Belgian Genetic Tests database

ANALYTE:
SOX3

NAME:	SRY-box transcription factor 3
SYMBOL:	SOX3
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)
- Congenital malformation gene panel
- Disorders of sex development - Primary Ovarian insufficiency - Hypogonadotropic Hypogonadism (gene panel)
- Endocrine Disorders - Hyper(Hypo)parathyroidism (gene panel - 24 genes)
- Endocrine Disorders - Hypothyroidism (gene panel - 42 genes)
- Intellectual Disability (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Short Stature (gene panel)
- Short stature/ Growth retardation/ (gene panel)

Related Diseases

- 46,XX testicular difference of sex development
- Non-acquired panhypopituitarism
- Septo-optic dysplasia spectrum
- X-linked congenital generalized hypertrichosis
- X-linked intellectual disability with isolated growth hormone deficiency

Related Gene Panels

- Congenital malformation (1721 genes) - ULB

- Congenital malformation gene panel - VUB
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - UGent
- Endocrine Disorders - Hyper(Hypo)parathyroidism (24 genes) - ULB
- Endocrine Disorders - Hypothyroidism (42 genes) - ULB
- Growth retardation/short stature (genepanel) - UZA
- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Intellectual Disability (104 genes) - IPG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Short Stature (46 genes) - IPG
