

**DISEASE:**

**Syndrome de Loeys-Dietz**

<b>NAME:</b>	Syndrome de Loeys-Dietz
<b>DESCRIPTION:</b>	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.
<b>ORPHACODE:</b>	60030
<b>XREF(S):</b>	<a href="#">ORPHANET</a>
<b>ANALYTE(S):</b>	<a href="#">IPO8</a> <a href="#">TGFBR1</a> <a href="#">TGFBR2</a>
<b>CREATED:</b>	19 Dec 2022 - 12:26
<b>CHANGED:</b>	19 Dec 2022 - 12:28

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- [Centrum Medische Genetica - UZ Gent](#)

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- [transforming growth factor beta receptor 1](#)
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- [Familial Thoracic Aortic Aneurysm \(21 genes\) - UGent](#)

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