

DISEASE:
Neuroferritinopathy

NAME:	Neuroferritinopathy
DESCRIPTION:	Neuroferritinopathy is a late-onset type of neurodegeneration with brain iron accumulation (NBIA; see this term) characterized by progressive chorea or dystonia and subtle cognitive deficits.
ORPHACODE:	157846
SYNONYMS:	Adult basal ganglia disease Ferritin-related neurodegeneration Hereditary ferritinopathy
XREF(S):	<u>Orphanet</u> <u>MeSH</u> <u>ICD-10</u> <u>OMIM</u>
ANALYTE(S):	<u>FTL</u>
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/disease/789>

RELATED CONTENT

Related Genetic Tests

- [Hemochromatosis \(17 genes\)](#)

Related Laboratories

- [Centre de Génétique Humaine - Erasme ULB](#)

Related Analytes

- [ferritin light chain](#)

Related Gene Panels

- [Hemochromatosis \(17 genes\) - ULB](#)

Source URL: <http://gentest.healthdata.be/disease/789>