ANALYTE:
KANSL1

NAME:	KAT8 regulatory NSL complex subunit 1
SYMBOL:	KANSL1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CENP-36 Centromere protein 36 DKFZP727C091 MSL1v1 NSL1 centromere protein 36
XREF(S):	Orphanet OMIM Reactome SwissProt Ensembl Genatlas HGNC
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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)
- Epilepsy gene panel
- Epilepsy, seizures (gene panel)
- Intellectual Disability (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- 17q21.31 microdeletion syndrome
- Koolen-De Vries syndrome due to a point mutation

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
- Epilepsy, seizures (196 genes) - IPG
- Intellectual Disability (104 genes) - IPG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Rare epilepsy with developmental delay (> 240 genes) - UZA
- test
