DISEASE:
Allan-Herndon-Dudley syndrome

NAME:	Allan-Herndon-Dudley syndrome
DESCRIPTION:	An X-linked intellectual disability syndrome with neuromuscular involvement characterized by infantile hypotonia, muscular hypoplasia, spastic paraparesis with dystonic/athetoic movements, and severe cognitive deficiency.
ORPHACODE:	59
SYNONYMS:	AHDS MCT8 deficiency Monocarboxylate transporter 8 deficiency X-linked intellectual disability-hypotonia syndrome
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	SLC16A2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/606

RELATED CONTENT

Related Genetic Tests

- Spastic Paraplegia (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- solute carrier family 16 member 2

Related Gene Panels

- Spastic Paraplegia (89 genes) - IPG