A case of inclusion body myositis with benign monoclonal gammopathy successfully responding to repeated immunoabsorption

T Nakayama, E Horiuchi, T Watanabe, S Murayama, H Nakase

Abstract
A 69 year old woman with inclusion body myositis is described. She presented with benign monoclonal gammopathy. She was resistant to steroid therapy, but responded to repeated immunoabsorption. Up to now, there has been no established therapy for inclusion body myositis, including IVIg. It is suggested that immunoabsorption could be an alternative therapy for inclusion body myositis, when it was accompanied by immunological abnormality.

(J Neurol Neurosurg Psychiatry 2000;68:230–233)

Keywords: inclusion body myositis, benign monoclonal gammopathy, steroid therapy, immunoabsorption, IgG

Inclusion body myositis is a common acquired myopathy in patients older than 50. The characteristic pathological features are inflammation, vacuolated muscle fibres, amyloid deposits and 15–18 nm tubulo filaments. Mendell et al postulated the diagnostic criteria of inclusion body myositis. They diagnosed patients who exhibited all the above pathological features as definite inclusion body myositis and if the muscle only showed inflammation without the other pathological features, as possible inclusion body myositis. Dalakas et al reported that inclusion body myositis was often associated with monoclonal gammopathies. We examined a 69 year old woman with inclusion body myositis associated with IgG-γ, BJP-γ type benign monoclonal gammopathy, which was successfully treated with repeated immunoabsorption.

Case report
A 69 year old woman developed muscle weakness from 66 years of age. There was no family history of neuromuscular disorders and no consanguinity. She was employed at a clothing shop as a deskworker. History disclosed dysesthesia on the palmer side of her right second to fourth fingers at 67. She was

Figure 1 Clinical course of the patient. MMT=Manual muscle strength test; arrows indicate immunoabsorption treatments; clinical improvements are underlined.
diagnosed with carpal tunnel syndrome from the delayed distal latency of the right median nerve shown in a nerve conduction study. Endoscopic ligament resection was performed and her dysesthesia disappeared.

At 66 years of age, she noticed weakness in her thigh muscles. At 67, she needed support standing up from a chair. At 68, she went up stairs one step at a time and she walked more slowly than before. Later she could not go up stairs and needed help with housework. She could not stand up from a squat position, so she was admitted to our hospital in September 1996.

On admission, her weight had decreased by 6 kg over 3 years. Neurological examination showed atrophy of her hip and hamstring muscles, proximal dominant muscle weakness (manual muscle strength test gave the following results: neck flexion 3, extension 4, arm 4, forearm 4, hip extension 3-, hip flexion 2+, knee extension 2+, knee flexion 2-, ankle dorsiflexion 4, ankle plantar flexion 5-), and dysesthesia at the bilateral palmer site of the second to fourth fingers. Laboratory studies showed the following: creatine kinase and lactate dehydrogenase were moderately increased to 550 IU/l and 400 IU/l, homogeneous type of antinuclear antibody was detected at 40 times dilution, IgG and BJP monoclonal gamaglobulins were detected. IgG was 650 mg/dl, IgA was 78.2 mg/dl, and IgM was 114 mg/dl. Urinalysis gave a %creatinine : creatinine ratio of 90%. Bone marrow aspiration showed hypocellular bone marrow and its plasma cell concentration was 8.2%. Bone radiography showed osteolytic lesions on the left temporal bone, the right fifth and sixth ribs and the left...
second and third ribs. Therefore, we concluded that her gammopathy was benign monoclonal gammopathy of uncertain significance (MGUS) from the Southwestern Oncology Group criteria for multiple myeloma.

Needle electromyography showed early recruitment of all examined muscles and at voluntary contraction polyphasic, low amplitude, and short duration unit potentials were revealed. At rest, a fibrillation and positive sharp wave was shown. We concluded that these were myogenic changes. Muscle CT showed severe atrophy of the bilateral gluteus maximum muscle and the hamstrings. Moderate atrophy of the paraspinal and bilateral posterior tibial muscles was also found.

Left semitendinosus muscle biopsy was performed. Haematoyxin-eosin staining showed increased connective tissue, variation in muscle fibre size, round muscle fibres in some groups, and increased intermyonuclei. We found basophilic round nuclei in some regenerating fibres; therefore, internuclear inclusions were suspected (fig 1 A). Immunochemistry showed mononuclear infiltration around one basophilic fibre, prominent CD3 positive T cells, and dominant CD 8 positive cells. Gomori-trichrome stain showed rimmed vacuoles in affected cells, and dominant CD3 positive T cells; therefore, internuclear inclusions were suspected. Electron microscopy in the cytoplasm or cytoplasmic filamentous inclusions were not disclosed. Tau-positive profiles were detected. Characteristic filamentous inclusions were not disclosed by electron microscopy in the cytoplasm or myonuclei. We concluded that the patient had inclusion body myositis by the clinical criteria Mendel et al.2

Prednisolone (50 mg/day) was given from December 1996. Muscle power increased, but on tapering prednisolone the power decreased again. The %creatinine : creatinine ratio of about 50% did not alter with the tapering of prednisolone. Steroid myopathy was excluded. In April 1997, prednisolone was tapered to 30 mg/day and methotrexate at 5 mg/week was administered. Her hip flexor power was decreased to 2- and we concluded that immunosuppressive therapy with prednisolone and methotrexate had little effect.

Next we examined monoclonal gammopathy; we started immunoabsorption at the end of May 1997, 2–3 times a month. After the fourth absorption, monoclonal gammaglobulin in the urine disappeared and that in serum reduced. After the seventh absorption her knee flexion power was increased to 3-. After the ninth absorption, she could flex the knee at a right angle, and keep her leg up in a prone position; with assistance she could stand up from a chair, and her creatine kinase concentration reduced to 50IU/l. We tapered prednisolone gradually to 20 mg/day from the seventh absorption onwards and she showed no progression of muscle weakness. She was discharged from our hospital at the end of August 1997 and we continued absorption once a month, because monoclonal gammaglobulin in her urine reappeared 3 to 4 weeks after absorption. Her clinical course is shown below (fig 2) In April 1998, she had maintained the strength of her hamstring muscles. Muscle MRI of her thigh showed muscle atrophy of the thigh similar to that before therapy; progression of muscle atrophy was not shown (fig 1 B).

Discussion

Many studies have reported myopathy associated with MGUS or multiple myeloma. Telerman-Topnet et al reported myositis with perifascicular atrophy associated with MGUS.3 Kiprov et al performed combination therapy of plasma exchange and treatment with prednisolone and cyclophosphamide for three cases of inflammatory myositis with perifascicular atrophy associated with IgG-γ type MGUS.4 The treatment produced clinical improvement in all three patients. Examination of muscle biopsy specimens by direct immunofluorescence showed linear deposits of IgG-γ along the sarcolemmal basement membrane. Muscle biopsy specimens after treatment showed no immune deposits. Sheehan-Dare and Simmons treated amyloid myopathy in association with IgG-γ chain myeloma by melphalan and prednisolone5 which resulted in the remission of both the myeloma and myopathy. Eymard et al reported that plasmapheresis and immunosuppressive agents produced partial clinical improvement of late onset rod myopathy with MGUS,6 however, therapy interruption led to remission. As shown above, reduction of serum monoclonal gammaglobulin by chemotherapy, plasma exchange, or immunoabsorption improved the myositis.

Dalakas et al reported that inclusion body myositis was often associated with monoclonal gammaglobulin,7 but there was no report of plasma exchange or immunoabsorption to inclusion body myositis with gammopathy. Chad et al reported that plasma exchange produced no improvement of inclusion body myositis in Sjögren’s syndrome8 but no control study was performed. Dalakas et al concluded that plasmapheresis was not helpful9 for inclusion body myositis because plasmapheresis was ineffective in a double blind controlled study conducted in polymyositis and dermatomyositis. Dalakas reported that IVIg treatment produced only modest improvement for inclusion body myositis10 and the efficacy of IVIg remains debatable.10

We examined inclusion body myositis with MGUS. Repeated immunoabsorption reduced urinary and serum monoclonal gammaglobulin and her muscle weakness was improved. The good response to immunoabsorption was similar to that of myositis with monoclonal gammaglobulin. It would be useful to look hard for immunological abnormalities such as MGUS or multiple myeloma in patients with inclusion body myositis and, if there, to consider immunoabsorption or plasmapheresis.

A case of inclusion body myositis with benign monoclonal gammopathy


A case of inclusion body myositis with benign monoclonal gammopathy successfully responding to repeated immunoabsorption

T Nakayama, E Horiuchi, T Watanabe, S Murayama and H Nakase

J Neurol Neurosurg Psychiatry 2000 68: 230-233
doi: 10.1136/jnnp.68.2.230