DISEASE:
Dentatorubral pallidoluysian atrophy

NAME:	Dentatorubral pallidoluysian atrophy
DESCRIPTION:	A rare subtype of autosomal dominant cerebellar ataxia type I characterized by involuntary movements, ataxia, epilepsy, mental disorders, cognitive decline and prominent anticipation.
ORPHACODE:	101
SYNONYMS:	DRPLA Dentatorubropallidoluysian atrophy Naito-Oyanagi disease
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	ATN1
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Dentatorubral pallidoluysian atrophy (DRPLA) - CAG repeat expansion
- Dentatorubral pallidoluysian atrophy (DRPLA) - CAG repeat expansion
- Dentatorubral pallidoluysian atrophy - ATN1 gene CAG repeat expansion
- Spinocerebellar ataxia (type 8, 17) + Dentatorubral pallidoluysian atrophy - 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

- atrophin 1

Related Gene Panels

- Spinocerebellar ataxia (type 8, 17 + ATN1) (5 genes) - VUB

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