sciensano **Belgian Genetic Tests database**

GENETIC TEST:
Congenital generalized lipodystrophy type 2 / Spastic paraplegia-17 / Hereditary motor neuronopathy type VA / Silver spastic paraplegia syndrome (hot spot mutation - p.Asn88Ser; p.Ser90; p.Arg96His)

FULL NAME:	Congenital generalized lipodystrophy type 2 / Spastic paraplegia-17 / Hereditary motor neuronopathy type VA / Silver spastic paraplegia syndrome (hot spot mutation - p.Asn88Ser; p.Ser90; p.Arg96His)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Sequence analysis: entire coding region
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565390-565401
ACCREDITATION (ISO 15189):	2022-10-07 / 2027-10-06
TURNAROUND TIME (MAXIMUM):	20 - 60 days
CREATED:	07 Aug 2019 - 12:10

CHANGED:	04 Nov 2022 - 10:44

RELATED CONTENT

Related Diseases

- Autosomal dominant spastic paraplegia type 17
- Congenital generalized lipodystrophy
- Distal hereditary motor neuropathy type 5
- Severe neurodegenerative syndrome with lipodystrophy

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- BSCL2 lipid droplet biogenesis associated, seipin