ANALYTE:
KMT2A

NAME:	lysine methyltransferase 2A
SYMBOL:	KMT2A
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ALL-1 CXXC7 HRX HTRX1 Histone-lysine N-methyltransferase 2A MLL1A TRX1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Intellectual Disability (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Kabuki syndrome (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Short stature/ Growth retardation/ (gene panel)

Related Diseases

- Acute myeloid leukemia with 11q23 abnormalities
- Acute myeloid leukemia with t(9;11)(p22;q23)
- Acute undifferentiated leukemia
- B-lymphoblastic leukemia/lymphoma with recurrent genetic abnormality
- B-lymphoblastic leukemia/lymphoma with t(v;11q23.3)
- Cornelia de Lange syndrome
- Mixed phenotype acute leukemia
- Mixed phenotype acute leukemia with t(9;22)(q34.1;q11.2)
- Mixed phenotype acute leukemia with t(v;11q23.3)
- Precursor B-cell acute lymphoblastic leukemia

- Wiedemann-Steiner syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Growth retardation/short stature (genepanel) - UZA
- Intellectual Disability (104 genes) - IPG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Kabuki (7 genes) - IPG
- 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
