DISEASE:
REN-related autosomal dominant tubulointerstitial kidney disease

NAME:	REN-related autosomal dominant tubulointerstitial kidney disease
DESCRIPTION:	A rare autosomal dominant tubulointerstitial kidney disease (ADTKD) of childhood due to REN mutations and characterized by early onset hypoproliferative anemia, hyperuricemia, gout, and slowly progressive tubulointerstitial kidney disease.
ORPHACODE:	217330
SYNONYMS:	ADTKD-REN FJHN type 2 Familial juvenile hyperuricemic nephropathy type 2 REN-associated FJHN REN-associated familial juvenile hyperuricemic nephropathy REN-associated kidney disease
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	REN
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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