

ANALYTE:
TBX4

NAME:	T-box transcription factor 4
SYMBOL:	TBX4
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
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RELATED CONTENT

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- [Congenital malformation \(gene panel - 1721 genes\)](#)
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- [Primary Arterial Hypertension \(gene panel\)](#)
- [Pulmonary Arterial Hypertension / Rendu Osler Weber disease \(gene panel - 24 genes\)](#)
- [Respiratory disorders \(gene panel\): non-CF bronchiectasis; pulmonary hypertension; interstitial lung disease](#)
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- [Skeletal dysplasia \(gene panel\)](#)

Related Diseases

- [17q23.1q23.2 microdeletion syndrome](#)
- [Coxopodopatellar syndrome](#)
- [Familial clubfoot due to 17q23.1q23.2 microduplication](#)
- [Heritable pulmonary arterial hypertension](#)

Related Gene Panels

- [Congenital malformation \(1721 genes\) - ULB](#)
- [Congenital malformation gene panel - VUB](#)
- [Intellectual disability \(gene panel\)](#)
- [Primary Arterial Hypertension \(19 genes\) - KUL](#)

- Pulmonary Arterial Hypertension (24 genes) - ULB
- Respiratory Disorders panel (137 genes) - Ugent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- chILD (34 genes) - KUL

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