Belgian Genetic Tests database

ANALYTE:
HIBCH

NAME:	3-hydroxyisobutyryl-CoA hydrolase
SYMBOL:	HIBCH
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/3570

RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Mitochondrial disorders (gene panel)
- Movement Disorders (gene panel)
- Movement Disorders (gene panel)

Related Diseases

- Neurodegeneration due to 3-hydroxyisobutyryl-CoA hydrolase deficiency

Related Gene Panels

- Ataxia (348 genes) - ULB
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Movement Disorders - UGent
- Movement Disorders - ULG
- mitochondrial disease, nuclear based (343 genes) - VUB

Source URL: http://gentest.healthdata.be/analyte/3570