A Case of Isolated Plexiform Neurofibroma in a Patient with Myasthenia Gravis

Seung Ju Back, M.D., Dae Hun Kim, M.D., Nari Kim, M.S., Young Lee, M.D., Young Joon Seo, M.D., Jang Kyu Park, M.D., Jeung Hoon Lee, M.D.

Department of Dermatology, College of Medicine, Chungnam National University, Daejeon, Korea

We report a case of an isolated plexiform neurofibroma occurring in a patient with myasthenia gravis. A 48-year-old man presented with asymptomatic skin-colored nodules on the tip of his 4th finger. Microscopically, a plexiform neurofibroma was identified located in the dermis that appeared to originate from small superficial nerves. He had a 20-year history of treated myasthenia gravis; otherwise, his personal and family histories were unremarkable. Given that myasthenia gravis is a disorder of the peripheral nerves, plexiform neurofibromas could be associated with myasthenia gravis. However, the development of an isolated plexiform neurofibroma in a case of myasthenia gravis has not yet been reported. The occurrence of a neurofibromas in a patient with myasthenia gravis suggests a link in the pathogenesis of these two diseases. (Ann Dermatol (Seoul) 21(1) 53 ∼ 55, 2009)

Keywords: Isolated, Myasthenia gravis, Plexiform neurofibroma

INTRODUCTION

Plexiform neurofibromas, which are histologically similar to discrete neurofibromas, are benign peripheral nerve sheath tumors that involve one or more nerve fascicles often arising from the branches of a major nerve. In this tumor, the nerve is converted into a convoluted mass which has been likened to a 'bag of worms'. The plexiform neurofibroma is considered to be pathognomonic of neurofibromatosis 1 (NF1). However, a few cases of isolated plexiform neurofibroma that were not associated with NF1 have been reported.

Myasthenia gravis is a relatively rare autoimmune disorder of the peripheral nerves associated with pemphigus, alopecia areata and psoriasis. The development of an isolated plexiform neurofibroma in myasthenia gravis has not been reported. Thus, we report a rare case of an isolated plexiform neurofibroma in a patient with a history of myasthenia gravis, and discuss the possible pathogenetic relationship between the two.
dermis (Fig. 2A). The tumor was well circumscribed and surrounded by concentric bands of collagen in an 'onion skin pattern'. It consisted of faintly eosinophilic, thin, collagen bundles, myxoid changes, and occasional blood vessels (Fig. 2B). The cell nuclei were spindle- or S-shaped and quite uniform in size. The nuclear chromatin was bland without significant polymorphism or increased mitotic activity. A fairly large proportion of the tumor cells were immunopositive for S-100 protein (Fig. 2C).

**DISCUSSION**

We have described an isolated plexiform neurofibroma that developed in a man with myasthenia gravis but no stigmata of von Recklinghausen’s disease. Plexiform neurofibromas consisting of a heterogeneous mix of various cell types, including Schwann cells and fibroblasts\(^6\),\(^7\), are reported only in patients with von Recklinghausen’s disease, and are considered to be pathognomic for NF1\(^1\),\(^2\). Enzinger and Weiss\(^8\), while acknowledging the extremely high association with von Recklinghausen’s disease, were somewhat more dubious about this relationship when the neurofibromas involved small nerves in the subcutis.

**Fig. 1.** 0.7×0.5 cm sized rubbery nodule on right fourth finger tip.

**Fig. 2.** (A) Well circumscribed numerous nerve fascicles with irregular configurations embedded in cellular matrices (H&E stain, ×40). (B) Spindle shaped cells with wavy collagen fibrils in myxoid stroma (H&E stain, ×100). (C) Numerous cells of the fascicles present intensive positivity for S-100 protein (S100 stain, ×200).
Moreover, several cases of isolated plexiform neurofibroma have recently been observed in otherwise healthy patients with no family history of NF or other anomalies, suggesting that the presence of a plexiform neurofibroma does not always indicate associated NF or structural anomalies\textsuperscript{9-11}. In our case, the tumor presented on the right 4th finger tip as a small rubbery nodule that was located superficially on the dermis. It involved a small superficial nerve, and was similar to recently described cases in which an isolated plexiform neurofibroma occurred as a superficial form involving small nerves in the dermis\textsuperscript{10,11}. The patient had no personal or family history of other abnormalities suggestive of NF, but had been treated for myasthenia gravis since 1989.

Myasthenia gravis is a relatively rare autoimmune disorder of peripheral nerves in which antibodies form against acetylcholine nicotinic postsynaptic receptors at the myoneural junction\textsuperscript{12}. It is characterized by weakness and rapid fatigue of the muscles under voluntary control. It may cause double vision, drooping eyelids, difficulties with speech, chewing, swallowing and breathing, as well as limb weakness. Myasthenia gravis is associated with other autoimmune diseases, such as thyroiditis, rheumatoid arthritis, systemic lupus erythematosus and thymoma\textsuperscript{3,13}. It is also associated with skin diseases such as pemphigus, alopecia areata, vitiligo and psoriasis\textsuperscript{3,5,14}.

There has been one other reported case of neurofibroma associated with myasthenia gravis. Waugh\textsuperscript{15} reported the development of myasthenia gravis in a case of multiple NF. However, no neurofibroma developing in a case of myasthenia gravis has been reported. The relationship between the diseases may be coincidental, but is worthy of further investigation due to the fact both developed from peripheral nerves.

In myasthenia gravis, autoantibodies block the postsynaptic receptor at the myoneural junction, preventing signal transduction which results in inflammation around the junction. This inflammation tends to be worse in areas of the body subject to frequent trauma. This inflammatory process enhances cell proliferation and, ultimately, the proliferation of nerve cells occurs. Based on this hypothesis, a clear mechanism for the development of the neurofibroma on our patient’s fingertip is evident. This case seems to confirm the premise that the superficial form of plexiform neurofibroma, involving small nerves in the dermis or subcutis, is not necessarily pathognomonic for von Recklinghausen’s disease, as previously reported\textsuperscript{10,11}. Furthermore, we have reported this case because of the rare occurrence of plexiform neurofibroma in myasthenia gravis.

REFERENCES