

**ANALYTE:**  
**TH**

<b>NAME:</b>	tyrosine hydroxylase
<b>SYMBOL:</b>	TH
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	DYT5b tyrosine 3-monooxygenase
<b>XREF(S):</b>	<u>Orphanet</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</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

- [Dystonia \(gene panel\)](#)
- [Hereditary spastic paraplegia \(gene panel - 249 genes\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Movement Disorders \(gene panel\)](#)
- [Movement Disorders \(gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)
- [Neuromuscular disorders \(548 genes\)](#)
- [Segawa syndrome \(TH gene\)](#)

### Related Diseases

- [Autosomal recessive dopa-responsive dystonia](#)

### Related Gene Panels

- [Dystonia \(68 genes\) - KUL](#)
- [Hereditary spastic paraplegia \(188 genes\) - ULB](#)
- [Intellectual disability & Epilepsy - UGent](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability/Epilepsy \(1091 genes\) - ULG](#)
- [Movement Disorders - UGent](#)

- Movement Disorders - ULG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Neuromuscular disorders (548 genes) - ULB

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