ANALYTE:
KAT6B

NAME:	lysine acetyltransferase 6B
SYMBOL:	KAT6B
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	MOZ-related factor MOZ2 Morf ZC2HC6B qkf querkopf
XREF(S):	Orphanet OMIM Reactome SwissProt Ensembl Genatlas HGNC
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Malformations of cortical development (235 genes)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Thyroid disgenesis (38 genes)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Blepharophimosis-intellectual disability syndrome, SBBYS type
- Genitopatellar syndrome
- Noonan 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
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Malformations of cortical development (235 genes) - VUB
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Thyroid disgenesis (38 genes) - VUB
