Case Report:
Thymoma with Myasthenia Gravis in Adolescent

Prasad K Shetty,
Balaiah K,
Ganesh Nayak,
Prasanna K Shetty,
Department of Pathology, Cardiovascular Surgery, Bhagwan Mahaveer Jain Hospital, Bangalore, India and Department of Pediatrics, Vaatsalya Hospital, Gokul Road, Hubli, India.

Address For Correspondence:
Dr. Prasad K Shetty,
Department of Pathology,
Bhagwan Mahaveer Jain Hospital,
Millers Road,
Bangalore-560052, India.
E-mail: dr.pkshetty@gmail.com

Citation: Shetty PK, Balaiah K, Nayak G, Shetty PK. Thymoma with Myasthenia Gravis in Adolescent. *Online J Health Allied Scs.* 2010;9(4):16
URL: http://www.ojhas.org/issue36/2010-4-16.htm

Submitted: Nov 1, 2010; Accepted: Nov 3, 2010; Published: Jan 20, 2011

Abstract:
Thymomas are exceedingly rare in the first 20 years of life, Thymic lesions comprise approximately 2–3% of all pediatric mediastinal tumors and include thymic cysts, hyperplasia, carcinoma, and thymomas. Fewer than 30 cases in children have been described in the literature. Thymomas in adults are commonly associated with other diseases, the most frequent being myasthenia gravis. However, this association has been rarely reported in childhood. These tumors are typically aggressive, with poor outcomes. We report a case of thymoma associated with myasthenia gravis in a 16-year-old girl and review the literature.

Key Words: Mediastinal mass; Thymoma; Myasthenia Gravis

Case Report:
A 16 year old girl presented with difficulty in breathing, difficulty in swallowing and on and off vomiting since 6 days, she gives history of herpes infection 3 months back. On examination her chest and cardiovascular exam were normal. Her neurological examination revealed neck stiffness, neck flexor weakness and poor reflex, an initial chest radiograph showed an anterior mediastinal mass extending to the right of the ascending aorta. A computed tomographic (CT) scan showed a lobulated 15x12cm non-fat-containing soft tissue mass in the anterior mediastinum. There were no areas of chest wall or pericardial invasion or lymphadenopathy.

Electroneuromyography (ENMG) was normal, repetitive nerve stimulation (RNS) test revealed neuromuscular junction dysfunction, her serum anti acetylcholinesterase receptor antibody level was 9.28nmol/L (Normal range being <0.3nmol/L). Based on radiological and biochemical features patient was diagnosed as thymoma associated with myasthenia gravis and was treated with neostigmine, glycopyrrolate and plasmapheresis 3 hours/day for 7 days followed by an exploratory thoracotomy and a resection of the mediastinal mass were performed. On gross examination it was a well encapsulated mass measuring 14x11x8cms, outer surface was bossulated and smooth and cut surface was gray white solid with zig zag nodules.

Figure 1: CT scan shows lobulated mass in the anterior mediastinum.

Figure 2: Well encapsulated thymoma with bossulated smooth surface.
Microscopy showed well encapsulated tumor, tumor cells were arranged in sharply defined, angular lobules with motting and trabeculation (caused by epithelial-lymphocyte), epithelial cells were spindly with fine chromatin with occasional mitosis admixed with lymphocytes. There was no evidence of capsular invasion.

Based on the pathological findings a diagnosis of Thymoma (World Health Organization classification Type AB and Masaoka classification Stage I) was made. Postoperatively patient was continued with treatment for myasthenia gravis and was discharged on 12th post operative day.

Discussion:
The association of thymoma and MG is well described in adults, occurring in anywhere from 10% to 59% of patients in adults. However, it has been stated that this association rarely occur in children. Most mediastinal tumors in the pediatric population are either neurogenic in origin (33%) or lymphomas (41%).[1] Primary thymic lesions (such as thymic cysts, thymolipomas, and thymic hyperplasia) represent only 2.5% of mediastinal tumors [2–4], while thymomas comprise about 1%. Presenting signs are from mass effect in the mediastinum, causing shortness of breath, superior vena cava syndrome, or non-specific complaints of discomfort.[1,2]
The incidence of Myasthenia gravis in the general population is thought to be 0.5/100,000 to 3/100,000, and approximately 1% of all patients with Myasthenia gravis are children.[3] The disease is characterized by easy fatigability and fluctuating strength in skeletal muscles. Patients weaken rapidly with exercise and generally worsen as the day progresses. The most common muscles affected in children are those supplied by the cranial nerves. The defect underlying the disease is thought to be a result of an autoimmune mechanism that causes a decrease in the number of available acetylcholine receptors on the postsynaptic membrane. This is manifested by the presence of measurable antibodies to acetylcholine receptorsY2 as demonstrated in our patient.[4] The disease is counteracted by the use of anticholinesterase drugs. As many as 25% of thymomas are not detected by chest radiographs but are identified by CT scanning. CT remains an essential part of tumor staging, providing information about tumor location and size and extent of involvement of, or invasion into, surrounding structures.[5]
The World Health Organization recently developed a classification system (Table 1), although most reports follow the general principles outlined by Masaoka et al (Table 2).[6]

The relationship between thymoma and MG remains unclear. Treatment in juvenile MG includes anticholinesterase drugs and prednisone. Thymectomy has been performed on patients who were refractory to drug therapy.[7] The natural course of MG in children is not well described, although several recent reviews would suggest a nearly normal life expectancy, with approximately 80% alive at 40 years of Respiratory failure is a common cause of death in MG and was a serious problem in our patient. [9,10] Several factors may precipitate respiratory compromise, including infections, surgery, emotional stress, decrease in dose of anticholinergic medication, or use of other medications such as aminoglycosides and muscle relaxants.[11,12]

In summary, the association of MG and thymoma, although rare, does occur in the pediatric population. We report a 16-year-old girl with this association.
References: