Belgian Genetic Tests database

ANALYTE:
MEIS2

NAME:	Meis homeobox 2
SYMBOL:	MEIS2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HsT18361 MRG1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM 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)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- 15q14 microdeletion syndrome

Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Heterotaxie PCD - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG