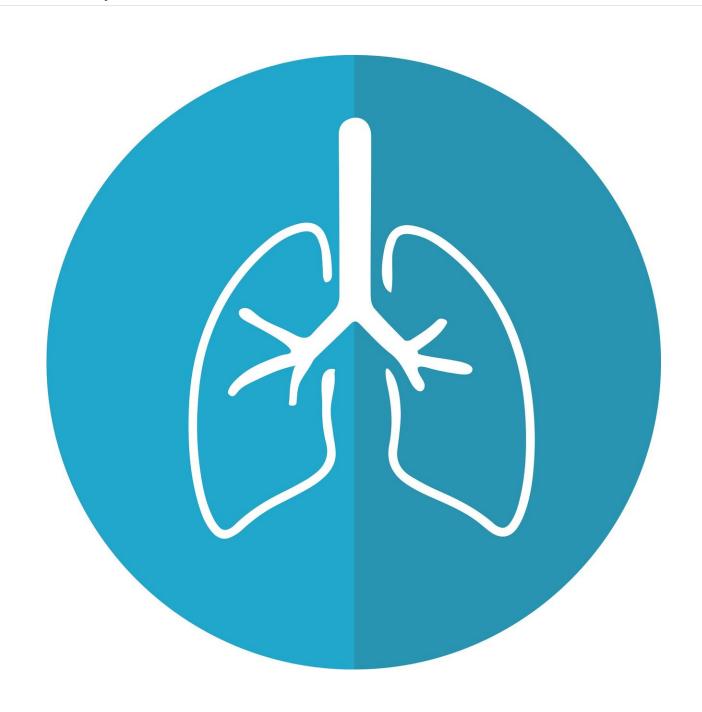


Scientists identify new features of lymphangioleiomyomatosis, a rare lung disease

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Researchers have discovered new insights about lymphangioleiomyomatosis (LAM), a rare lung disease, which affects about 1 in 200,000 Americans and commonly features the growth of abnormal cells in different tissues and organs, including the lungs.

As researchers studied LAM cells in the lab, they discovered a "mixed phenotype," or differences in physical expression in the lymphatic endothelial cells. These cells appeared to resemble cells in lymphatics as well as blood endothelial cells. The characteristics were also noted in a small percentage of lung cells provided by people living with idiopathic pulmonary fibrosis (IPF), another rare lung disease. However, the researchers didn't observe this phenotypical crossover when studying cells from healthy volunteers or from people living with Kaposi's sarcoma, a blood-related cancer.

When LAM cells accumulate in the lungs, cysts can form and impede airflow, making it hard to breathe. To help people living with LAM and other rare lung conditions, researchers have been studying physical characteristics of LAM. LAM cells in the lung reside in nodules, which are small clusters of LAM and other cells that line the cysts and may be disseminated. These lung nodules also contain lymphatics, part of the lymphatic system, which supports circulation and immune function. Lymphatic endothelial cells line the nodules and provided new insight about LAM.

Researchers are hopeful that ongoing research and future studies will lead to personalized therapies for people living with LAM and other rare



lung conditions. They also recommend studying and will continue to follow the role lymphatics may have in <u>disease progression</u>, which could support the development of new and improved therapies.

The paper, "A mixed blood-lymphatic endothelial cell phenotype in LAM and IPF but not in Kaposi's sarcoma or TSC," published Feb. 1 in the *American Journal of Respiratory Cell and Molecular Biology*.

More information: Gustavo Pacheco-Rodriguez et al, A Mixed Blood-Lymphatic Endothelial Cell Phenotype in LAM and IPF but not in Kaposi's Sarcoma or TSC, *American Journal of Respiratory Cell and Molecular Biology* (2022). DOI: 10.1165/rcmb.2021-0293LE

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