Adult pulmonary Langerhans cell histiocytosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Adult pulmonary Langerhans cell histiocytosis. ORPHA:99874

Adult pulmonary Langerhans Cell Histiocytosis (PLCH) is a rare histiocytic lung disease characterized by the accumulation of Langerhans and other inflammatory cells in the small airways, resulting in the formation of nodular inflammatory lesions.