Belgian Genetic Tests database

ANALYTE:
CNNM2

NAME:	cyclin and CBS domain divalent metal cation transport mediator 2
SYMBOL:	CNNM2
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt Reactome
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RELATED CONTENT

Related Genetic Tests

- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Familial primary hypomagnesemia with normocalciuria and normocalcemia
- Primary hypomagnesemia-generalized seizures-intellectual disability-obesity syndrome

Related Gene Panels

- Early onset epileptic encephalopathy (845 genes) - ULB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

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