ANALYTE:
AIFM1

NAME:	apoptosis inducing factor mitochondria associated 1
SYMBOL:	AIFM1
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SYNONYMS:	AIF CMTX4 DFNX5
XREF(S):	Orphanet OMIM SwissProt Ensembl Genatlas HGNC Reactome
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RELATED CONTENT

Related Genetic Tests

- Ataxia Spasticity (gene panel)
- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- Congenital malformation (gene panel - 1721 genes)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Hearing loss (deafness), (gene panel)
- Hereditary Spastic Paraplegia (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Mitochondrial disorders (gene panel)
- Myopathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Neuromuscular disorders (548 genes)
- Neuropathy (gene panel)
- Neuropathy (gene panel)
- Peripheral neuropathy (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Leukoencephalopathy-spondyloepimetaphyseal dysplasia syndrome
- Severe X-linked mitochondrial encephalomyopathy

- Spondyloepimetaphyseal dysplasia, Bieganski type
- X-linked Charcot-Marie-Tooth disease type 4
- X-linked hereditary sensory and autonomic neuropathy with deafness

Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL
- Ataxia Spasticity - UGent
- Congenital malformation (1721 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hearing loss (deafness) (genepanel) - UZA
- Hereditary Spastic Paraplegia & ataxia (genepanel) - UZA
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Leukodystrophy - UGent
- Myopathy (332 genes) - IPG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Neuromuscular disorders (548 genes) - ULB
- Neuropathy (148 genes) - IPG
- Neuropathy (genepanel) - UZA
- Neuropathy panel - UGent
- Skeletal dysplasia (genepanel) - UZA
- mitochondrial disease, nuclear based (343 genes) - VUB
