Belgian Genetic Tests database

ANALYTE:
SLC39A13

NAME:	solute carrier family 39 member 13
SYMBOL:	SLC39A13
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FLJ25785
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
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Source URL: http://gentest.healthdata.be/analyte/1652

RELATED CONTENT

Related Genetic Tests

- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Ehlers-Danlos syndroom, EDS (gene panel)
- Intellectual disability (virtual gene panel)
- Skeletal dysplasia (gene panel)
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Related Diseases

- SLC39A13-related spondylodysplastic Ehlers-Danlos syndrome

Related Gene Panels

- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Ehlers-Danlos syndrome -UGent
- Intellectual disability (gene panel)
- Recessive Ehlers-Danlos Syndrome (11 genes) - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent