DISEASE:
Familial visceral myopathy

NAME:	Familial visceral myopathy
DESCRIPTION:	Familial visceral myopathy is a rare hereditary myopathic degeneration of both gastrointestinal and urinary tracts that causes chronic intestinal pseudo-obstruction. It usually presents after the first decade of life with megaduodenum, megacystis and symptoms such as abdominal distension and/or pain, vomiting, constipation, diarrhea, dysphagia, and/or urinary tract infections.n.
ORPHACODE:	2604
SYNONYMS:	Familial hollow visceral myopathy Hereditary hollow visceral myopathy Megaduodenum and/or megacystis
XREF(S):	Orphanet OMIM OMIM ICD-10
ANALYTE(S):	ACTG2
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