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### IDENTIFICATION

|                         |            |
|-------------------------|------------|
| Causal gene(s)          | ATXN3      |
| Repeat size or mutation | 64 CAG     |
| Cell type               | Fibroblast |

### DONOR INFORMATION

|                              |      |
|------------------------------|------|
| Donor gender                 | Male |
| Age at disease onset (years) | 39   |
| Age at collection (years)    | 45   |

### SOURCE & PUBLICATIONS

|   |   |
|---|---|
| Originating lab / institution             | University Medical Center Groningen (UMCG)  |
| Links to publications or public resources | Levels of DNAJB family members (HSP40) correlate with disease onset in patients with spinocerebellar ataxia type 3 - PubMed<br><a href="https://pubmed.ncbi.nlm.nih.gov/20726892/">https://pubmed.ncbi.nlm.nih.gov/20726892/</a><br>The HSPB8-BAG3 chaperone complex is upregulated in astrocytes in the human brain affected by protein aggregation diseases - PubMed<br><a href="https://pubmed.ncbi.nlm.nih.gov/21696420/">https://pubmed.ncbi.nlm.nih.gov/21696420/</a> |