# DISEASE:
# Spinocerebellar ataxia type 17

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| **NAME:** | Spinocerebellar ataxia type 17 |
| **DESCRIPTION:** | Spinocerebellar ataxia type 17 (SCA17) is a rare subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term). It is characterized by a variable clinical picture which can include dementia, psychiatric disorders, parkinsonism, dystonia, chorea, spasticity, and epilepsy. |
| **ORPHACODE:** | 98759 |
| **SYNONYMS:** | HDL4<br>Huntington disease-like 4<br>SCA17 |
| **XREF(S):** | Orphanet<br>OMIM<br>ICD-10 |
| **ANALYTE(S):** | TBP |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# RELATED CONTENT

## Related Genetic Tests

- Spinocerebellar ataxia (SCA) type 17 - CAG/CAA repeat expansion
- Spinocerebellar ataxia (SCA) types 8, 10, 12, 17 - repeat expansion
- Spinocerebellar ataxia (type 8, 17) + Dentatorubral pallidoluysian atrophy - repeat expansion
- Spinocerebellar ataxia (types 1, 2, 3, 6, 7, 17) - CAG repeat expansion

## Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Brussel VUB

## Related Analytes

- TATA-box binding protein

## Related Gene Panels

- Spinocerebellar ataxia (type 8, 10, 12, 17) (4 genes) - UZA
- Spinocerebellar ataxia (type 8, 17 + ATN1) (5 genes) - VUB