Belgian Genetic Tests database

DISEASE:
Infantile-onset ascending hereditary spastic paralysis

NAME:	Infantile-onset ascending hereditary spastic paralysis
DESCRIPTION:	Infantile-onset ascending hereditary spastic paralysis (IAHSP) is a very rare motor neuron disease characterized by severe spasticity of the lower limbs in early life, progression of spasticity to the upper limbs in late childhood, and dysarthria.
ORPHACODE:	293168
SYNONYMS:	IAHSP
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	ALS2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/1962

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)
- Spastic Paraplegia (gene panel)

Related Laboratories

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

Related Analytes

- alsin Rho guanine nucleotide exchange factor ALS2

Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB
- Spastic Paraplegia (89 genes) - IPG