DISEASE:
Hemoglobin H disease

NAME:	Hemoglobin H disease
DESCRIPTION:	An intermediate form of alpha-thalassemia characterized by increased hemolysis and mild to severe anemia with marked microcytosis and hypochromia. Hemoglobin H disease (HbH) disease belongs to the group of nontransfusion-dependent thalassemia.
ORPHACODE:	93616
SYNONYMS:	Alpha-thalassemia intermedia HbH disease
XREF(S):	Orphanet OMIM MedDRA ICD-10
ANALYTE(S):	HBA2 HBA1
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