ANALYTE:
SIN3A

NAME:	SIN3 transcription regulator family member A
SYMBOL:	SIN3A
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DKFZP434K2235 KIAA0700
XREF(S):	Orphanet OMIM SwissProt Ensembl Reactome Genatlas HGNC
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- 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

Related Diseases

- 15q24 microdeletion syndrome
- SIN3A-related intellectual disability syndrome due to a point mutation

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- 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

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