DISEASE:
Juvenile primary lateral sclerosis

NAME:	Juvenile primary lateral sclerosis
DESCRIPTION:	A very rare motor neuron disease characterized by progressive upper motor neuron dysfunction leading to loss of the ability to walk with wheelchair dependence, and subsequently, loss of motor speech production.
ORPHACODE:	247604
SYNONYMS:	JPLS Juvenile PLS
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	ALS2 ERLIN2
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RELATED CONTENT

Related Genetic Tests

- Amyotrophic Lateral Sclerosis (ALS) (gene panel)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)
- Spastic Paraplegia (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

- alsin Rho guanine nucleotide exchange factor ALS2
- ER lipid raft associated 2

Related Gene Panels

- Amyotrophic Lateral Sclerosis (ALS) - UGent
- Neuromuscular disorders (166 genes) - VUB
- Spastic Paraplegia (89 genes) - IPG

Source URL: http://gentest.healthdata.be/disease/844