Belgian Genetic Tests database

ANALYTE:
ACTB

NAME:	actin beta
SYMBOL:	ACTB
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ß-actin
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/2019

RELATED CONTENT

Related Genetic Tests

- Becker nevus
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Dystonia (gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy (gene panel)
- Epilepsy gene panel
- Epilepsy, seizures (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Malformations of cortical development (235 genes)
- Microphthalmia / Anophthalmia / Coloboma-Anterior Segment Dysgenesis (MAC-ASD) (gene panel)
- Movement Disorders (gene panel)
- Movement Disorders (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Skeletal dysplasia (gene panel)
- Trombosis - Hemostasis (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Baraitser-Winter cerebrofrontofacial syndrome
- Becker nevus syndrome
- Developmental malformations-deafness-dystonia syndrome

Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Dystonia (68 genes) - KUL
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Epilepsy, seizures (196 genes) - IPG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Malformations of cortical development (235 genes) - VUB
- Microphthalmia/Anophthalmia/Coloboma - Anterior Segment Dysgenesis - UGent
- Movement Disorders - UGent
- Movement Disorders - ULG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Primary immune deficiencies (444 genes) - KUL
- Primary immune deficiencies - UGent
- Rare epilepsy with developmental delay (> 240 genes) - UZA
- Skeletal dysplasia (genepanel) - UZA
- Trombosis - Hemostasis (107 genes) - KUL
- test
