

ANALYTE:
NF1

NAME:	neurofibromin 1
SYMBOL:	NF1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	Watson disease neurofibromatosis von Recklinghausen disease
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
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Related Gene Panels

- [Congenital malformation \(1721 genes\) - ULB](#)
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- [Epilepsy, seizures \(196 genes\) - IPG](#)
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- [Intellectual Disability \(104 genes\) - IPG](#)
- [Intellectual disability & Epilepsy - UGent](#)

- Intellectual disability (gene panel)
- Maffucci syndrome (65 genes) - KUL
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Onco-endocrine pathologies (50 genes) - UCL
- Overgrowth & vascular anomalies (65 genes) - KUL
- Paranganglioma and pheochromocytoma (29 genes) - UCL
- Pediatric oncopredisposition - UGent
- Rare epilepsy with developmental delay (> 240 genes) - UZA
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent
- Stroke - UGent
- Sturge-Weber syndrome (65 genes) - KUL
- test

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