DISEASE:
Ataxia-oculomotor apraxia type 1

NAME:	Ataxia-oculomotor apraxia type 1
DESCRIPTION:	A rare autosomal recessive cerebellar ataxia, characterized by progressive cerebellar ataxia associated with oculomotor apraxia, severe neuropathy, and hypoalbuminemia.
ORPHACODE:	1168
SYNONYMS:	AOA1
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	APTX
CREATED:	13 May 2019 - 01:02
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