
GENETIC TEST:
Neurodegeneration with Brain Iron Accumulation (gene panel)

FULL NAME:	Neurodegeneration with Brain Iron Accumulation (gene panel)
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 Whole Exome Sequencing (WES) Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565493-565504
TURNAROUND TIME (MAXIMUM):	6 months
CREATED:	31 Jul 2019 - 14:17
CHANGED:	16 Dec 2022 - 12:41

RELATED CONTENT

Related Diseases

- [Aceruloplasminemia](#)
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- [Autosomal recessive spastic paraplegia type 28](#)
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- [X-linked spastic paraplegia type 34](#)

Related Laboratories

- [Centrum Medische Genetica - UZ Gent](#)

Related Analytes

- [adaptor related protein complex 4 subunit mu 1](#)

- [ATPase cation transporting 13A2](#)
- [chromosome 19 open reading frame 12](#)
- [Coenzyme A synthase](#)
- [ceruloplasmin](#)
- [carnitine acetyltransferase](#)
- [DDB1 and CUL4 associated factor 17](#)
- [DDHD domain containing 1](#)
- [fatty acid 2-hydroxylase](#)
- [ferritin light chain](#)
- [GTP-binding protein 2](#)
- [pantothenate kinase 2](#)
- [phospholipase A2 group VI](#)
- [RALBP1 associated Eps domain containing 1](#)
- [sterol carrier protein 2](#)
- [WD repeat domain 45](#)

Related Gene Panels

- [Neurodegeneration with Brain Iron Accumulation \(NBIA\) - UGent](#)

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