Belgian Genetic Tests database

ANALYTE:
ESCO2

NAME:	establishment of sister chromatid cohesion N-acetyltransferase 2
SYMBOL:	ESCO2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	EFO2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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Source URL: http://gentest.healthdata.be/analyte/1356

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Stroke (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Juberg-Hayward syndrome
- Roberts syndrome

Related Gene Panels

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

- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Stroke - UGent
