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## Patient 3

### IDENTIFICATION

Causal gene(s)	SPG7
Repeat size or mutation	chr16:g.89623293A > G, c.2182-2A > G (intron 16- exon 17) and c.1529C>T (exon 11)
Cell type	hiPSC

### DONOR INFORMATION

Donor gender	Male
Age at collection (years)	47

### SOURCE & PUBLICATIONS

Originating lab / institution	University of Sydney
Links to publications or public resources	Pharmacological rescue of mitochondrial and neuronal defects in SPG7 hereditary spastic paraplegia patient neurons using high throughput assays - PubMed <a href="https://pubmed.ncbi.nlm.nih.gov/37766787/">https://pubmed.ncbi.nlm.nih.gov/37766787/</a>