ANALYTE:
MAGI2

NAME:	membrane associated guanylate kinase, WW and PDZ domain containing 2
SYMBOL:	MAGI2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ACVRIP1 AIP1 ARIP1 KIAA0705 MAGI-2
XREF(S):	Orphanet Genatlas HGNC Reactome OMIM SwissProt Ensembl
CREATED:	20 Aug 2021 - 01:08
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/index.php/index.php/analyte/4724

RELATED CONTENT

Related Genetic Tests

- Early onset epileptic encephalopathy (gene panel - 845 genes)
- End-stage renal disease, ESRD (gene panel)
- Epilepsy gene panel
- Epilepsy, seizures (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)

Related Diseases

- Genetic steroid-resistant nephrotic syndrome

Related Gene Panels

- Early onset epileptic encephalopathy (845 genes) - ULB
- End-stage renal disease (106 genes) - IPG
- Epilepsy gene panel - VUB
- Epilepsy, seizures (196 genes) - IPG
- Intellectual disability (gene panel)
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG

- Panel Nephro-ULG-V1
