

## A PATIENT WITH MYASTHENIA GRAVIS AND ALLERGIC RHINITIS

### *MYASTHENİA GRAVİS VE ALLERJİK RİNİTLİ BİR HASTA*

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#### **ABSTRACT**

Myasthenia gravis is an autoimmune disorder of the neuromuscular junction. In most cases, auto antibodies that are formed against acetylcholine receptor (AChR) are the leading cause of the disease. Association of Myasthenia Gravis and allergic disorders is uncommon. Herein we report a patient with myasthenia gravis and allergic rhinitis.

**Key Words:** Allergic, rhinitis, atopic, Myasthenia Gravis, prick test

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## ÖZET

Myasthenia Gravis nöromusküler bileşkenin otoimmün hastalığıdır. Asetilkolin reseptörlerine (AChR) karşı gelişen otoantikorlar hastaların çoğunda saptanan en sık nedendir. Myasthenia Gravisin allerjik hastalıklarla birlikteliği nadir bir durumdur. Biz burada myasthenia gravis ve allerjik riniti bulunan bir vakayı sunuyoruz.

**Anahtar Kelimeler:** Allerjik, rinit, atopik, myasthenia gravis, prik test

## INTRODUCTION

Myasthenia gravis (MG) is an autoimmune disorder of the neuromuscular junction (NMJ) (1). In most cases, auto antibodies directed towards the skeletal muscle acetylcholine receptor (AChR) was the leading cause of the disease. In non-AChR patients, other components of the postsynaptic muscle endplate might be the other targets of the disease (2). Association of allergic disorders and MG is uncommon. Herein we report a patient with myasthenia gravis and allergic rhinitis.

## CASE

A 20-year-old boy with a history of ocular myasthenia gravis, referred to our allergy clinic for his nasal complaints.

He was suffering from nasal blockage and itching during spring for the last five years. He had no history of cough, dispnea or wheezing. Besides he had no symptoms of conjunctivitis. His mother had a history of allergic asthma. At the age of 14, he suffered from ptosis at his left eye and diplopia. He diagnosed with ocular MG. Anti-acetylcholine receptor antibody was negative (0.1 nmol/L). Two years before he underwent thymectomy. The pathology revealed thymic hyperplasia. After surgery his complaints about myasthenia gravis were resolved. The patient was not taking any medication for MG after thymectomy.

Physical examination was normal except for allergic shiner and Dennie Morgan lines. On the chest auscultation there were no pathologic findings. A complete neurological exam revealed no abnormalities.

Laboratory tests including complete blood count and biochemical parameters were normal. The eosinophil count was  $600/\text{mm}^3$ . Routine skin prick tests with aeroallergens were positive for grass pollen and also for cat and dog epithelia and negative for all other allergens tested (mites, moulds and latex). Total IgE was high (627 IU/ml). Lung function tests were at normal range.

## DISCUSSION

The prevalence of childhood MG in Europe and North America is 10–15% of MG cases (3). It is much more common in Asian countries such as China. Disease onset age is under 15 years at 50% of patients, many with purely ocular manifestations (4). Consistent with literature, our country is at Asia and our patient diagnosed at the age of 14 with ocular manifestations.

In the literature one study indicated a significant association of allergic conjunctivitis with MG especially for ocular or seronegative MG in cases without thymoma (5). Our case was seronegative without thymoma but he had only nasal symptoms, no symptoms of conjunctivitis.

There is very limited data about association of MG with allergic disorders. Kai et al. reported a case with limb-girdle type myasthenia gravis and atopic dermatitis, whose symptoms of both disease improved after thymectomy (6). Recently a case of near fatal asthma associated with MG was published. They reported the patient's lung function test declined overall after administration of the bronchodilator suggesting respiratory muscle weakness (7).

It is reported that there is an over expression of CD23 (the low affinity receptor for IgE) in the germinal centers of the thymus of MG patients (8). The CD23 receptor (Fc $\epsilon$ RII) plays an important role in IgE-related reactions, in the control of IgE synthesis (9). Thus, CD23 may be the common immunologic factor between MG and allergic disorders.

To the best of our knowledge our case was the only case that had myasthenia gravis and allergic rhinitis in the English literature.

In conclusion, there is limited data about association of MG and allergic rhinitis at the literature. There may be a common immunologic factor between them or it could be a random situation. In the light of this case we reviewed the literature.

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