

**ANALYTE:**  
**BMPR1B**

<b>NAME:</b>	bone morphogenetic protein receptor type 1B
<b>SYMBOL:</b>	BMPR1B
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	ALK6 CDw293
<b>XREF(S):</b>	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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## RELATED CONTENT

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### Related Genetic Tests

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [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](#)
- [Skeletal dysplasia \(gene panel\)](#)
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- [Skeletal dysplasia \(gene panel\)](#)

### Related Diseases

- [Acromesomelic dysplasia, Grebe type](#)
- [Brachydactyly type A1](#)
- [Brachydactyly type A2](#)
- [Brachydactyly type C](#)
- [Fibular aplasia-complex brachydactyly syndrome](#)

### 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

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