Belgian Genetic Tests database

ANALYTE:
ALMS1

NAME:	ALMS1 centrosome and basal body associated protein
SYMBOL:	ALMS1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	KIAA0328
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ciliopathy (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Early-onset severe obesity
- End-stage renal disease, ESRD (gene panel)
- Epilepsy gene panel
- Hepatology (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Lipodystrophy and/or hyperinsulinism (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Retinal dystrophy / RETNET (gene panel)

Related Diseases

- Alström syndrome

Related Gene Panels

- Ciliopathy (120 genes) - UGent

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Early-onset severe obesity (44 genes) - ULG
- End-stage renal disease (106 genes) - IPG
- Epilepsy gene panel - VUB
- Hepatology panel - UGent
- Heterotaxie PCD - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Lipodystrophy and/or hyperinsulinism (30 genes) - IPG
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Retinal dystrophy - UGent
