ANALYTE:
CCDC39

NAME:	coiled-coil domain containing 39
SYMBOL:	CCDC39
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CFAP59 CILD14 DKFZp434A128 FAP59
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Ciliopathy (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital structural heart defects (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Primary Ciliary Dyskinesia (gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Respiratory disorders (gene panel): non-CF bronchiectasis; pulmonary hypertension; interstitial lung disease

Related Diseases

- Primary ciliary dyskinesia

Related Gene Panels

- Ciliopathy (120 genes) - UGent
- Congenital malformation (1721 genes) - ULB
- Congenital structural heart defects - UGent
- Heterotaxie PCD - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG

- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Primary Ciliary Dyskinesia (61 genes) - KUL
- Respiratory Disorders panel (137 genes) - Ugent
