Belgian Genetic Tests database

DISEASE:
Sandhoff disease, infantile form

NAME:	Sandhoff disease, infantile form
ORPHACODE:	309155
SYNONYMS:	Hexosaminidases A and B deficiency, infantile form Infantile GM2 gangliosidosis 0 variant
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	HEXB
CREATED:	13 May 2019 - 01:02
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Source URL: http://gentest.healthdata.be/disease/2092

RELATED CONTENT

Related Genetic Tests

- Dystonia (gene panel)
- Lysosomal Storage Disease (gene panel)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)
- Sandhoff disease

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- hexosaminidase subunit beta

Related Gene Panels

- Dystonia (68 genes) - KUL
- Lysomal Storage (64 genes) - VUB
- Neuromuscular disorders (166 genes) - VUB