

# Lymphangioleiomyomatosis (LAM) - Pipeline Insight, 2020

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#### **Abstracts**

This report can be delivered to the clients within 24-48 Hours

DelveInsight's, "Lymphangioleiomyomatosis— Pipeline Insight, 2020," report provides comprehensive insights about 10+ companies and 10+ pipeline drugs in Lymphangioleiomyomatosis pipeline landscape. It covers the pipeline drug profiles, including clinical and nonclinical stage products. It also covers the therapeutics assessment by product type, stage, route of administration, and molecule type. It further highlights the inactive pipeline products in this space.

Geography Covered

Global coverage

Lymphangioleiomyomatosis Understanding

Lymphangioleiomyomatosis: Overview

Lymphangioleiomyomatosis (LAM) also known as lymphangiomyomatosis is a condition that affects the lungs, the kidneys, and the lymphatic system. The lymphatic system consists of a network of vessels that transport lymph fluid and immune cells throughout the body. Lymph fluid helps exchange immune cells, proteins, and other substances between the blood and tissues. LAM is found almost exclusively in women. It often occurs as a feature of an inherited syndrome called tuberous sclerosis complex. When LAM occurs alone it is called isolated or sporadic LAM.



#### **Symptoms**

Signs and symptoms of LAM most often appear during a woman's thirties. Affected women have an overgrowth of abnormal smooth muscle-like cells (LAM cells) in the lungs, resulting in the formation of lung cysts and the destruction of normal lung tissue. They may also have an accumulation of fluid in the cavity around the lungs (chylothorax). The lung abnormalities resulting from LAM may cause difficulty breathing (dyspnea), chest pain, and coughing, which may bring up blood (hemoptysis). Many women with this disorder have recurrent episodes of collapsed lung (spontaneous pneumothorax).

#### Diagnosis

Routine investigations can be supportive but not diagnostic in LAM. The chest radiograph often appears normal in early disease, although may show a pneumothorax or pleural effusion. The most common abnormalities are reticulonodular shadowing and cysts or bullae. The lung volumes are generally preserved and the combination of preserved lung volumes and interstitial changes occurs in a small number of conditions including LAM, Langerhans' cell histiocytosis, sarcoidosis and chronic hypersensitivity pneumonitis.

#### **Treatment**

Standard treatment of lymphangioleiomyomatosis is lung transplantation, but the disorder can recur in transplanted lungs. Rapamune (sirolimus) has been approved to treat lymphangioleiomyomatosis (LAM), a rare progressive lung disease that primarily affects women of childbearing age. Rapamune is manufactured by Wyeth Pharmaceuticals, Inc., a subsidiary of Pfizer, Inc. Treatment with sirolimus is recommended for patients with abnormal or declining lung function. Alternative treatments, such as hormonal manipulation with progestins, tamoxifen, and oophorectomy, are largely ineffective and not recommended.

#### Lymphangioleiomyomatosis Emerging Drugs Chapters

This segment of the Lymphangioleiomyomatosis report encloses its detailed analysis of various drugs in different stages of clinical development, including phase III, II, I and preclinical. It also helps to understand clinical trial details, expressive pharmacological action, agreements and collaborations, and the latest news and press releases.



Lymphangioleiomyomatosis Emerging Drugs

Rapamycin- Al Therapeutics

LAM-001 is the world's first inhaled mTOR inhibitor and designed to treat the rare lung disease, lymphangioleiomyomatosis. LAM is a genetic-based disease found primarily in women and characterized by hyperactivation of mTOR signaling. LAM-001 has completed clinical trials in normal healthy volunteers and in patients with LAM disease.

Saracatinib: AstraZeneca

Saracatinib is a small molecule, highly-potent and selective inhibitor of src tyrosine kinase. The drug is currently in phase II stage of development for the treatment of LAM.

Further product details are provided in the report......

Lymphangioleiomyomatosis: Therapeutic Assessment

This segment of the report provides insights about the different Lymphangioleiomyomatosis drugs segregated based on following parameters that define the scope of the report, such as:

Major Players in Lymphangioleiomyomatosis

There are approx. 10+ key companies which are developing the therapies for Lymphangioleiomyomatosis. The companies which have their Lymphangioleiomyomatosis drug candidates in the most advanced stage, i.e. phase II include, AI Therapeutics.

Phases

DelveInsight's report covers around 10+ products under different phases of clinical development like

Late stage products (BLA Filed and Phase III)



Mid-stage products (Phase II and							
Early-stage products (Phase I) along with the details of							
Pre-clinical and Discovery stage candidates							
Discontinued & Inactive candidates							
Route of Administration							
Lymphangioleiomyomatosis pipeline report provides the therapeutic assessment of the pipeline drugs by the Route of Administration. Products have been categorized under various ROAs such as							
Infusion							
Intradermal							
Intravenous							
Intravesical							
Oral etc.							
Molecule Type							
Products have been categorized under various Molecule types such as							
Antineoplastics							
Vaccine							
Gene therapies							
Immunotherapy							



/letal	
Monoclonal antibodies	
Nanoparticle	
Oncolytic viruses	
Peptide	
Plasmid	
Protein	
Small molecule	
igand	
Bacteria and others.	
Product Type	

Drugs have been categorized under various product types like Mono, Combination and Mono/Combination.

Lymphangioleiomyomatosis: Pipeline Development Activities

The report provides insights into different therapeutic candidates in phase III, II, I and preclinical stage. It also analyses Lymphangioleiomyomatosis therapeutic drugs key players involved in developing key drugs.

Pipeline Development Activities

The report covers the detailed information of collaborations, acquisition and merger, licensing along with a thorough therapeutic assessment of emerging Lymphangioleiomyomatosis drugs.

Report Highlights



The companies and academics are working to assess challenges and seek opportunities that could influence Lymphangioleiomyomatosis R&D. The therapies under development are focused on novel approaches to treat/improve Lymphangioleiomyomatosis.

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Therapeutic Assessment
Pipeline Assessment
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Unmet Needs
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Current Treatment Scenario and Emerging Therapies:
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How many Lymphangioleiomyomatosis drugs are developed by each company?

How many emerging drugs are in mid-stage, and late-stage of development for



the treatment of Lymphangioleiomyomatosis?

What are the key collaborations (Industry–Industry, Industry–Academia), Mergers and acquisitions, licensing activities related to the Lymphangioleiomyomatosis therapeutics?

What are the recent trends, drug types and novel technologies developed to overcome the limitation of existing therapies?

What are the clinical studies going on for Lymphangioleiomyomatosis and their status?

What are the key designations that have been granted to the emerging drugs?

#### **Key Players**

Al Therapeutics

AstraZeneca

#### **Key Products**

Rapamycin

Saracatinib



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**Product Development Activities** 

Saracatinib: AstraZeneca

**Product Description** 

Research and Development

**Product Development Activities** 

Drug profiles in the detailed report.....

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Comparative Analysis



Drug Name: Company name

**Product Description** 

Research and Development

**Product Development Activities** 

Drug profiles in the detailed report.....

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Comparative Analysis

Drug Name: Company name

**Product Description** 

Research and Development

**Product Development Activities** 

Drug profiles in the detailed report.....

**Inactive Products** 

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Non Muscle Invasive Bladder Cancer Key Companies

Non Muscle Invasive Bladder Cancer Key Products

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