

DISEASE:**X-linked sideroblastic anemia and spinocerebellar ataxia**

NAME:	X-linked sideroblastic anemia and spinocerebellar ataxia
DESCRIPTION:	A rare syndromic, inherited form of sideroblastic anemia characterized by mild to moderate anemia (with hypochromia and microcytosis) and early-onset, non- or slowly progressive spinocerebellar ataxia.
ORPHACODE:	2802
SYNONYMS:	Pagon-Bird-Detter syndrome X-linked sideroblastic anemia with ataxia XLSA-A
XREF(S):	<u>Orphanet</u> <u>OMIM</u> <u>ICD-10</u>
ANALYTE(S):	<u>ABCB7</u>
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/disease/1026>

RELATED CONTENT

Related Analytes

- [ATP binding cassette subfamily B member 7](#)

Source URL: <http://gentest.healthdata.be/disease/1026>