

DISEASE:**Body skin hyperlaxity due to vitamin K-dependent coagulation factor deficiency**

NAME:	Body skin hyperlaxity due to vitamin K-dependent coagulation factor deficiency
DESCRIPTION:	A rare genetic skin disease characterized by severe skin laxity affecting the trunk and limbs.
ORPHACODE:	91135
SYNONYMS:	PXE-like syndrome Pseudoxanthoma elasticum-like syndrome
XREF(S):	<u>Orphanet</u> <u>OMIM</u> <u>ICD-10</u>
ANALYTE(S):	<u>GGCX</u>
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Source URL: <http://gentest.healthdata.be/index.php/index.php/disease/3216>

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