

[Cell Lines](#) [SCA3](#) [Publication](#)

# Pa2-SCA3

## IDENTIFICATION

Causal gene(s)	ATXN3
Repeat size or mutation	78 CAG
Cell type	hiPSC

## DONOR INFORMATION

Donor gender	Male
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## SOURCE & PUBLICATIONS

Originating lab / institution	Chinese Academy of Sciences
Links to publications or public resources	CRISPR/Cas9-mediated genetic correction reverses spinocerebellar ataxia 3 disease-associated phenotypes in differentiated cerebellar neurons - PubMed <a href="https://pubmed.ncbi.nlm.nih.gov/39872157/">https://pubmed.ncbi.nlm.nih.gov/39872157/</a>