ANALYTE:
CSF1R

NAME:	colony stimulating factor 1 receptor
SYMBOL:	CSF1R
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	C-FMS CD115 CSFR
XREF(S):	Orphanet Reactome Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Dementia, young onset (gene panel)
- Erythrocytoses, polycythémies, thrombocytoses et neutropénies congénitales (gene panel)
- Hereditary spastic paraplegia (gene panel - 249 genes)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Maffucci syndrome (gene panel)
- Movement Disorders (gene panel)
- Movement Disorders (gene panel)
- Neurodegeneration (gene panel)
- Neuromuscular disorders (548 genes)
- Overgrowth & vascular anomalies (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Sturge-Weber syndrome (gene panel)

Related Diseases

- Early-onset calcifying leukoencephalopathy-skeletal dysplasia
- Hereditary diffuse leukoencephalopathy with axonal spheroids and pigmented glia

Related Gene Panels

- Congenital malformation (1721 genes) - ULB

- Dementia, young onset (gene panel)
- Erythocyoses, polycythémies, thrombocytoses congénitales (gene panel) - ULG
- Hereditary spastic paraplegia (188 genes) - ULB
- Intellectual disability (gene panel)
- Leukodystrophy - UGent
- Maffucci syndrome (65 genes) - KUL
- Movement Disorders - UGent
- Movement Disorders - ULG
- Neurodegeneration (99 genes) - IPG
- Neuromuscular disorders (548 genes) - ULB
- Overgrowth & vascular anomalies (65 genes) - KUL
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Sturge-Weber syndrome (65 genes) - KUL
