ANALYTE:
TREM2

NAME:	triggering receptor expressed on myeloid cells 2
SYMBOL:	TREM2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	TREM-2 Trem2a Trem2b Trem2c
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Dementia, young onset (gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Leukodystrophy (gene panel)
- Movement Disorders (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Amyotrophic lateral sclerosis
- Behavioral variant of frontotemporal dementia
- Early-onset autosomal dominant Alzheimer disease
- NON RARE IN EUROPE: Alzheimer disease
- Nasu-Hakola disease
- Progressive non-fluent aphasia
- Semantic dementia

Related Gene Panels

- Congenital malformation (1721 genes) - ULB

- Congenital malformation gene panel - VUB
- Dementia, young onset (gene panel)
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Leukodystrophy - UGent
- Movement Disorders - ULG
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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