DISEASE:
Hereditary hyperferritinemia-cataract syndrome

NAME:	Hereditary hyperferritinemia-cataract syndrome
DESCRIPTION:	A rare genetic disease characterized by the association of early onset cataract with persistently raised plasma ferritin concentrations in the absence of iron overload.
ORPHACODE:	163
SYNONYMS:	Bonneau-Beaumont syndrome HHCS Hereditary hyperferritinemia with congenital cataracts
XREF(S):	Orphanet OMIM ICD-10 MeSH
ANALYTE(S):	FTL
CREATED:	13 May 2019 - 01:02
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