ANALYTE:
NIPA1

NAME:	NIPA magnesium transporter 1
SYMBOL:	NIPA1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	MGC35570 SLC57A1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Ataxia Spasticity (gene panel)
- Cerebral palsy (gene panel)
- Epilepsy (gene panel)
- Hereditary Spastic Paraplegia (94 genes)
- Hereditary Spastic Paraplegia (gene panel)
- Hereditary spastic paraplegia (gene panel - 249 genes)
- Intellectual disability (virtual gene panel)
- Neuromuscular disorders (548 genes)
- Spastic Paraplegia (gene panel)

Related Diseases

- 15q11.2 microdeletion syndrome
- Autosomal dominant spastic paraplegia type 6

Related Gene Panels

- Ataxia Spasticity - UGent
- Cerebral palsy (212 genes) - UZA
- Hereditary Spastic Paraplegia & ataxia (genepanel) - UZA
- Hereditary Spastic Paraplegia (94 genes) - KUL
- Hereditary spastic paraplegia (188 genes) - ULB
- Intellectual disability (gene panel)
- Neuromuscular disorders (548 genes) - ULB

- Rare epilepsy with developmental delay (> 240 genes) - UZA
- Spastic Paraplegia (89 genes) - IPG
- test
