Ultrastructural Mitochondrial Abnormalities in Patients With Sporadic Amyotrophic Lateral Sclerosis

A large number of neurodegenerative diseases are caused by impairment of mitochondrial function. Mutations in genes that encode proteins responsible for the shape and dynamics of mitochondria have been associated with some genetic neurodegenerative diseases, which implies that mitochondrial shape plays an important role in the health of neurons and muscle.

Neurons are highly dependent on mitochondria because they have high energy demands and are unable to switch to glycolysis when mitochondrial oxidative phosphorylation is impaired. An ultrastructural hallmark of the synapse is the abundance of mitochondria, which are essential to maintaining calcium homeostasis and adequate levels of adenosine triphosphate (critical for nerve transmission). Neurons have extraordinarily long cellular processes, and tight control of mitochondrial dynamics facilitates the distribution of active mitochondria to dendrites and axon terminals. Higgins et al described vacuolated mitochondria in the early phases of motor neuron degeneration in transgenic mice with familial amyotrophic lateral sclerosis (ALS) and the SOD1 gene; they found that mutant SOD1 extends the outer mitochondrial membrane and expands the intermembrane space.

Much less is known about the involvement of mitochondria in muscle of patients with ALS. Defects of the mitochondrial respiratory chain have been described in several patients with ALS. Higgins et al described a patient with early-onset and rapidly progressive motor neuron disease who harbored a heteroplasmic microdeletion of the mitochondrial DNA (mtDNA)—encoded subunit 1 of cytochrome-c oxidase (COX). Finsterer described a mother and 2 daughters with symptoms compatible with ALS. All 3 patients showed COX-negative muscle fibers, ultrastructurally abnormal mitochondria, and no mutations in SOD1, but they all harbored 3 mtDNA mutations, one in the transfer RNAIle gene, one in the cytochrome b gene, and one in the adenosine triphosphatase 6 gene.

Recently, we reviewed the muscle biopsy specimens from 50 patients with typical sporadic ALS. Histochemical data showed variably severe COX deficiency in 23 of the 50 patients (46%). Of these 23 patients, 7 (30%) showed severe deficiency (≥10 COX-negative fibers of 100), and in these 7 patients, the biochemical defect of respiratory chain enzymes paralleled the histochemical defect.

Methods. To verify and extend our histochemical and biochemical data, we have now examined ultrastructurally the muscle biopsies from the 7 patients with severe...
results. We found ultrastructural mitochondrial abnormalities in 6 patients. The derangement of mitochondrial ultrastructure included disruption of the cristae and dilution of the matrix. In some cases, the mitochondria with abnormal cristae and matrices were also greatly increased in size (giant mitochondria). Also, some of the giant mitochondria showed a normal aspect. In a few mitochondria, the cristae were converted to bizarre multilamellar whorls (Figure). These ultrastructural findings support the concept that dysfunction of the oxidative metabolism may play an important role in the pathogenic pathway in sporadic ALS.

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Comentarios y Opiniones

Central Nervous System Problems With Eosinophilia

Seth and Schmidley⁷ comment in their report of 3 cases of cerebral infarcts in the setting of eosinophilia that neurological problems in hypereosinophilic syndrome (HES) include stroke, neuropathy, and encephalopathy. They note that the strokes are small and occur in the arterial border zones but may progress. Specifically, they comment that generalized encephalopathy occurs in a small percentage of patients with HES,⁸ producing behavioral disturbances and upper motor neuron signs. An early report showed that 3 patients with encephalopathy had multiple lesions in the border zones, worsened with higher eosinophil counts, and improved with treatment.⁹ Sethi and Schmidley speculate that HES encephalopathy is due to multiple small cerebral artifacts. There is a clear alternative explanation for the encephalopathy. We studied a patient with cerebrospinal fluid (CSF) eosinophilia who had reversible dementia and HES and suggested an explanation for the encephalopathy. In 1933, Gordon⁴ noted that neurotoxic effects from eosinophil-laden lymph node preparations after intracerebral inoculation produced encephalopathy, termed the Gordon phenomenon. In our case, computed tomography of the head showed periventricular white matter hypodensities, while magnetic resonance imaging (MRI) revealed confluent, increased signal around the ventricles.³

The literature cites several reports of encephalopathy with HES in which confusion, delusions, psychosis, or coma occurred.⁶ Although possible mechanisms of cerebral injury include direct eosinophil infiltration or sludging of cells in small vessels, a direct toxic effect of eosinophil-derived neurotoxin has been invoked.⁷ In our patient with CSF eosinophilia, the steroid-induced improvement in encephalopathy and dementia with reversal of CSF eosinophilia lends support to a Gordon phenomenon in humans.

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