DISEASE:
Aceruloplasminemia

NAME:	Aceruloplasminemia
DESCRIPTION:	A rare adult-onset disorder of neurodegeneration with brain iron accumulation (NBIA) characterized by anemia, retinal degeneration, diabetes and various neurological symptoms.
ORPHACODE:	48818
SYNONYMS:	Hereditary ceruloplasmin deficiency
XREF(S):	Orphanet MeSH ICD-10 OMIM
ANALYTE(S):	CP
CREATED:	13 May 2019 - 01:02
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