

**ANALYTE:**  
**IDH1**

<b>NAME:</b>	isocitrate dehydrogenase (NADP(+)) 1
<b>SYMBOL:</b>	IDH1
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">Ensembl</a> <a href="#">Genatlas</a> <a href="#">HGNC</a> <a href="#">OMIM</a> <a href="#">Reactome</a> <a href="#">SwissProt</a>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	26 Oct 2023 - 23:49

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## RELATED CONTENT

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### Related Genetic Tests

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Epidermal nevus syndrome \(gene panel\)](#)
- [Maffucci syndrome \(gene panel\)](#)
- [Overgrowth & vascular anomalies \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Sturge-Weber syndrome \(gene panel\)](#)
- [Vascular malformations \(somatic\)](#)

### Related Diseases

- [Acute myeloid leukaemia with myelodysplasia-related features](#)
- [Giant cell glioblastoma](#)
- [Gliosarcoma](#)
- [Maffucci syndrome](#)
- [Metaphyseal chondromatosis with D-2-hydroxyglutaric aciduria](#)
- [Ollier disease](#)

### Related Gene Panels

- [Congenital malformation \(1721 genes\) - ULB](#)
- [Maffucci syndrome \(65 genes\) - KUL](#)
- [Overgrowth & vascular anomalies \(65 genes\) - KUL](#)

- Skeletal dysplasia (394 genes) - VUB
  - Skeletal dysplasia (genepanel) - UZA
  - Skeletal dysplasia - UGent
  - Sturge-Weber syndrome (65 genes) - KUL
  - Vascular malformations (somatic) (19 genes) - UCL
  - epidermal nevus syndrome (65 genes) - KUL
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