Belgian Genetic Tests database

ANALYTE:
RIN2

NAME:	Ras and Rab interactor 2
SYMBOL:	RIN2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	RASSF4
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt Reactome
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Source URL: http://gentest.healthdata.be/analyte/2318

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Cutis Laxa / Geroderma osteodysplasticum (gene panel)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Ehlers-Danlos syndroom, EDS (gene panel)
- Skin disorders (gene panel)

Related Diseases

- RIN2 syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Cutis Laxa / Geroderma osteodysplasticum - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Ehlers-Danlos syndrome -UGent
- Recessive Ehlers-Danlos Syndrome (11 genes) - UGent
- Skin disorders - UGent