Superior Mediastinal Teratoma- A Case Report with review of literature

Elaf Abdulwahhab Hamdi*, Rabea Salim Abd Aljabbar Sofi Ali**, Professor Wahda Mohammad Taib Al-Nuaimy*

*Department of Pathology, College of Medicine, University of Mosul, ** Department of Cardiothoracic and Vascular Surgery, Al-Jumhuri Teaching Hospital, Nineveh-Health Office, Mosul, Iraq

Correspondence: ela@uomosul.edu.iq

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ABSTRACT

"A teratoma is a germ cell tumor composed of tissue derived from two or three germ layers-ectoderm, mesoderm, and endoderm". Teratomas are mostly found in children & young adults & usually originated in gonads & extragonadal midline tissues such as retroperitoneum, sacrococcygeal region & mediastinum. Mediastinal teratomas are relatively rare representing 3% of all tumors within the chest with an average incidence of 8 cases per year and the anterior mediastinum being the most common site for these tumors, benign teratomas form only 10% of them. We are presenting a case of 39 years old male who presented with progressive exertional dyspnea, orthopnea with frequent attacks of cough for 3 years and feeling of pulsatile mass in left upper parasternal area in the last 4 months. Chest x-ray showed superior mediastinal shadow projecting to left upper chest zone and deviating the trachea to the right side of chest, provisional diagnosis was aneurysm of aortic arch. Computed tomography (CT) scan of chest and CT angiography done and revealed a superior mediastinal mass of heterogeneous opacity containing fluid, soft tissue, bones and calcification encroaching the right pulmonary artery and compressing and deviating the trachea to the right. Complete surgical removal done through left thoracotomy, Final diagnosis of a mature cystic teratoma was given on histopathology. This case is being presented here because mediastinum is a rare site for teratoma and superior mediastinal teratoma was reported in only few cases from review of literature.

Keywords: teratoma, germ cell tumor, mediastinum, Superior mediastinal Teratoma.

ورم المنصف الصدری العلوي المسخي : تسجيل حالة مع مراجعة مقال

إلف عبد الوليد حمدي* ، ربيع سالم عبد الجبار** ، الاستاذ وحدة محمد طيطي العبد

*فرع الأمراض / كلية الطب / جامعة الموصل ، **قسم جراحة القلب والصدر والأورام الدموية / مستشفى الجمهوري التعليمي / دائرة صحة نينوى / الموصل / العراق

الخلاصة

الورم المسخي هو ورم الخليط الجوئي ويكون من نسيج مشتق من طبقتين أو ثلاث طبقات جوية - الأديم الظهار، والأديم الوسط والأديم الباطن. توجد الأورام المسخية في الغالب عند الأطفال والشباب ولذا يوجد في نسبته العدد التناسلي على امتداد خط الوسط في الجسم. وقد تم الإبلاغ عن مواقع خارج العقد التناسلي، مثل منطقة عجان المصع، خلف الصفاق والمنصف الصدر. تعتبر الأورام المسخية في المنصف الصدر نادرة نسبيا حيث تشكل 8-13% من جميع الأورام في المنصف الصدر و 1-2% فقط من أورام الخلايا الجوئية التي تتطور في المنصف. نقدم حالة لرجل يبلغ من العمر 39 عاما يعاني من ضيق التنفس المجدد المتزايدي، والاختناق الليلي مع نوبات متكررة من السعال لمدة 3 سنوات، والشعور بوجود كتلة نابضة في منطقة جانبي القص العلوي الأيسر في الأشهر الأربعة الأخيرة من العدد. أظهرت الأشعة السينية للصدر ظل كتلة في المنصف الصدرى يعود على منطقه الصدر العلوي اليسرى ويحرف القصية الهوائية إلى الجانب الأيمن. وكان التشخيص النهائى للورم المسخي الصدرى الناتج عن التصوير الأوعية القطاعي (CT) التفاضلي عبر علاج الدم الأوربي. وقد أجري التصوير المقطعي للصور الأوربي المحويب وأظهر كتلة منصف علوي غير متجانسة تحتوي على السوائل والأورام الرخوة والظام والتلك التي تتفاوت على الارض الأخرى الأيمن وتشوه وتحرف القصية الهوائية إلى اليمين. تم إجراء الإزالة الجراحية الكاملة من خلال عملية فتح
INTRODUCTION

The term “teratoma” derives its origin from the Greek words “terato” and “oncoma” meaning “monster” and “swelling,” respectively. Their unpredictable behavior, obscure origin and bizarre microscopic appearance, make them interesting. Primary mediastinal tumors are rare representing 3% of all tumors within the chest, benign teratomas form only 10% of them. The anterior mediastinum being the most common site for these teratomas. After complete surgical excision benign cystic teratomas have excellent prognosis.

Case Report

A 39 years old male patient presented with progressive exertional dyspnea and orthopnea with frequent attacks of cough for 3 years duration and feeling of pulsatile mass in left upper parasternal area in the last 4 months. Chest x-ray showed superior mediastinal shadow projecting to left upper chest zone and deviating the trachea to the right side of the chest, differential diagnosis was aneurysm of aortic arch and lymphoma. Computed tomography (CT) scan of chest and CT angiography (Fig.1) done and revealed a superior mediastinal cystic mass of heterogeneous opacity containing fluid, soft tissue, bones and calcification encroaching the right pulmonary artery and compressing and deviating the trachea to the right.

Complete surgical removal done through a left thoracotomy. Intraoperatively there was a mass 13 cm × 9 cm × 7.5 cm present in the superior mediastinum protruding to left hemithorax (Fig.2). The mass was encroaching on great vessels, and severely adherent to pericardium, lung, and chest wall. Complete surgical resection of the mass was done including pieces of pericardium and mediastinal pleura. Patient post operatively developed hoarseness of voice due to left recurrent laryngeal nerve injury.

Grossly multiple fragments of soft tissue with hair and sebaceous material, bone and cartilage (Fig. 3). Microscopical examination revealed a cyst wall with marked hyaline degenerative changes and calcification, filled with hair, sebum, mature cartilage, respiratory epithelium with mucus secreting glands (Fig. 4). Histopathology reported was benign mature cystic teratoma. The case is being presented here because mediastinum is an uncommon site for teratoma and superior mediastinum is not being reported as a site for teratoma in our region.

DISCUSSION

Teratomas are tumors with either benign or malignant behavior. They may contain tissues of germ layers either mesodermal or endodermal or ectodermal or mixed elements. Mediastinal teratomas are an extragonadal germ cell tumor, derived from pluripotent cell with the multi-directional potential of differentiation into different types of cell. Furtherly teratomas are classified into mature (cystic or solid) teratomas, immature teratomas, and monodermal (highly specialized) teratomas. This reported case was a case of mature cystic teratoma.

The incidence of teratoma is about 1 per 4,000 live births. The various sites in decreasing order of the frequency of occurrence are as follows: Sacrococcygeal 40%, ovary 25%, other sites including neck and mediastinum 18 %, testis 12% & brain 5%. Men and women are equally affected by teratoma with an age ranging from 1 - 73 years, the average age at presentation is 28 years. This case is a superior mediastinal mature cystic teratoma in a 39 years old male.

Most cases of mediastinal mature teratoma are asymptomatic and found incidentally. When symptoms are present, it is due to the compression of adjacent structures. These include cough, dyspnea, chest pain, and respiratory distress. Occasionally the teratoma can erode and rupture into the pleural space, pericardium or tracheobronchial tree causing hemoptysis and trichoptysis which is pathognomonic for teratoma. For the case being discussed, the patient had a history of progressive exertional dyspnea and orthopnea with frequent attacks of cough for 3 years duration and feeling of pulsatile mass in left upper parasternal area in the last 4 months.

The imaging technique of choice for evaluation of an abnormal mediastinum is chest CT scan. It shows the site and extension of the mass in relation to the surrounding structures and to detect the vascularity of the mass. The typical appearance is a sharply demarcated, lobulated, heterogeneous mass and the intrinsic components of the mass, including fat, bone, soft tissue, areas...
of cystic calcifications, and fluid which is pathognomonic for diagnosing teratoma. In this case, superior mediastinal mass was reported on CT scan that was suggestive of mediastinal tumor mostly teratoma.

The treatment of choice for mediastinal teratoma is complete surgical resection. It helps in confirming the diagnosis and ensuring a long-term cure rate with little chance of recurrence. Median sternotomy is the most commonly used method because of the excellent exposure. When it affects the hemithorax, then lateral thoracotomy is the approach of choice. In this case complete surgical excision was done through left thoracotomy.

Mediastinal teratoma is usually not fatal by itself but the complications of major surgical procedures (such as pneumonectomy) or bleeding from major vessels, can cause death. In this case, complete surgical excision was done and the patient post operatively developed hoarseness of voice due to left recurrent laryngeal nerve injury which was reported as a complication in extensive left mediastinal dissection.

Malignant transformation of mature cystic teratomas (MCTs) is rare and occur in 0.17%–2% of them. Squamous cell carcinoma is the most common type of malignant transformation, and there are few reports about multiple malignancies developing in a single MCT. Extensive tissue sampling was done for this case to exclude any immature or any malignant component but it did not show any evidence of malignancy.

Though benign, it is important to diagnose mediastinal teratomas as they can cause symptoms resulting from compression of adjacent structures such as threatening respiratory distress. Mature teratomas are benign tumors and have an excellent prognosis after complete resection, and so there is no role for adjuvant radiotherapy or chemotherapy in their management.

**CONCLUSION**

Superior mediastinal teratoma has been rarely reported previously, and this is the first reported case in our locality. complete excision of the tumor is possible and curative. The patient is followed up for 6 months and he is doing well apart of mild hoarseness of voice with no evidence of recurrence.
Fig. 4: Superior mediastinal mature teratoma. Showing: a- cartilage, b- respiratory mucosa, c- skin and sebaceous glands, d-glands and fibro-collagenous tissue. (H & E stain 100X)

REFERENCES