Lymphangioleiomyomatosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Lymphangioleiomyomatosis. ORPHA:538

Lymphangioleiomyomatosis (LAM) is a multiple cystic lung disease characterized by progressive cystic destruction of the lung and lymphatic abnormalities, frequently associated with renal angiomyolipomas (AMLs). LAM occurs either sporadically or as a manifestation of tuberous sclerosis complex (TSC).