Belgian Genetic Tests database

ANALYTE:
DNMT3A

NAME:	DNA methyltransferase 3 alpha
SYMBOL:	DNMT3A
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Maffucci syndrome (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Onco-endocrine pathologies (gene panel)
- Overgrowth & vascular anomalies (gene panel)
- Overgrowth (gene panel)
- Paraganglioma and pheochromocytoma (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Sturge-Weber syndrome (gene panel)

Related Diseases

- Acute myeloid leukaemia with myelodysplasia-related features
- Sporadic pheochromocytoma/secreting paraganglioma
- Tatton-Brown-Rahman syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB

- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Maffucci syndrome (65 genes) - KUL
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Onco-endocine pathologies (50 genes) - UCL
- Overgrowth & vascular anomalies (65 genes) - KUL
- Overgrowth (24 genes) - IPG
- Paraganglioma and pheochromocytoma (29 genes) - UCL
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Sturge-Weber syndrome (65 genes) - KUL
