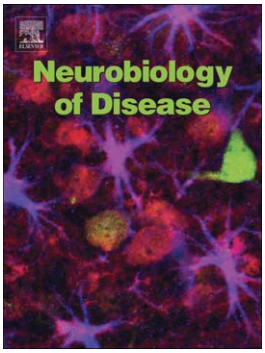

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Dynein mutations associated with hereditary motor neuropathies impair mitochondrial morphology and function with age

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Footnote:

Abbreviations

CMT	Charcot Marie Tooth disease
Cra	*Cramping* allele of the *Dync1h1* gene (p.Y1055C mutation)
IPGTT	intra-peritoneal glucose tolerance test
LPL	lipoprotein lipase
MEF	Mouse embryonic fibroblast
MFN	Mitofusin
PPARα	peroxisome proliferation activated receptor, alpha
SMA-LED	Spinal Muscular Atrophy with lower extremity dominance
UCP3	uncoupling protein 3