A Case of Ocular Myasthenia Gravis Presenting as Double Depressor Palsy

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A 65-year-old man who had been experiencing diplopia in front and down gaze for 15 days visited our hospital. Hypertropia was noted in the patient’s left eye, and limitation of depression was found in the adduction, primary gaze, and abduction. Brain magnetic resonance imaging showed no remarkable findings. Two weeks after the first visit, the patient complained of ptosis in the left eye. An ice test was performed and the ptosis was resolved after the test. Then, anti-acetylcholine receptor binding antibody levels were checked and found to be slightly elevated. We prescribed methylprednisolone per os 24 mg for 2 weeks, and his symptoms improved after the 2-week treatment. Five weeks after his first visit, the patient showed an ortho result in the alternate prism cover test and normal ocular movements. This may be the first case in which ocular myasthenia gravis presented as double depressor palsy, and in such cases, the possibility of ocular myasthenia gravis should be considered to rule out double depressor palsy.

Key Words: Double depressor palsy, Myasthenia gravis, Strabismus

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Discussion

Although congenital DDP patients usually do not complain of diplopia, acquired cases have diplopia commonly. DDP is a very rare condition. The relatively low incidence of this condition can be attributed to the fact that the inferior rectus muscle is supplied by the inferior branch of the 3rd cranial nerve while the superior oblique muscle is supplied by the 4th cranial nerve [2]. Etiology of DDP is not clear and it results from various conditions such as a primary paralysis of the inferior rectus muscle (congenital or acquired), a primary supranuclear palsy of depression, and secondary dysfunction of the inferior rectus due to ipsilateral superior rectus contracture [3]. To define the pathogenesis, a careful history taking is needed to rule out secondary causes including orbital wall fracture, thyroid orbitopathy and previous vertical rectus muscle surgery.

OMG is an autoimmune disorder that leads to ptosis and/or diplopia that is caused by weakness of the extraocular eye muscles, levator palpebrae superioris, and orbicularis oculi, without dysfunction of other muscles. A history of variable and fatigable ocular muscle weakness in the presence of normal pupillary function should raise the index of suspicion for OMG [2]. The ice test is a convenient, specific, and relatively sensitive technique for diagnosing myasthenia gravis [4]. In the present case, the ice test proved useful in diagnosing OMG.

In conclusion, this may be the first case in which OMG presented as DDP, and in such cases, the possibility of OMG should be considered to rule out DDP.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.
Fig. 3. Three weeks after oral steroid medication administration, the limitation of depression was completely resolved.

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