

**ANALYTE:**  
**HSPG2**

|                             |                                                                                                                           |
|-----------------------------|---------------------------------------------------------------------------------------------------------------------------|
| <b>NAME:</b>                | heparan sulfate proteoglycan 2                                                                                            |
| <b>SYMBOL:</b>              | HSPG2                                                                                                                     |
| <b>VERSION OF ORPHANET:</b> | 2023-06-22 14:14:43                                                                                                       |
| <b>SYNONYMS:</b>            | PRCAN<br>endorepellin<br>perlecan<br>perlecan proteoglycan                                                                |
| <b>XREF(S):</b>             | <u>Orphanet</u><br><u>Ensembl</u><br><u>Genatlas</u><br><u>HGNC</u><br><u>OMIM</u><br><u>Reactome</u><br><u>SwissProt</u> |
| <b>CREATED:</b>             | 13 May 2019 - 01:01                                                                                                       |
| <b>CHANGED:</b>             | 22 Jun 2023 - 16:14                                                                                                       |

## RELATED CONTENT

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### Related Genetic Tests

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Congenital structural heart defects \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Myopathy \(gene panel\)](#)
- [Neuromuscular disorders \(232 genes \(= myopathy, metabolic myopathy, ion channel muscle diseases, muscular dystrophy, myotonic dystrophy, rhabdomyolysis, myasthenia\)\)](#)
- [Neuromuscular disorders \(548 genes\)](#)
- [Neuromuscular disorders \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)

### Related Diseases

- [1p36 deletion syndrome](#)
- [Dyssegmental dysplasia, Silverman-Handmaker type](#)
- [Schwartz-Jampel syndrome](#)

### Related Gene Panels

- [Congenital malformation \(1721 genes\) - ULB](#)
- [Congenital malformation gene panel - VUB](#)

- Congenital structural heart defects - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Myopathy (332 genes) - IPG
- Neuromuscular disorders (548 genes) - ULB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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