“Carotid body tumour: A case report”

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Abstract: Carotid body tumours are rare, slowly growing benign neoplasm arising from paraganglionic cells at the region of carotid bifurcation. They are the rare tumours of head and neck region derived from neural crest. We report here a case of carotid body tumour which was successfully treated surgically.

Key Word: Carotid body tumour; Paraganglioma; Chemodectoma

I. Introduction

Carotid body tumour is a rare benign paraganglioma which present as neck swelling. Because of its rarity we sometimes misdiagnose it as other disease. It is also known as chemodectoma. It is a highly vascular tumour and arises from the carotid bifurcation. It has a autosomal dominant inheritance pattern with incomplete penetrance.

II. Case Report

A 25 years female presented to the department of ENT of our institution with Painless slowly growing swelling in the right side of upper part of neck for 2 years. There was no pressure symptoms and no history of swelling elsewhere in the body. No past history of tuberculosis and no family history of such swelling was there.

On examination, an oval, around 6 X 4 cm swelling was present in the right side upper part of neck just below and behind the angle of mandible. On palpation, it was non tender, smooth surface, firm in consistency, pulsatile and more mobile transversely than vertically. No regional lymphadenopathy was there.

Routine blood examination, stool and urine examination were within normal limits. FNAC revealed hamartoma. CT Scan (Fig. I) revealed a well defined avidely enhancing isodense mass in right carotid space which displaced the great vessels posterolaterally.

Under GA, neck was explored by a transverse cervical incision at the level of hyoid bone. The mass was medial to carotid bifurcation extending from transverse process of cervical vertebra to mastoid process on the right side. The mass was excised after ligation of external carotid artery keeping internal carotid artery intact. Histopathology (Fig. II) revealed Zellballens with a prominent fibrovascular connective tissue stroma suggestive of carotid body tumour. Postoperative period was uneventful. Patient had no recurrence after 2 years follow up.

III. Discussion

Carotid body tumour are usually benign lesions first mentioned in 1743 by Von Haller and actually described in 1862 by Lushka [1]. The carotid body itself is a small chemoreceptor organ buried posteriorly in the adventitia of the common carotid bifurcation and have a chemoreceptor role by modulating respiratory and cardiovascular function in response to fluctuations in arterial pH, O2, and CO2. Neoplastic growth of this area is known as chemodectomas or paragangliomas.

Incidence of carotid body tumour is only 0.012% [2]. They can be unilateral or bilateral (7%). In approximately 10% a familial trend exists [3]. Familial cases inherited with an autosomal dominant inheritance pattern with incomplete penetrance. There is female gender predominance noted 8.3:1 at high altitudes as compared to a modest one of 2:1 in lower altitudes [4]. The average age of presentation is between 35 to 50 years. Increased prevalence is also associated with high altitudes. 10% of cases are multicentric.

Carotid body tumour is the most common type of paraganglioma and usually presents as an asymptomatic mass in the anterolateral region of the neck. Pressure symptoms may develop with
increasing size of the mass. On physical examination, this tumour is laterally mobile but vertically fixed because of its attachment to the carotid bifurcation [5]. Mass is frequently pulsatile and a bruit may be auscultated over the mass. The large majority of carotid paragangliomas are non-secreting but enquiry must be made about any symptoms of excessive catecholamine secretion like uncontrolled hypertension, tachycardia, facial flushing or excessive sweating.

Majority of carotid body tumours are benign, but may infiltrate locally. About 3% are malignant [6]. The commonest overt malignant manifestation is local recurrence. It grows slowly and rarely metastasizes.

The investigation of choice is bilateral angiography [7] where the tumour appears as a hypervascular oval mass. It substantiates the diagnosis, indicates the size and vascularity of the tumour and determines bilaterality [8,9]. On sonography, paraganglioma present as a well-defined, solid hypoechoic mass and on colour Doppler imaging, hypervascularity with a low-resistance flow pattern is seen [10]. Colour Doppler and angiography are crucial in showing vascularisation details of the tumour [11]. Computed tomography and magnetic resonance imaging reveal a well circumscribed mass at or above the carotid bifurcation, which splays the internal and external carotid arteries. On post contrast scan there is intense enhancement [12,13]. The mass usually shows a salt and pepper appearance caused by vessels with signal void within the tumour stroma.

Fine-needle aspiration has a beneficial role in the diagnosis of carotid body tumour [14]. Paragangliomas consist of clusters of Type I or chief cells and Type II or sustentacular cells. These clusters of cells make up the histologic structure termed Zellballen. Nuclear pleomorphism and cellular hyperchromatism is common in paragangliomas. Malignancy cannot be determined histologically but is reserved for presence of clinical evidances like local, regional, or distant metastasis.

Tissue diagnosis is not necessary and biopsy may be dangerous[15]. Once diagnosed, excision with subadventitial dissection is the treatment of choice. In inoperable cases radiation therapy is recommended [16].

IV. Figures

Figure I: CT Scan (axial view) of neck showing a well-defined avidely enhancing isodense mass (X) in the right carotid space.

Haematoxilin & Eosin stain

(10x10)
Figure II: Photomicrography showing Zellballens (Y) with a prominent fibrovascular connective tissue stroma (Z).

References

ABBREVIATIONS
FNAC - Fine needle aspiration cytology.
CT - Computed tomography.
GA - General anaesthesia.
CTVS - Cardiothoracic vascular surgeon.
pH - Power of hydrogen.
O2 - Oxygen.
Co2 - Carbon di-oxide.