DISEASE:
Amyotrophic lateral sclerosis type 4

NAME:	Amyotrophic lateral sclerosis type 4
DESCRIPTION:	A rare, genetic motor neuron disease characterized by late childhood- or adolescent-onset of slowly progressive, severe, distal limb muscle weakness and wasting, in association with pyramidal signs, normal sensation, and absence of bulbar involvement, leading to degeneration of motor neurons in the brain and spinal cord.
ORPHACODE:	357043
SYNONYMS:	ALS4 Distal hereditary motor neuropathy with upper motor neuron signs dHMN with upper motor neuron signs
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	SETX
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RELATED CONTENT

Related Genetic Tests

- Amyotrophic Lateral Sclerosis (ALS) (gene panel)
- Amyotrophic lateral sclerosis (ALS) / Frontotemporal demention (FTD) - GGGGCC repeat expansion in C9ORF72
- Neurodegeneration (gene panel)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)
- Neuropathy (gene panel)

Related Laboratories

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

Related Analytes

- senataxin

Related Gene Panels

- Amyotrophic Lateral Sclerosis (ALS) - UGent
- Neurodegeneration (99 genes) - IPG
- Neuromuscular disorders (166 genes) - VUB

- Neuropathy (148 genes) - IPG

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