ANALYTE:
SMARCB1

NAME:	SWI/SNF related, matrix associated, actin dependent regulator of chromatin, subfamily b, member 1
SYMBOL:	SMARCB1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	BAF47 Ini1 PPP1R144 RDT SNF5 Sfh1p Snr1 hSNFS integrase interactor 1 malignant rhabdoid tumor suppressor protein phosphatase 1, regulatory subunit 144 sucrose nonfermenting, yeast, homolog-like 1

XREF(S):	Orphanet Reactome SwissProt Ensembl Genatlas HGNC OMIM
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Maffucci syndrome (gene panel)
- Meningioma (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Overgrowth & vascular anomalies (gene panel)
- Pediatric oncopredisposition (gene panel)
- Rhabdoid Tumor Predisposition Syndrome (SMARCA4; SMARCB1 genes)
- Schwannoma Predisposition Syndrome (gene panel)
- Short stature/ Growth retardation/ (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Sturge-Weber syndrome (gene panel)

Related Diseases

- Atypical teratoid rhabdoid tumor

- Coffin-Siris syndrome
- Familial multiple meningioma
- Full schwannomatosis
- Meningioma
- Rhabdoid tumor predisposition syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Growth retardation/short stature (genepanel) - UZA
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Maffucci syndrome (65 genes) - KUL
- Meningioma (3 genes) - KUL
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Overgrowth & vascular anomalies (65 genes) - KUL
- Pediatric oncopredisposition - UGent
- Rhabdoid Tumor Predisposition Syndrome (2 genes) - KUL
- Schwannoma Predisposition (3 genes) - KUL
- Schwannoma Predisposition Syndrome (3 genes) - KUL
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Sturge-Weber syndrome (65 genes) - KUL
