DISEASE:
Spinocerebellar ataxia type 2

NAME:	Spinocerebellar ataxia type 2
DESCRIPTION:	Spinocerebellar ataxia type 2 (SCA2) is a subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term) characterized by truncal ataxia, dysarthria, slowed saccades and less commonly ophthalmoparesis and chorea.
ORPHACODE:	98756
SYNONYMS:	SCA2
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	ATXN2
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RELATED CONTENT

Related Genetic Tests

- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)
- Spinocerebellar ataxia (SCA) types 1, 2, 3, 6, 7 - CAG repeat expansion
- Spinocerebellar ataxia (types 1, 2, 3, 6, 7) - CAG repeat expansion
- Spinocerebellar ataxia (types 1, 2, 3, 6, 7) - CAG 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
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- ataxin 2

Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB
- Spinocerebellar ataxia (types 1, 2, 3, 6, 7) (5 genes) - KUL
