
DISEASE:
Syndrome de Loeys-Dietz

NAME:	Syndrome de Loeys-Dietz
DESCRIPTION:	Loeys-Dietz syndrome is a rare genetic connective tissue disorder characterized by a broad spectrum of craniofacial, vascular and skeletal manifestations with four genetic subtypes described forming a clinical continuum.
ORPHACODE:	60030
XREF(S):	<u>ORPHANET</u>
ANALYTE(S):	<u>IPO8</u> <u>TGFBR1</u> <u>TGFBR2</u>
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