop, which starded six months ago. A hard elastic oval mass smaller than 4 cm in diameter was palpable on the right side of the neck during the last three months. She had hypertension. No vascular bruit or lymph node swelling was evident. Neurological examinations revealed no abnormality including the function of the cranial nerves. A carotid compression test done to determine the possibility of combined

resection of the tumor with the carotid bifurcation revealed no noticeable changes in EEG. She underwent preoperative ultrasonography and computerized tomography. Carotid echo color Doppler revealed a hypervascular tumor originating from the carotid bifurcation (Fig.1,2,3).

SURGICAL TECHNIQUE

The operation was performed under general anaesthesia. A anterior incision to the sternocleidomastoid muscle with a wide preparation of the common carotid artery and then of the external and internal carotid arteries were done. Attention was given not to

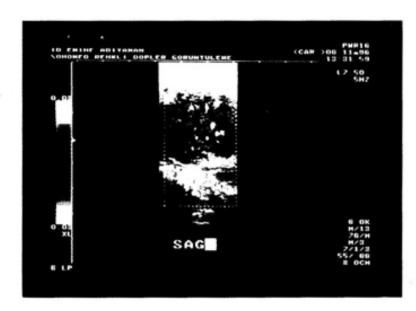


Fig.1- Echocolor Doppler of a carotid body tumor showing a solid vascular mass on the right side at the level of the carotid bifurcation. It encircles external carotid artery almost totally and the internal carotid artery partially, measuring 48x40x30 mm. This vascular mass is mainly perfused by the proximal branches of the external carotid artery. The mass displays the internal carotid artery anterolateral side, but it does not cose any significant obstruction to the blood flow.

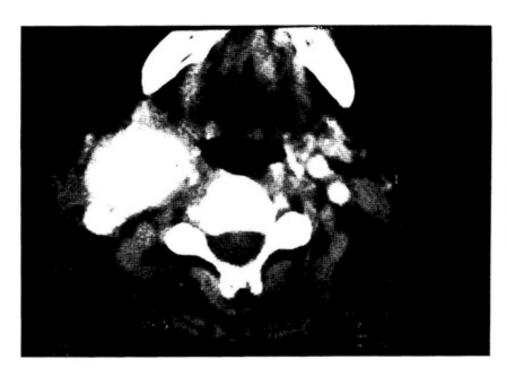


Fig.2- CT scanning of the neck shows that the mass is lokalized at the right carotid bulbus and both internal and external carotid arteries can not be distinguished from the mass.

damage cranial nerves. Because potential bleeding from the tumor during dissection, an autotransfussion system was established. The tumor, which arising from carotid bifurcation, 4x4.5 cm in size, was explored (Fig.4-A). It

had extensive vascular structures, and all the peripheral vascular exisions were ligated and cut. Although it was difficult to remove the entire tumor from the carotid bifurcation, and possible invasion of the tumor into the vessel

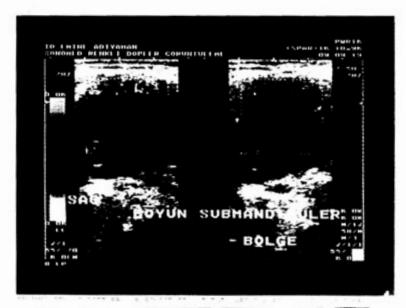


Fig.3- Echo color Doppler, which is performed after CT scanning of the neck with contrast infusion, reveales that the mass is in hypervascular character.



Fig.4- Operative Views.

A- The carotid body tumor is originated from carotid bifurcation.



B- After resection of the tumor a Goro-Tex tubuler graft is anastomozed to the proximal and distal ends of the carotid artery.

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wall was suspected, the tumor was completely resected with the carotid bifurcation. A temporary shunt was not used, because of the absence of noticeable changes in EEG on the carotid compression test. After systemic heparinization, proximal anastomosis of the Gore-Tex tubuler graft was first performed in end-to-end fashion with 5/0 polypropylene sutures (Fig.4-B). Then distal anastomosis was performed. The common carotid artery occlusion time was 15 minutes. Total blood loss was estimated to be 450 ml, and any blood transfusion was not performed. A Hemovac drain was placed after surgical hemostasis. The patient's neurological status was stable after surgery, and no spesific treatment was required. Microscopic examination of the resected specimen revealed a carotid body tumor (Fig.5). Histologically the exuberant growth of paraganglionic cells were identified.

DISCUSSION

Although carotid body tumors may grow slowly and be asymptomatic for many years, they eventually cause symptoms in nearly 75% of patients6, such as carotid artery and cranial nerve compression due to local effect. If the tumor has hormonal activity, it may cause headache, palpitation, hypertension, photofobias, swelling and dysrhytmia7. The first surgical treatment of the carotid body tumor was performed in the end of the 19th century8. During the past 50 years, several trends in the surgical management of cervical paragangliomas have become clear. Instead of only biopsy or partial excision, complete surgical resection is now achieved in nearly all patients⁶. Especially in young patients such as this one, immidiate and complete resection of the tumor is indicated because of the higher incidence of malignant manifestations9. Currently percutaneous needle or incisional

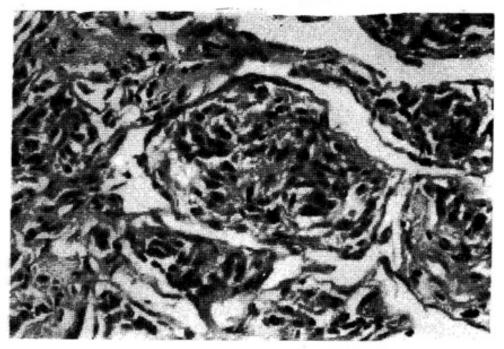


Fig.5- Microscopic view (Hematoxylen-eosin, x400) The exuberant growth of paraganglionic cells are seen.

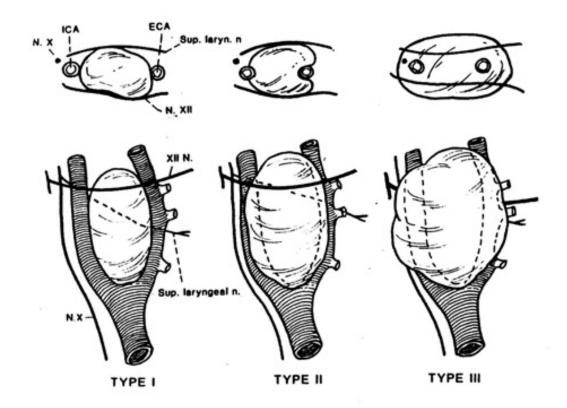


Fig.6- Shamblin classification of difficulty of surgical resection. Group-I tumors are localized and easily resected. Group-II includes tumors adherent or partially surrounding vessels. Group-III paragangliomas intimately surround or encase the vessels. (From Hallett JW et al: J Vasc Surg 7:284-291;1988)

biopsy should be absolutely avoided because of the high incidence of complications⁴. Balfour and Wildner emphasized the difficulty of surgical resection of such tumors¹⁰. Complication rates which include cranial nerve injury, bleeding, carotid artery injury, and stroke have been reported between 10-40% ⁶⁻¹¹.

Shamblin classified the tumors the difficulty of surgical resection according to ¹² (Fig.6):

type-1: These tumors should be easily resected from the adjacent vessel without significant trauma to the vessel wall or tumor capsula,

type-2: They seem to partially surround the vessel and are much more adherent to the vessel adventisia. Dissection of these tumors should be carried out only by surgeons who are prepared to restore the cerebral circulation if it becomes necessary,

type-3: These tumors are intimately adherent to the entire circumference of the carotid

bifurcation, and surgical resection may be impossible. They can be excised only by completely interrupting the internal carotid circulation, and cerebral circulation must be reestabilished by bypass grafting.

If the tumor size smaller than 3-4 cm, it is called small tumor and it can be resected with the surrounding capsula without damaging peripheral vital sutructure. Surgical exposure enough from anterior border is sternocleidomastoid muscle. If the tumor is greater than 4-5cm in size, it is called a great tumor. The surgical exposure could be more agressive such as a T-radical neck incision (Fig.7). During surgical resection, surrounding tissue could be damaged. Preoperative embolisation could be performed to reduce postoperative bleeding. External carotid artery and its branchs could be ligated to reduce bleeding during surgical resection7.

To minimize cranial nerve injury, tumor dissection could be considered in three anatomic zones⁷ (Fig.8):

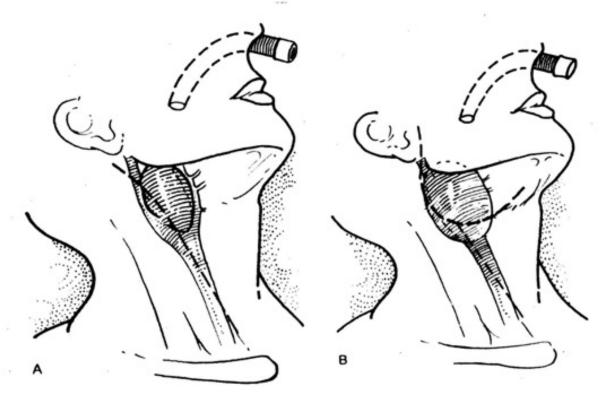


Fig.7- Neck incision for carotid body tumor resection. A nasotracheal tube allows greater displacement of the floor of the mouth during retraction and dissection beneath the mandible. A: Insicion for small tumor (<3-4 cm, Shamblin I). B: Incision for medium and large tumors (>4-5 cm, Shamblin II and III). (From Bergen JJ, Yao JST: Techniques in Arterial Surgery. WB Saunders Company, Philadelphia, 1990; p 217)

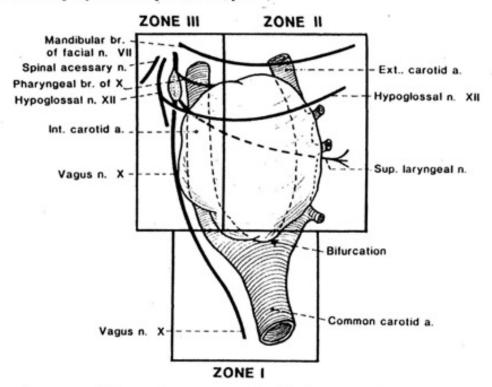


Fig.8- Dissection zones. Most serios nuerovascular injuries occour in Zone-3. (From Hallett JW et al: J Vasc Surg 7:284-291, 1988)

zone-1: Include the carotid artery bifurcation and adjacent vagus nerve. The vagus nerve can be injured during exposure or clamping of the

common carotid artery,

zone-2: Encompasse the external carotid artery territory, the overlying hypoglossal nerve, and the underlying superior laryngeal nerve. The hypoglossal nerve can usually be dissected from the tumor surface. The superior laryngeal nerve can be saved by dissecting right on the tumor surface,

zone-3: Containe the internal carotid artery and confluence of many cranial nerves. Most of the serious neurovascular injuries occur in

this crowded zone.

Alternative treatment method is tumor embolization and radiotherapy. Embolization performed the introduction of by the polyvinylalchohol beads microvasculature of the paraganglioma and gelfoam into artery branch itself to obliterate the flow^{4,8}. Preoperative embolization reduce the blood circulation and tends to cause a necrosis with reduction of the tumoral size4. It reduces tumor bleeding and provides surgical comfort⁸. Even the radiation therapy plays a complementary and limited role; in cases where extirpation is not possible, the radiotherapy can be employed even though the carotid body tumor has a relatively poor radiosensitivity¹³. In our case we have not performed preoperative embolization.

The surgical treatment of the carotid body

tumor has three goals:

1- for long term survey the tumor must to be

resected fully,

2- to diagnose the tumor digital selective angiography is the must important invasive study, and it has advantage for the planning a preoperative embolization,

3- full attention to preserve internal carotid artery during surgical resection, will be postoperative morbidity reduced

mortality.

The result of the surgical treatment of carotid body tumor are successful with a recurrence rate after complete resection of about 5% and a low incidence of metastatic disease 1.7-2% 14,15. We think that almost every carotid body tumor can be removed with excellent long-term survival and with low incidence of stroke.

REFERENCE

- Glenner GG, Grimley PM: Tumours of the extraadrenal paraganglion system (including chemoreceptors). Atlas of pathology. Armed Forces tumour Pathology.2nd series. Institute of 1974:9.
- Hamilton JR, Barros A: Radionucleide 2angiography and surgery for familial bilateral chemodectomas. Eur J Vasc Surg 1987;1:97-105.
- Grufferman S, Gillman MW, Posternak 3-LR: Familial carotid body tumours. Case report and epidemiologic review. Cancer 1980;46: 2116-2122.
- Matticari S, Credi G, Pratesi C, Bertini 4-D: Diagonesis and surgical treatment of the carotid body tumors. J Cardiovasc Surg 1995;36: 233-239.
- Martin CE, Rosenfeld L, McSwain B: 5-Carotid body tumors. A 16 year fallow-up of seven malignant cases. South Med J 1973;63:1236-1243.
- Hallett JW, Nora JD, Hallier LH, Cherry 6-PC: Trends KJ, Pairolero nuerovascular complications of surgical management for carotid body and cervical paragangliomas: A fifty-year experience with 153 tumors. J Vasc Surg 1988;7:284-291.
- Hallett JW: Carotid body 7tumor Techniques in Arterial resection. WB Saunders · Company Surgery. 1990;214-222.
- Smith RF, Shetty PC, Reddy DJ: 8-Surgical treatment of carotid unusual paragangliomas presenting technical difficulties: The value of preoperative embolezation. J Vasc Surg 1988;7:631-637.
- 9-Daudi FA, Thurston OG: Carotid body tumors: the University of Alberta Hospital experience. Canad J Surg 1989;32:159-161.
- 10- Balfour DC, Wildner F: The intercarotid paraganglion and its tumors. Surg Gynecol Obstet 1914;18:203-213.
- 11- McCale DP, Vaccors PS, James AG: Treatment of carotid body tumors. J Cardiovasc Surg 1990;31:356-358.

- 12- Shamblin WR, ReMine WH, Sheps SG, Hattinson FG Jr: Carotid body tumor (chemodectoma): clinicopathologic analysis of ninety cases. Am J Surg 1971;122:735-739.
- 13- Mitchell DC, Cleyne CAC: Chemodectomas of the neck the response to radiotherapy. Br J Surg 1985;72:903-908.
- 14- Lees CD, Levine HL, Beven EG, Tucker HM: Tumours of the carotid body, experience with 41 operative cases. Am J Surg 1981;142: 362-365.
- 15- Rosen IB, Palmer JA, Goldberg M, Mustard RA: Vascular problems associated with carotid body tumors. Am J Surg 1981;68:433-438.