

ANALYTE:
SHOX

NAME:	SHOX homeobox
SYMBOL:	SHOX
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	GCFX PHOG SHOXY SS
XREF(S):	<u>Orphanet</u> <u>OMIM</u> <u>OMIM</u> <u>SwissProt</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u>
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
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- [Neurodevelopmental disorders \(1300 genes\)](#)
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- [Short stature/ Growth retardation/ \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
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Related Diseases

- [Langer mesomelic dysplasia](#)
- [Léri-Weill dyschondrosteosis](#)
- [SHOX-related short stature](#)

Related Gene Panels

- [Congenital malformation \(1721 genes\) - ULB](#)
- [Growth retardation/short stature \(genepanel\) - UZA](#)

- 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
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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