Cell Lines SPG7 Publication

04/009

IDENTIFICATION

Causal gene(s)	SPG7
Repeat size or mutation	c.1529C>T (het) and p.Ala510V (exon 11) and c.1745G>A and p.Gly582Asp (exonl1)
Cell type	Fibroblast

DONOR INFORMATION

Donor gender	Female
Age at collection (years)	44

SOURCE & PUBLICATIONS

Originating lab / institution	University of Sydney
Links to publications or public resources	Single cell morphology distinguishes genotype and drug effect in Hereditary Spastic Paraplegia - PubMed https://pubmed.ncbi.nlm.nih.gov/34404843/