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- e107 Helix: October 2016 issue  
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- e94 MME mutation in dominant spinocerebellar ataxia with neuropathy (SCA43)  
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- e96 KCNQ2 encephalopathy: Features, mutational hot spots, and ezogabine treatment of 11 patients  
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- e98 Complicated spastic paraplegia in patients with AP5Z1 mutations (SPG48)  
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- e97 Mendelian randomization shows a causal effect of low vitamin D on multiple sclerosis risk  
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- e105 Loss of MUNC13-1 function causes microcephaly, cortical hyperexcitability, and fatal myasthenia  
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- e102 Mutated CTSF in adult-onset neuronal ceroid lipofuscinosis and FTD  
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- e95 Novel TK2 mutations as a cause of delayed muscle maturation in mtDNA depletion syndrome  
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- e99 Novel mutation in mitochondrial DNA in 2 siblings with Leigh syndrome  
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- e101 Late-onset Lafora disease with prominent parkinsonism due to a rare mutation in EPM2A  
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- e104 GBA p.T369M substitution in Parkinson disease: polymorphism or association? A meta-analysis  
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#### CORRECTION

e111 Complicated spastic paraplegia in patients with *AP5Z1* mutations (SPG48)

 Podcast  Video  LOE classification  LOE recommendation

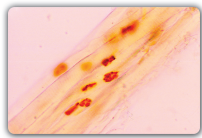


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**Cover image:** Cytochemical reaction for acetyl cholinesterase (AChE) in longitudinally oriented fibers shows pretzel-shaped or linearly arrayed endplate regions. Colorized by Kaitlyn Aman Ramm, *Neurology*<sup>®</sup> Editorial Assistant. See "Loss of *MUNC13-1* function causes microcephaly, cortical hyperexcitability, and fatal myasthenia."

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