ANALYTE:
RAI1

NAME:	retinoic acid induced 1
SYMBOL:	RAI1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DKFZP434A139 KIAA1820 MGC12824 SMS
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Autism (gene panel)
- 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 (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)
- test

Related Diseases

- 17p11.2 microduplication syndrome
- PMP22-RAI1 contiguous gene duplication syndrome
- Smith-Magenis syndrome

Related Gene Panels

- Autism (57 genes) - IPG

- 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 (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
- test1
