

ANALYTE:
SMC1A

NAME:	structural maintenance of chromosomes 1A
SYMBOL:	SMC1A
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DXS423E KIAA0178 SB1.8 Smcb
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

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- [Early onset epileptic encephalopathy \(gene panel - 845 genes\)](#)
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Related Diseases

- [Atypical Rett syndrome](#)
- [Cornelia de Lange syndrome](#)

- Semilobar holoprosencephaly
- Wiedemann-Steiner syndrome

Related Gene Panels

- Autism (57 genes) - IPG
- Cardiomyopathy, hereditary (208 genes) - VUB
- 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
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- test

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