Myopathy due to primary systemic amyloidosis

An 85 year old man presented with a 6 month history of progressive proximal muscle weakness, with associated paraesthesia, and dysphagia. Clinical examination showed signs of a peripheral neuropathy, proximal muscle wasting, and weakness of the lower limbs. Electrophysiological examination demonstrated a generalised axonal sensori-motor neuropathy, with a superimposed proximal myopathy. Creatinine kinase concentration and erythrocyte sedimentation rate were normal, as were blood lactate and carotene concentrations. A deltoid muscle biopsy showed ragged red fibres, with no features of an inflammatory process. The patient's condition deteriorated rapidly, culminating in a cardiac respiratory arrest.

On postmortem examination, there was extensive deposition of amyloid multiple organs including the heart, lungs, oesophagus, liver, spleen, kidney, prostate, and thyroid. Sections of nerves from the brachial plexus, and sural and sciatic nerves showed amyloid deposition in the vessel walls, as well as focal deposition within nerve bundles (figure A). The bone marrow appearance was consistent with multiple myeloma, with a diffuse infiltrate of atypical plasma cells containing abundant cosinophilic cytoplasmic inclusions (figure B). The plasma cells stained strongly for kappa light chains and negatively for lambda light chains, indicating monoclonality. The deltoid and quadriceps muscles showed coagulative necrosis with amyloid deposits within the vessel walls and the interstitium between muscle fibres (figure C).

Amyloid myopathy should be considered when a patient presents with progressive proximal muscle weakness, even in the absence of accompanying pseudohypertrophy of muscles, hoarseness of voice, macroglossia, or palpable nodules within muscles. Deposited amyloid is often patchy and may not be seen on a limited muscle biopsy.

ARUN AGGARWAL ALASTAIR CORBETT
Department of Neurology, Concord Repatriation General Hospital,
Hospital Road, Concord, NSW, Australia 2139

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ARUN AGGARWAL and ALASTAIR CORBETT

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