DISEASE:
Autosomal dominant cutis laxa

NAME:	Autosomal dominant cutis laxa
DESCRIPTION:	A rare connective tissue disorder characterized by wrinkled, redundant and sagging inelastic skin associated in some cases with internal organ involvement.
ORPHACODE:	90348
SYNONYMS:	ADCL
XREF(S):	Orphanet ICD-10 OMIM OMIM OMIM
ANALYTE(S):	ALDH18A1 FBLN5 ELN
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

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- fibulin 5

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