BRACHIO-CERVICAL INFLAMMATORY MYOPATHY WITH ASSOCIATED SCLERODERMA PHENOTYPE AND LUPUS SEROLOGY

Brachio-cervical inflammatory myopathy (BCIM) is a unique clinicopathologic entity characterized by neck and upper extremity weakness with relative sparing of lower extremities and commonly associated with connective tissue diseases or myasthenia gravis and serum autoantibodies (e.g., antinuclear antibody [ANA], anti–double stranded DNA [dsDNA], and anti–acetylcholine receptor).\textsuperscript{1} Muscle pathology is distinctive, with prominent B-cell infiltrates and endomysial membrane attack complex (MAC; C5b-9) deposition. Despite the detailed original series, there have been no subsequent reports (besides abstracts\textsuperscript{2,3}) demonstrating the full clinicopathologic features of BCIM.

We report an exemplary case of BCIM associated with clinical features of scleroderma and lupus serology. We review isolated reports suggestive of BCIM and contrast its pathology with other idiopathic inflammatory myopathies (IIMs).

Case report. A 31-year-old woman experienced unintentional 32-pound weight loss during pregnancy. Following delivery, the patient reported painless progressive weakness in her proximal upper extremities and dysphagia. Four months later, examination revealed thickening and soft tissue swelling in the face and distal upper extremities. There was acro-osteolysis of the fingers and abnormal dilated nail fold capillaries. There was severe bilateral, right worse than left, upper extremity weakness (2 to 3/5) with sparing of the lower extremities and moderate neck flexor and extensor weakness.

Bloodwork revealed elevated creatine kinase [CK] (5,085 IU/L) level and positive serology including ANA, extractable nuclear antigen (anti-Ro), anti-dsDNA, anti-Ro 52, and anti-Ku. Myositis-specific antibodies were negative. CT of the chest/abdomen/pelvis and MRI of the cervical spine were normal. Baseline pulmonary function tests were unremarkable. Needle electromyography of the right arm showed fibrillation potentials and early recruitment of motor unit potentials in all muscles. Muscle biopsy of the left biceps showed a florid inflammatory myopathy with the distinctive prominent B-cell infiltrates and endomysial MAC deposition (figure).

The patient was treated with high-dose IV methylprednisolone for 4 days, followed by 1 mg/kg oral prednisone and a gradual taper and azathioprine. She received monthly IV immunoglobulin. At 5-month follow-up, strength improved 4+/5 in the affected muscles. The CK level declined to 167 IU/L. Dysphagia improved, and G-tube feeding could be discontinued with resumption of solid oral diet.

Discussion. Our case is a prototypical example of BCIM, demonstrating the clinicopathologic features first described by Pestronk.\textsuperscript{1} They reported that the most commonly associated conditions were myasthenia gravis (40%) and rheumatoid arthritis (20%). Muscle biopsies showed extensive inflammatory infiltrates with at least 1 prominent CD20\textsuperscript{+} B-cell focus, major histocompatibility complex class I expression on muscle sarcolemma, and endomysial deposition of MAC (9/10). Corticosteroid treatment resulted in improvement in 7/7 patients. We extend the original description of BCIM by reporting a patient with associated features of scleroderma, including thickening of the skin in the face and upper extremities and acro-osteolysis of the fingers. Of interest, her serology showed antibodies relatively specific for systemic lupus erythematosus (anti-dsDNA), along with myositis-associated antibodies anti-Ro52 and anti-Ku.

In the literature, there have been sporadic reports compatible with BCIM. A series of 14 patients with “cervicobrachial polymyositis” described clinical features consistent with BCIM, but muscle biopsy only described interstitial mononuclear inflammatory infiltrates without elaboration.\textsuperscript{4} Consistent with our case, several reports of scleroderma have been associated with brachio-cervical weakness. One described B-cell inflammatory infiltrates on muscle biopsy but did not describe MAC deposition,\textsuperscript{5} while a series of 5 patients showed inflammatory infiltrates without elaboration.\textsuperscript{6}

In the larger series,\textsuperscript{1,4} the referring diagnoses were rarely myopathy (with motor neuron disease and myasthenia gravis being the most common), highlighting that the brachio-cervical pattern of weakness is an under-recognized presentation of inflammatory...
myopathy. This has important clinical implications, given the treatable nature of this condition.

Based on a pathologic classification of IIMs, BCIM can be placed within the category of IIM with endomysial pathology. It is distinguished from dermatomyositis by the absence of perifascicular pathology, lack of MAC deposition on capillaries, and a more prominent B-cell component. Among IIMs, endomysial MAC deposition is a distinctive feature of BCIM. The diagnosis of IIMs has become increasingly complex with a rapid discovery of myositis-specific and myositis-associated autoantibodies and more sophisticated and detailed pathologic examination of muscle biopsies. Careful clinical-serologic-pathologic correlation will be needed for future classification systems, a central requirement of which will be muscle biopsy. Pathologic findings may have treatment implications: prominent B-cell foci suggest a possible role for B-cell depleting agents such as rituximab, and a therapeutic response has been reported in 1 steroid-refractory case. Our report well demonstrates that even with myriad overlapping clinical and serologic features, muscle biopsy can reveal a very distinct pattern of pathology that defines a unique clinicopathologic entity.

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