Belgian Genetic Tests database

ANALYTE:
ARSA

NAME:	arylsulfatase A
SYMBOL:	ARSA
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	metachromatic leucodystrophy
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Ataxia (gene panel)
- Ataxia Spasticity (gene panel)
- Cerebral palsy (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Dystonia (gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Hereditary Spastic Paraplegia (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Lysosomal Storage Disease (gene panel)
- Movement Disorders (gene panel)
- Movement Disorders (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodegeneration (gene panel)
- Neuropathy (gene panel)
- Peripheral neuropathy (gene panel)

Related Diseases

- Metachromatic leukodystrophy, adult form
- Metachromatic leukodystrophy, juvenile form

- Metachromatic leukodystrophy, late infantile form
- NON RARE IN EUROPE: Pseudoarylsulfatase A deficiency

Related Gene Panels

- Ataxia (141 genes) - KUL
- Ataxia (348 genes) - ULB
- Ataxia Spasticity - UGent
- Cerebral palsy (212 genes) - UZA
- Congenital malformation (1721 genes) - ULB
- Dystonia (68 genes) - KUL
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hereditary Spastic Paraplegia & ataxia (genepanel) - UZA
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Leukodystrophy - UGent
- Lysomal Storage (64 genes) - VUB
- Movement Disorders - UGent
- Movement Disorders - ULG
- Nephropathy panel - UGent
- Neurodegeneration (99 genes) - IPG
- Neuropathy (genepanel) - UZA
- Neuropathy panel - UGent
