DISEASE:
Glycogen storage disease due to hepatic glycogen synthase deficiency

NAME:	Glycogen storage disease due to hepatic glycogen synthase deficiency
DESCRIPTION:	A genetically inherited anomaly of glycogen metabolism and a form of glycogen storage disease (GSD) characterized by fasting hypoglycemia. This is not a glycogenosis, strictly speaking, as the enzyme deficiency decreases glycogen reserves.
ORPHACODE:	2089
SYNONYMS:	GSD due to hepatic glycogen synthase deficiency GSD type 0a Glycogen storage disease due to liver glycogen synthase deficiency Glycogen storage disease type 0a Glycogenosis type 0a
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	GYS2
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