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SCA7 KO

IDENTIFICATION

Causal gene(s)	ATXN7
Repeat size or mutation	ATXN7 Knock-out
Cell type	hiPSC

SOURCE & PUBLICATIONS

Originating lab / institution	University of California
Links to publications or public resources	Metabolic and Organelle Morphology Defects in Mice and Human Patients Define Spinocerebellar Ataxia Type 7 as a Mitochondrial Disease - PubMed https://pubmed.ncbi.nlm.nih.gov/30699348/