DISEASE:
Autosomal dominant primary hypomagnesemia with hypocalciuria

NAME:	Autosomal dominant primary hypomagnesemia with hypocalciuria
DESCRIPTION:	A mild form of familial primary hypomagnesemia (FPH), characterized by extreme weakness, tetany and convulsions. Secondary disturbances in calcium excretion are observed.
ORPHACODE:	34528
SYNONYMS:	HOMG2 Isolated autosomal dominant hypomagnesemia Isolated renal magnesium wasting Renal hypomagnesemia type 2
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	FXYD2
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