Introduction:
Carotid body tumors (CBT) are distinctly uncommon hypervascular neck tumors.\(^1\)\(^-\)\(^3\) They arise from glomus bodies (paraganglia) located in the crotch of the external and internal artery, at the level of carotid bifurcation. Carotid paraganglia are composed of chemoreceptor cells derived from the carotid primitive neural crest.\(^1\)\(^,\)\(^4\) In general, tumors arising from any paraganglionic tissue are best called paragangliomas (PGLs) and are arbitrarily classified by their relationship to the adrenal gland (adrenal medullary tumors or pheochromocytomas and extra-adrenal PGLs). Carotid body tumour is the most common form of paraganglioma of the head and neck region.\(^5\) It is usually benign and non-functional. It grows and expands slowly and rarely metastasises.\(^5\)\(^,\)\(^6\)

Case Report:
A 50 years male patient was admitted in a private hospital of Dhaka, Bangladesh in June, 2011 with a large (70 × 50 × 45 mm) sized swelling on the right side of the neck. The swelling was non-tender having restricted mobility particularly in vertical axis, lying anterior to the sternocleidomastoid muscle having no fixity to the muscle or skin either. It had transmitted pulsation but there was systolic bruit over the swelling. There was no cranial nerve involvement. Neck glands were not enlarged. The lower limit of the swelling was quite perceptible but the upper limit was indistinct and appeared to have buried under the angle of the mandible and sternomastoid muscle. The patient was a smoker and had an episode of anterior myocardial infarction 4 months back. Coronary angiogram was done which revealed less than 50% proximal left anterior descending artery lesion and medical treatment was advised for him. He was non-diabetic and his blood pressure was within normal limit. He was taking nitrates, lipid lowering agents, antiplatelet drug and low dose beta-blockers. His antiplatelet drug was discontinued for 5 days before operation. His haematological and biochemical investigations did not reveal any abnormality. Twenty four hour Vanilyl mandelic acid (VMA) excretion was normal. Chest X-ray showed mild cardiomegaly and electrocardiogram showed old anterior myocardial infarction. Echocardiogram showed mild left ventricular dysfunction with ejection fraction of 50%. CT carotid angiogram revealed a tumour mass involving right external carotid artery with tumour blush indicating high vascularity.

Abstract:
Carotid body tumor, an extra-adrenal paraganglioma, represents an uncommon entity arising in chemoreceptor tissue located at the carotid bifurcation. They commonly present as asymptomatic neck masses and careful preoperative evaluation is required to find out the functional and vascular status of these tumours. An unusual case of a large (70 × 50 × 45 mm) carotid body tumor of Shamblin type III in a 50 year old male is being presented here. The evaluation, management and the final outcome of this case of carotid body tumour is being reported here.

Keywords: carotid body tumour, paragangliomas.
The operation was done under general anaesthesia with endotracheal intubation. A proximal transverse incision was made in the neck to take control of the right common carotid artery. The tumour was then exposed by another generous distal transverse skin crease incision over the most prominent part of the tumour and proximal control of the external carotid artery and distal control just beyond the tumour was taken. There were lot of collateral large unusual communication between the tumour mass and tributaries of the internal jugular veins and the main trunks which ultimately made sacrificing part of internal jugular vein mandatory, despite the risk of cerebral oedema. The tumour along with the external carotid artery was excised with much difficulty. Haemostasis was ensured and the wound was closed keeping a closed suction drainage system in situ. Histological examination of the tumor mass was done and it revealed typical features of non-secreting paraganglioma.

The patient recovered well from the effect of anaesthesia and there were no neurological deficit, but the patient developed left ventricular failure in the immediate postoperative period. The patient required two days ventilator and inotropic support in the ICU. He was then transferred to the general ward and was discharged home 5 days later. On first follow up visit after 3 weeks he complained of severe headache with diplopia of vision. There was mild hemiatrophy of the right half of the tongue and diplopia of vision. He was referred to a neurosurgeon who advised magnetic resonance venogram of the cerebral sinuses. The MRA-V did not reveal any abnormality but considering his symptom a course of antibiotic, diuretic and steroid was given to reduce the raised intracranial pressure. The decision to put a CSF shunt to peritoneal cavity was deferred. The patient improved substantially, his headache and diplopia were completely relieved after 1 month treatment and his tongue muscle function also improved leaving a little bit of slurring of speech. The patient is alright now.

Fig.2: Digital subtraction angiogram (DSA showing normal contralateral neck vessel.

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Discussion:
Paragangliomas (PGLs) are extra-adrenally located in 10% of cases. 90% of these extra-adrenal PGLs are located in the abdomen, mainly in chromaffin cells of the organ of Zuckerkandl. The remaining 10% of the extra-abdominal PGL include CBT and glomus tumours. The carotid body is a vascular reddish-brown structure about the size of a grain of rice located within adventitia posteromedial to bifurcation of the common carotid artery. The carotid body acts as a chemoreceptor responding to variations in oxygen tension and carbon dioxide concentrations; henceforth alternatively named chemodectomas.

CBT usually presents in the fourth to sixth decades of life (age range: 3 months–89 years). The average tumour size is 4.5 cm × 3.5 cm × 3 cm, with the largest ones exceeding 15 cm in diameter and weighing 200 g. Three different types of carotid body tumors (CBTs) are described: sporadic, familial, hyperplastic. The sporadic form is the most common type, representing approximately 85% of carotid body tumors (CBTs). The familial type (10-50%) is more common in younger patients. The hyperplastic form is very common in patients with chronic hypoxia, which includes those patients living at a high altitude (> 5000 feet above sea level), like those patients living in New Mexico, Peru and Colorado. The hyperplastic form is also seen in patients who have chronic obstructive pulmonary disease (COPD) or cyanotic heart disease. The incidence of bilateral CBT varies from 10% to 25% in different series. Here a case of sporadic CBT measuring 70 × 50 × 45 mm in size in a 50 years male patient has been described.

Most of the lesions are benign, but malignant lesions are seen in 6%-12% of cases. The diagnosis of malignancy is reserved for tumours with local, regional and distant metastasis. There are no histological features that distinguish benign from malignant lesions. Between 10% and 50% of PGLs are hereditary (autosomal dominant) due to alterations in genes coding for succinate-ubiquinone oxidoreductase subunit D (SDHD), B (SDHB) and C (SDHC).

Patients with CBT typically present with a painless mass in the angle of the jaw that may be partially covered by the sternocleidomastoid muscle. These vascular tumours may transmit pulsations from nearby carotid arteries or may be pulsatile inherently. At the time of diagnosis, cranial nerve palsies are present in 10% of cases with the X, XII, V and VII cranial nerves commonly involved in decreasing order of frequency, while the involvement of cervical sympathetic chain produces ipsilateral Horner’s syndrome. Carotid sinus syndrome with bradycardia, hypotension and unconsciousness can occur spontaneously or secondary to head movement or pressure on the tumour. Episodic symptoms of phaeochromocytoma are seen in 1%-3% of cases.

The size of the tumour is important not only for the clinical manifestations, but also for a decision on treatment. In 1971, Shamblin introduced a classification system based on the tumour size, classifying small tumours which could easily be resected from the vessels as Group I. Group II includes tumours that are intimately associated and compressed carotid vessels, but that could be resected with careful subadventitial dissection. Group III consists of tumours that are large and typically encease the carotid artery, requiring complete or partial vessel resection and replacement. Here a case of CBT of Shamblin Group III has been presented.

Ultrasound studies in both sides of the neck, may exclude the presence of lymph nodes, thyroid or brachial cysts. In most cases, color-coded Doppler sonography evaluates the hypervascularity and upward intratumoural blood flow in a neck mass at carotid angle, suggesting a CBT. Accurate diagnosis is based on angiographic criteria, the most reliable of these being the separation and splaying of internal and external carotid arteries (known as the lyre sign). On CT scanning, CBT is easily recognized as hypervascular mass located at the carotid bifurcation, changing the architecture of the angle between the internal and external carotid artery, bringing them apart from each other resembling a saddle. Arterial anatomy of carotid vessels and their relationship to an enhancing lesion of the neck can also evaluated by CT angiography. MRI is the most important imaging modality in evaluating CBTs in relation to surrounding soft tissue and vascular structures. Total body scintigraphy using 123I-MIBG is used as a screening tool to detect distant additional primary or metastatic lesion in patients or their relatives. Biochemical assays are routinely employed for the diagnosis of a functioning tumour, irrespective of symptoms. The usual methods are measurement of urinary free catecholamines and its metabolites VMA and MNs in a 24-hour urine sample. Histopathologically, CBT are firm, rubbery well-encapsulated masses composed of nests of epithelioid cells (chief cells), with clear cytoplasm separated by delicate stroma (zellballen pattern).

The success in treating benign and malignant CBT is based on early diagnosis, complete resection of the tumour after an adequate catecholamine blockade, which is given prophylactically. Complete resection of the primary mass is the treatment of choice. Postoperative cranial nerve deficits
and arterial injury have remained a significant problem. Patients with larger tumours have an increased incidence of complications. Endovascular embolisation may reduce the operative time and limit blood loss. Though we do not employ this technique preoperatively most surgeons now prefer this technique for larger tumours of Shamblin Grade III. In patients who are not suited for surgery, irradiation has been attempted for symptomatic relief with varying success. Chemotherapy using vincristine, dacarbazine and cyclophosphamide along with 131I-MIBG therapy has been attempted for patients with systemic metastasis. Radiotherapy may be effective for large tumours, recurrent tumours and patients who are poor candidates for surgery.

References: