Complete Remission of Lambert-Eaton Myasthenic Syndrome after Successful Treatment of Small-Cell Lung Cancer

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Introduction

Lambert-Eaton myasthenic syndrome (LEMS) is a rare presynaptic autoimmune disease of the neuromuscular transmission with an incidence of 0.5 per million and a prevalence of 3.4 per million [1]. It is mediated by antibodies directed mainly against P/Q type of voltage-gated calcium channels (VGCC) which are detected in about 85%-90% of the patients [2]. A tumor, mainly a small cell lung cancer (SCLC) is detected in about 60% of the patients [3,4] while other tumors (thymoma, lymphoma, leukemia, prostate cancer) are rarely diagnosed. Remission of symptoms is rare in the SCLC-LEMS. In the non-tumor-LEMS (NT-LEMS) the prognosis is favorable [5] while in the tumor LEMS (T-LEMS) it is related to tumor treatment [6].

Case Report

A 58 years old man, ex smoker, was referred for electrodiagnostic consultation because of fluctuating proximal weakness for the last 6 months, dry mouth, sexual dysfunction and mild constipation. Neurologic examination showed weakness of the proximal limb muscles and absent tendon reflexes. Compound muscle action potential (CMAP) from abductor digiti minimi (ADM) was 3.8 mV at rest and showed increment 138% (9mV) immediately after 10 seconds of maximal muscle contraction (Figure 1). Repetitive nerve stimulation (RNS) showed a decrement of CMAP 24% at 3Hz and an increment 176% at 30Hz. With the diagnosis of LEMS the patient was admitted to the hospital. SCLC was diagnosed by chest CT and biopsy (the patient had no symptoms of SCLC). VGCC antibodies were positive. The patient was transferred to an oncologic department and the tumor was treated with chemotherapy and radiotherapy. Amifampridine 10 mg x3 was prescribed. When back at home the patient felt well and referred no weakness. We saw him 3 months later. There was no muscle weakness and the tendon reflexes were reduced. Amifampridine was withdrawn and a new electrophysiological study was performed one week later. CMAP amplitude at rest from ADM was 22 mV and RNS was normal and VGCC antibodies were not found. On follow up, on July 2015, one and half year from amifampridine withdrawal, the patient is still asymptomatic.

Discussion

LEMS is a presynaptic autoimmune or paraneoplastic neuromuscular junction disorder. About 60% of patients with LEMS have a SCLC [4]. The syndrome is frequently misdiagnosed and the time to diagnosis is 0.6-40 months after the onset of the symptoms in the SCLC-LEMS [3].

Clinically it presents proximal muscle weakness, absent tendon reflexes and autonomic symptoms mainly dry mouth, erectile dysfunction and constipation. Electrophysiologically it is characterized by a triad of findings: Low amplitude of compound muscle action potential (CMAP) from ADM at rest and after 10sec of muscular contraction at diagnosis. C= RNS in ADM after cancer treatment. D= CMAP from ADM after cancer treatment. ADM= Abductor digiti minimi, RNS= Repetitive nerve stimulation.

Figure 1: A. RNS in ADM at diagnosis. B=Compound muscular action potential (CMAP) from ADM at rest and after 10sec of muscular contraction at diagnosis. C= RNS in ADM after cancer treatment. D= CMAP from ADM after cancer treatment. ADM=Abductor digiti minimi, RNS= Repetitive nerve stimulation.

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electrophysiologically and serologically) after successful treatment of the tumor. The remission of LEMS and the absence of VGCC antibodies after SCLC treatment is a clear evidence of the paraneoplastic nature of the disease. The reduction or removal of the antigen with the treatment of the SCLC may improve or completely cure the LEMS [6].

References:


