ANALYTE:
SETD2

NAME:	SET domain containing 2, histone lysine methyltransferase
SYMBOL:	SETD2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FLJ23184 HIF-1 HYPB KIAA1732 KMT3A
XREF(S):	Orphanet OMIM Reactome SwissProt Ensembl Genatlas HGNC
CREATED:	13 May 2019 - 01:01
CHANGED:	26 Oct 2023 - 23:49

Source URL: http://gentest.healthdata.be/index.php/index.php/analyte/862

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Early-onset severe obesity
- Epilepsy (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Neurodevelopmental disorders (1300 genes)
- Overgrowth (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Luscan-Lumish syndrome
- SETD2-related microcephaly-severe intellectual disability-multiple congenital anomalies syndrome
- Sotos syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Early-onset severe obesity (44 genes) - ULG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)

- Intellectual disability/Epilepsy (1091 genes) - ULG
- Neurodevelopmental disorders (1300 genes) - ULB
- Overgrowth (24 genes) - IPG
- Rare epilepsy with developmental delay (> 240 genes) - UZA
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- test
