DISEASE:
Spinocerebellar ataxia type 10

NAME:	Spinocerebellar ataxia type 10
DESCRIPTION:	Spinocerebellar ataxia type 10 (SCA10) is a subtype of type I autosomal dominant cerebellar ataxia (ADCA type I; see this term). It is characterized by slowly progressive cerebellar syndrome and epilepsy, sometimes mild pyramidal signs, peripheral neuropathy and neuropsychological disturbances.
ORPHACODE:	98761
SYNONYMS:	SCA10
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	ATXN10
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RELATED CONTENT

Related Genetic Tests

- Spinocerebellar ataxia (SCA) types 8, 10, 12, 17 - repeat expansion
- Spinocerebellar ataxia (type 8, 17) + Dentatorubral pallidoluysian atrophy - repeat expansion

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- ataxin 10

Related Gene Panels

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