

**DISEASE:**  
**Lymphangioleiomyomatosis**

<b>NAME:</b>	Lymphangioleiomyomatosis
<b>DESCRIPTION:</b>	A rare, multiple cystic lung disease characterized by progressive cystic destruction of the lung and lymphatic abnormalities, frequently associated with renal angiomyolipomas (AMLs).
<b>ORPHACODE:</b>	538
<b>SYNOMYS:</b>	LAM
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">OMIM</a> <a href="#">MeSH</a> <a href="#">MedDRA</a> <a href="#">ICD-10</a>
<b>ANALYTE(S):</b>	<a href="#">TSC1</a> <a href="#">TSC2</a>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

## RELATED CONTENT

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### Related Genetic Tests

- Primary lymphedema / fetal hydrops (gene panel)

### Related Laboratories

- Centre de Génétique Médicale UCL

### Related Analytes

- TSC complex subunit 1
- TSC complex subunit 2

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Source URL: <http://gentest.healthdata.be/disease/1468>