DISEASE:
AApoAII amyloidosis

NAME:	AApoAII amyloidosis
DESCRIPTION:	A rare hereditary amyloidosis with primary renal involvement characterized by variable onset of renal insufficiency with edema, hypertension, proteinuria, and azotemia, eventually leading to end-stage renal disease. Amyloid cardiomyopathy and histopathological evidence of amyloid deposition in other organs, such as the spleen, liver, adrenal glands, and pancreas, among others, have also been described.
ORPHACODE:	238269
SYNONYMS:	Apolipoprotein A-II amyloidosis Familial amyloid nephropathy due to apolipoprotein A-II variant Familial renal amyloidosis due to apolipoprotein A-II variant Hereditary amyloid nephropathy due to apolipoprotein A-II variant Hereditary renal amyloidosis due to apolipoprotein A-II variant
XREF(S):	Orphanet ICD-10
ANALYTE(S):	APOA2
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