DISEASE:
Primary hypomagnesemia with hypercalciuria and nephrocalcinosis without severe ocular involvement

NAME:	Primary hypomagnesemia with hypercalciuria and nephrocalcinosis without severe ocular involvement
DESCRIPTION:	Familial primary hypomagnesemia with hypercalciuria and nephrocalcinosis without severe ocular involvement (FHHN) is a form of familial primary hypomagnesemia (FPH; see this term), characterized by recurrent urinary tract infections, nephrolithiasis, bilateral nephrocalcinosis, renal magnesium (Mg) wasting, hypercalciuria and kidney failure.
ORPHACODE:	31043
SYNONYMS:	FHHNC without severe ocular involvement HOMG3 Renal hypomagnesemia type 3
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	CLDN16
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