Cell Lines SCA14 Publication

PRKCG01

IDENTIFICATION

Causal gene(s)	RKCG
Repeat size or mutation	H36R (exon 1)
Cell type	Fibroblast

DONOR INFORMATION

Donor gender	Female
Age at collection (years)	71

SOURCE & PUBLICATIONS

Originating lab / institution	University of Oxford
Links to publications or public resources	Neurodegeneration in SCA14 is associated with increased PKCu03b3 kinase activity, mislocalization and aggregation - PubMed https://pubmed.ncbi.nlm.nih.gov/30249303/