Belgian Genetic Tests database

ANALYTE:
SOX9

NAME:	SRY-box transcription factor 9
SYMBOL:	SOX9
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	SRA1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Disorders of sex development - Primary Ovarian insufficiency - Hypogonadotropic Hypogonadism (gene panel)
- Intellectual disability (virtual gene panel)
- Malformations of cortical development (235 genes)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- 46,XX ovotesticular difference of sex development
- 46,XX testicular difference of sex development
- 46,XY complete gonadal dysgenesis
- 46,XY partial gonadal dysgenesis
- Campomelic dysplasia
- Isolated Pierre Robin 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

- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - UGent
- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Intellectual disability (gene panel)
- Malformations of cortical development (235 genes) - VUB
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
