

Clinical features, epidemiology, and therapy of lymphangioleiomyomatosis

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Abstract: Lymphangioleiomyomatosis (LAM) is a multisystem disease of women, characterized by proliferation of abnormal smooth muscle-like LAM cells, leading to the formation of lung cysts, fluid-filled cystic structures in the axial lymphatics (eg, lymphangioleiomyomas), and renal angiomyolipomas. LAM is caused by mutations of the *TSC1* or *TSC2* genes, which encode, respectively, hamartin and tuberin, two proteins with a major role in control of the mammalian target of rapamycin (mTOR) signaling pathway. LAM occurs sporadically or in association with tuberous sclerosis complex, an autosomal-dominant syndrome characterized by widespread hamartomatous lesions. LAM may present with progressive dyspnea, recurrent pneumothorax, or chylothorax. Pulmonary function tests show reduced flow rates (forced expiratory volume in the first second) and diffusion capacity. Exercise testing may reveal gas exchange abnormalities, ventilatory limitation, and hypoxemia. The severity and progression of disease may be assessed by lung histology scores, quantification of computed tomography, pulmonary function testing, 6-minute walk tests, cardiopulmonary exercise testing, and measurement of serum vascular endothelial growth factor D levels. Sirolimus and everolimus, two mTOR inhibitors, are effective in stabilizing lung function and reducing the size of chylous effusions, lymphangioleiomyomas, and angiomyolipomas. However, inhibition of mTOR complex 1 increases autophagy, possibly enhancing LAM cell survival. Inhibition of autophagy with hydroxychloroquine, in combination with sirolimus, has been proposed as a possible treatment for LAM. Deficiency of tuberin results in increased RhoA GTPase activity and cell survival, an effect that is mediated through mTOR complex 2 signaling. Because sirolimus and everolimus only affect the activity of mTOR complex 1, therapies targeting RhoA GTPases with simvastatin, which inhibits Rho GTPases and promotes apoptosis, are being investigated. As in the case of cancer, LAM may be best treated with multiple drugs targeting signaling pathways considered important in the pathogenesis of disease.

Keywords: lymphangioleiomyomatosis, tuberous sclerosis, TSC1 and TSC2 mutations, mammalian target of rapamycin signaling pathway

Introduction

Lymphangioleiomyomatosis (LAM), a multisystem disorder affecting predominantly women, is characterized by cystic lung destruction and extrapulmonary disease consisting of angiomyolipomas (AMLs), lymphatic tumors, eg, lymphangioleiomyomas, and chylous effusions.¹⁻⁶ The pathologic features of LAM result from the proliferation of neoplastic cells (LAM cells), which have characteristics of both smooth muscle cells and melanocytes.^{2,4-6} LAM occurs with increased frequency in patients with tuberous sclerosis complex (TSC), an autosomal-dominant disorder caused by mutations in the *TSC1* or *TSC2* genes, and characterized by mental retardation, autism, seizures,

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and hamartomatous lesions in the brain, heart, skin, kidney, eyes, lungs, and liver.⁷⁻¹⁰ A noninherited form of LAM called sporadic LAM is caused by somatic mutations of the *TSC2* gene⁸⁻¹⁰ and is estimated to have a prevalence of approximately 3.3–7.7 per 1,000,000 women.¹¹

Epidemiology

LAM was once considered to be a fatal disease of women of childbearing age for which there was no effective treatment except for lung transplantation.¹²⁻¹⁴ Thanks to intensive study in the past 2 decades,^{1,3,15} LAM is now considered to be a chronic disease that can affect both pre- and postmenopausal women, with a median transplant-free survival of approximately 29 years from the onset of symptoms and a 10-year transplant-free survival of 86%.^{1,15,16}

TSC is a multisystem, autosomal-dominant disorder that occurs in 1 in 12,000–14,000 children aged <10 years or 1 in 6,000 live births.⁷ TSC is characterized by facial angiofibroma, periungual fibromas, Shagreen patches, cortical tubers, cardiac rhabdomyomas, giant cell astrocytomas, mental retardation, and seizures, in addition to clinical features found in sporadic LAM.⁷ The prevalence of LAM in TSC was once thought to be 1%–4%,¹⁷⁻²⁰ but subsequent studies showed that the occurrence of cystic lung disease in women with TSC ranges from 26% to 38%.²¹⁻²³ In a recent study, it was estimated that as many as 80% of women with TSC will develop lung cysts.²⁴ Lung cysts occur in only about 13% of men with TSC,²⁵ who show much less clinically significant disease than women.

Pathology

Histologically, lung lesions consist of LAM cells in the walls of cysts and along blood vessels, lymphatics, and bronchioles, leading to narrowing of the airways, thickening of the vascular walls, lymphatic disruption, and venous occlusion.^{5,6} Noncystic lesions consist of nodular infiltrates of LAM cells. The center of the nodules contains a preponderance of small, spindle-shaped cells, whereas epithelioid cells with large cytoplasm predominate in the periphery.^{5,6} Both types of cells react with antibodies against smooth muscle antigens, eg, α -actin, vimentin, and desmin. Epithelioid cells react with human melanin black antibody (HMB-45), a monoclonal antibody that recognizes gp100, a premelanosomal protein encoded by the *Pmel17* gene.^{5,6} The spindle-shaped cells react with antiproliferation cell nuclear antigen antibodies, indicating a proliferative phenotype.^{5,6}

AMLs are characterized histologically by proliferation of smooth muscle-like LAM cells, poorly differentiated

blood vessels, and adipocytes.⁴⁻⁶ The blood supply of AML frequently originates from the renal arteries or aberrant blood vessels.^{4-6,26} Tumors may vary in size from a few millimeters to >20 cm in diameter.²⁶

Lymphangioliomyomas consist of chyle-filled lymphatic masses of varying sizes that are most frequently localized to the retroperitoneum, pelvis, and posterior mediastinum.^{4-6,26} Proliferation of epithelioid cells in the walls of lymphatic vessels leads to obstruction of chyle flow and formation of chyle-filled masses.^{4-6,26}

Clinical presentation

Patients often present with a history of recurrent pneumothoraces or progressive dyspnea (Table 1).¹⁻³ Less commonly, the first sign of LAM is a chylous effusion, hemoptysis, abdominal hemorrhage originating from an AML, or the discovery of abdominal or pelvic tumor masses.¹⁻⁶ Pneumothoraces occur in about 50%–60% of patients and tend to recur.¹⁻³ Patients with larger cysts are more likely to present with pneumothoraces.²⁷ Another frequent mode of presentation is dyspnea on exertion, which worsens with time. These patients show more advanced disease than those presenting with pneumothorax, perhaps because the occurrence of a pneumothorax prompts lung imaging studies that uncover the presence of lung cysts and lead to diagnosis.¹⁵

Other modes of presentation include chylothorax, chylous ascites, hemoptysis, chyluria, chyloptysis, abdominal lymphangioliomyomas suggesting malignancy, and abdominal hemorrhage caused by renal AMLs (Table 1).^{1,4,26}

Lymphatic involvement in LAM occurs in the posterior mediastinal, retroperitoneal, and pelvic areas and consists of lymphangioliomyomas, lymphadenopathy, pleural effusions, and ascites.^{4,26,28}

Table 1 Clinical features of LAM

Presenting signs or symptoms	Sporadic LAM	TSC-LAM
Pneumothorax	57%	47%
Dyspnea on exertion	73%	71%
Abdominal angiomyolipomas	32%	93%
Pleural effusions	12%	6%
Lymphangioliomyomas	29%	9%
Hemoptysis	32%	21%
Ascites	10%	6%

Note: Adapted with permission from Ryu JH, Moss J, Beck GJ, et al. The NHLBI Lymphangioliomyomatosis Registry. Characteristics of 230 patients at enrollment. *Am J Respir Crit Care Med.* 2006;173(1):105–111. Copyright © 2006 American Thoracic Society.¹ And Avila NA, Dwyer AJ, Rabel A, Moss J. Sporadic lymphangioliomyomatosis and tuberous sclerosis complex with lymphangioliomyomatosis, comparison of CT features. *Radiology.* 2007;242(1):277–285. Copyright © 2007 Radiological Society of North America.²⁸

Abbreviations: LAM, lymphangioliomyomatosis; TSC, tuberous sclerosis complex.



Figure 1 Skin manifestations of tuberous sclerosis complex in adult women.
Notes: (A) Two hypomelanotic macules located on the back. (B) Facial angiofibromas involving the nose and adjacent cheek. (C) Shagreen patch involving the lower back. (D) Periungual fibroma.

In patients with TSC, typical skin lesions include Shagreen patches and facial and periungual angiofibromas (Figure 1); neurologic involvement includes tubers and giant cell astrocytomas.⁷ The prevalence of extrapulmonary LAM in TSC-LAM patients is different from that in patients with sporadic LAM. In a study comparing computed tomography (CT) scan features of TSC-LAM and sporadic LAM patients, AMLs were found in approximately 93% of patients with TSC and LAM, and 32% of those with sporadic LAM (Table 1).^{1,4,28} Lymphangiomyomas and chylous effusions were more common in sporadic LAM.²⁸ The number of sclerotic bone lesions was greater in patients with TSC-LAM than in those with sporadic LAM.²⁸ Lung disease in TSC may consist of only a few cysts scattered throughout the lungs. These patients may have no clinical symptoms and are usually diagnosed by CT scans. However, those patients with TSC who initially present with pulmonary symptoms may have a clinical course similar to patients with sporadic LAM.²⁴

Pathogenesis

LAM in the presence of TSC is caused by proliferation of neoplastic LAM cells that have mutations or deletions in one of the tuberous sclerosis genes, *TSC1* localized on chromosome 9q34 or, more frequently, *TSC2*, which is localized on chromosome 16p13.^{8–10,29} Only *TSC2* abnormalities have been found to be associated with sporadic LAM.^{8–10,29} Loss of heterozygosity of *TSC2* has been demonstrated in LAM cells isolated from lung, AMLs, blood, chyle, and urine from patients with sporadic LAM and TSC-LAM.^{8–10,30,31} LAM cell clusters, consisting of LAM cell aggregates covered by lymphatic endothelial cells, have been identified in chylous

fluid.³² Although LAM cells exhibit low-grade features of malignancy, they have metastatic properties. Indeed, LAM cells have been detected in donor lungs of patients who had lung transplantation, suggesting migration to the lungs of cells from other sites, such as the kidney, lymphatic system, or uterus.^{33–35} Identical *TSC2* mutations have been found in the lungs and kidneys of the same patient with sporadic LAM.⁹

Dysregulation of the mammalian target of rapamycin (mTOR) signaling pathway is the cause of abnormal LAM cell proliferation. *TSC1* and *TSC2* genes encode, respectively, hamartin and tuberin (Figure 2),^{36–39} two proteins that regulate the intracellular serine/threonine kinase mTOR signaling path-

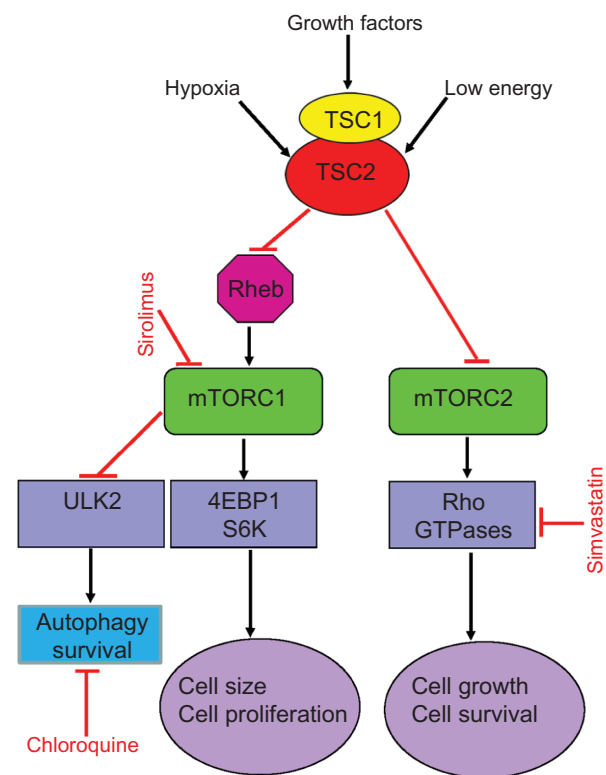


Figure 2 Simplified scheme of the mTOR signaling pathways showing potential sites of action of some agents tested or in use in LAM.

Notes: *TSC1/2* integrates multiple signals, such as those from growth factors, energy state, and hypoxia, to control cell size and proliferation. *TSC1/2* regulates mTORC1 negatively through its actions on Rheb. Activation of mTORC1 leads to cell growth and proliferation. mTORC1 is also a major regulator of autophagy. Blockade of mTORC1 by sirolimus augments autophagy, which may lead to increased cell survival. In vitro, this effect can be inhibited by chloroquine. mTORC2 regulates the actin cytoskeleton through Rho GTPases, which affect cell migration, morphogenesis, and apoptosis. Simvastatin reduces Rheb and Rho activities and promotes apoptosis. Combined therapy with sirolimus, chloroquine, or simvastatin may act synergically to inhibit LAM cell growth. Absence of *TSC1/2*, eg, hamartin and tuberin, results in loss of Rheb inhibition, leading to activation of mTORC1, enhanced 4EBP1/S6K signaling, and increased cell size and proliferation. Sirolimus blocks mTORC1, decreasing cell size and proliferation.

Abbreviations: mTOR, mammalian target of rapamycin; LAM, lymphangiomyomatosis; TSC, tuberous sclerosis complex; mTORC1, mTOR complex 1; mTORC2, mTOR complex 2; Rheb, Ras homologue enriched in brain; 4EBP1, factor 4E-binding protein 1; S6K, S6 kinase 1; ULK2, Unc-51-like kinase 2.

way. mTOR regulates cell size, proliferation, and survival by integrating signals from growth factors, energy, and stress.^{36,37} Tuberin is a GTPase-activating protein for the guanine nucleotide-binding protein Rheb (Ras homologue enriched in brain), which promotes the formation of inactive Rheb-GDP from active Rheb-GTP.³⁶ Inhibition or absence of tuberin, as occurs with *TSC2* gene mutations, results in accumulation of active Rheb-GTP, stimulation of mTOR, phosphorylation of S6 kinase and eukaryotic initiation factor 4E-binding protein, and increased translation, cell size, and proliferation.³⁷

Since LAM occurs almost exclusively in women, a role of estrogens in the pathogenesis of LAM was suspected.² Further, estrogen and progesterone receptors were seen in lung nodules and AML,^{40–42} predominance of LAM was observed in premenopausal women,¹ and worsening of pulmonary symptoms was observed during pregnancy or following the administration of exogenous estrogens.^{43,44} Moreover, the rate of decline in lung function tends to be greater in premenopausal women than in postmenopausal women.⁴⁵ Furthermore, in vitro and animal data have shown that estrogens promote the proliferation of TSC-null rat ELT3 leiomyoma-derived cells and human AML *TSC2*^{-/-} cells in vitro and the growth of subcutaneous tumors in animal models of TSC.^{46–49} However, there is yet no definite proof that antiestrogen therapies are effective in the treatment of LAM.⁴⁵

The mechanisms by which LAM cell proliferation causes formation of lung cysts and destruction of the lung parenchyma are under investigation.^{29,50} Compression of the airways by LAM cells, leading to distention of the terminal airspaces upstream from the occluded airway, has been proposed as a cause of cyst formation.^{2,5,6} Another potential cause of cystic lesions is degradation of lung elastic fibers caused by proteinases.^{29,50} This degradation may result from an increase in matrix metalloproteinase (MMP) activity in the lungs, where they have an important role in lung remodeling and lymphangiogenesis.^{50–52} LAM nodules contain MMP2, MMP9, MMP1, MMP activators (MT1-MMP), and MMP inhibitors (TIMPs).^{51–53} A reduction in levels of an MMP inhibitor (TIMP-3) was reported in LAM lesions,^{53,54} and serum levels of MMP-9 were reported to be elevated in patients with LAM.⁵⁵ An imbalance between MMP and MMP inhibitors may be a contributing factor to destruction of the lung parenchyma.⁵⁶

Disorganized lymphangiogenesis has an important role in the pathogenesis of LAM.^{32,50,57–60} Lymphatic spaces are present in LAM lung nodules that have, in addition, immunoreactivity toward vascular endothelial growth factor (VEGF)-C and VEGF-D.^{32,50,57–60} Vascular endothelial growth factor receptor (VEGFR)-3 and other markers of lymphatic

endothelial cells are also noted in the channels found in the lung nodules.^{50,57–60}

Physiologic features

According to the National Heart, Lung, and Blood Institute Lymphangioliomyomatosis Registry, airflow obstruction was seen initially in approximately 61% of patients with sporadic LAM.^{1,45} Normal spirometry was present in about 31% of patients with sporadic LAM, whereas normal lung function was observed in 53% of the patients with TSC-LAM (Table 2).¹ Air trapping may be associated with severe airflow obstruction. Reduced lung diffusing capacity (DL_{CO}) was reported in about 57% of patients with sporadic LAM,¹ with some patients having reduced DL_{CO} and normal flow rates.⁴⁵

Gas exchange abnormalities, especially during exercise, are frequently observed in LAM (Table 3).^{61,62} These abnormalities are characterized by reduced exercise capacity, decreased peak oxygen uptake (VO_2 max), reduced breathing reserve, increased ventilatory equivalent for carbon dioxide (VE/VCO_2), and hypoxemia (Table 3).^{61,62}

Airflow limitation in LAM had been attributed to alveolar destruction with consequent loss of elastic recoil.⁶³ Studies of lung mechanics showed, however, that airway resistance was increased and lung elastic recoil was not significantly reduced.⁶⁴ The primary causes of dyspnea and exercise limitation are reduced breathing reserve, dynamic hyperinflation, and an exaggerated ventilatory response to exercise because of limitation in oxygen transfer due to loss of alveolar capillary surface area.^{61,62,65} Exercise-induced pulmonary hypertension seen in patients with severe disease may also be a factor in reducing oxygen transfer.⁶⁶

Radiologic findings

Chest radiographic findings may be normal or show an interstitial pattern or cystic changes. CT shows well-defined,

Table 2 Pulmonary function abnormalities in LAM

Functional abnormality	Sporadic LAM	TSC-LAM
Decreased FEV_1	61%	38%
Decreased FEV_1/FVC ratio	61%	38%
Decreased DL_{CO}	57%	38%
Increased RV/TLC ratio	7%	3%
Normal lung function	30%	53%

Note: Adapted with permission from Ryu JH, Moss J, Beck GJ, et al. The NHLBI Lymphangioliomyomatosis Registry. Characteristics of 230 patients at enrollment. *Am J Respir Crit Care Med.* 2006;173(1):105–111. Copyright © 2006 American Thoracic Society.¹

Abbreviations: LAM, lymphangioliomyomatosis; TSC, tuberous sclerosis complex; FEV_1 , forced expiratory volume in the first second; FVC, forced vital capacity; DL_{CO} , lung diffusion capacity for carbon monoxide; RV, residual volume; TLC, total lung capacity.

Table 3 Cardiopulmonary exercise abnormalities in LAM

Decreased VO ₂ maximum
Decreased work rate
Decreased breathing reserve
Decreased SaO ₂
Increased VE/VCO ₂ at AT
Increased skeletal muscle fatigue

Abbreviations: LAM, lymphangioleiomyomatosis; VO₂ max, peak oxygen uptake; SaO₂, arterial oxygen tension; VE/VCO₂, ventilatory equivalent for CO₂; AT, anaerobic threshold.

round, thin-walled cysts scattered throughout the lungs, which may vary in size from a few millimeters to 2 cm (Figure 3).^{67–69} In patients with lymphatic involvement, pleural effusions and lung infiltrates due to chyle in the interstitium may be present. There is correlation between the extent of the cystic destruction, pulmonary function, and exercise performance.^{62,67–72}

AMLs, lymphadenopathy, lymphangioleiomyomas, and ascites may be visualized by ultrasonography, CT scans, or magnetic resonance imaging.^{26,28} AMLs occur predominantly in the kidney and liver, appearing as tumor masses comprising fatty areas, mixed with kidney parenchyma (Figure 3).^{26,28} Atypical AMLs lack fatty tissue, comprising predominantly epithelioid LAM cells that may suggest renal cell carcinoma.^{4–6,73}

Lymphangioleiomyomas appear as well-circumscribed masses of variable dimensions, comprising an outer wall and a fluid-rich region central region (Figure 3).^{26,28,74–76} The size

of lymphangioleiomyomas tends to be greater in the evening than in the morning, which may help differentiating them from malignant tumors.^{75,76}

Diagnosis

The diagnosis of LAM should be strongly suspected in any woman who presents with progressive dyspnea, recurrent pneumothoraces, or chylous pleural effusions.^{1,4,6} The diagnosis of definite LAM may be made in the presence of a characteristic CT scan and a transbronchial, thoracoscopic, or open lung biopsy^{29,77–79} showing the characteristic histological features of LAM and reactivity with monoclonal antibody HMB-45.^{4–6} Alternatively, the presence of a characteristic lung CT and AML, chylous effusions, lymphangioleiomyoma, or TSC are sufficient to establish the diagnosis.⁷⁹ The presence of a characteristic CT scan in the absence of extrapulmonary findings is not diagnostic of LAM.⁷⁹

An important biomarker of value in diagnosing LAM is VEGF-D, a lymphangiogenic growth factor that is increased in the serum of patients with LAM, especially those with lymphatic involvement.^{58–60} In the appropriate clinical and radiologic setting, a VEGF-D serum level ≥ 800 pg/mL is rarely found in other cystic lung diseases.⁶⁰ VEGF-D may be of value in grading the severity of disease and monitoring therapeutic response to mTOR inhibitors.⁶⁰

The differential diagnosis of LAM includes emphysema, sarcoidosis, Langerhans cell histiocytosis, Birt–Hogg–Dubé

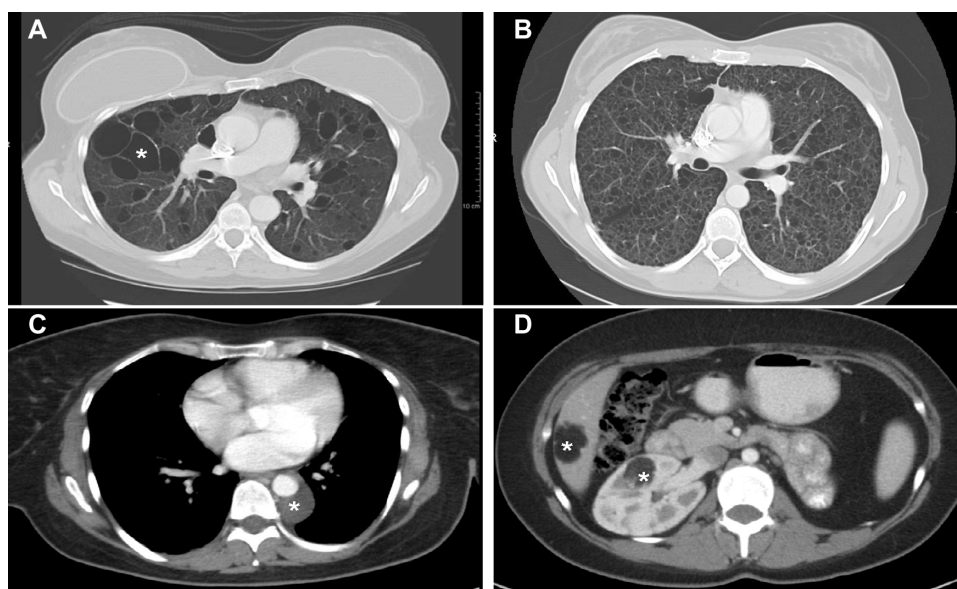


Figure 3 Computed tomography scans showing pulmonary and extrapulmonary images of patients with LAM.

Notes: (A) Multiple large (white asterisk) thin-walled cysts scattered throughout the lungs. Between the cystic areas there is normal-appearing lung parenchyma. (B) Numerous small thin-walled cysts have completely replaced the normal lung parenchyma. (C) Small posterior mediastinal lymphangioleiomyoma surrounding the descending aorta (white asterisk). (D) Right renal (white asterisk) and liver (white asterisk) angiomyolipomas in a patient with TSC-LAM. The fatty, low-density component is clearly visualized.

Abbreviations: LAM, lymphangioleiomyomatosis; TSC, tuberous sclerosis complex.

syndrome, follicular bronchiolitis, light chain disease, Sjögren's syndrome, allergic alveolitis, bronchiectasis, Mounier-Kuhn syndrome, pneumoconiosis, and chronic lung infectious processes. In general, most of these conditions can be easily excluded on the basis of clinical, histopathological, and radiologic findings.^{2,79}

Prognosis

Clinical, pathologic, and physiologic data may assist in estimating the prognosis of newly diagnosed patients with LAM. Premenopausal patients may have a faster decline in lung function than older, postmenopausal women.^{45,80} Patients who give a history of pneumothorax as the first sign of LAM present with better lung function than those experiencing dyspnea on exertion.^{15,79} Indeed, patients presenting with dyspnea have lower survival rates than patients whose initial symptom is related to a pneumothorax.⁸¹ Moreover, in patients with forced expiratory volume in the first second (FEV₁) and DL_{CO} >40% predicted, the rate of functional decline in patients presenting with dyspnea on exertion was greater than in those presenting with pneumothorax.⁸¹

In patients who have had a lung biopsy, a predominance of cystic lesions instead of cellular infiltrates is associated with worse prognosis.^{82,83} The presence of hemosiderin-laden macrophages in the biopsy tissue is also associated with more severe disease.⁸² Patients with more cystic disease tend to have lower FEV₁ and DL_{CO}, lower VO₂ max, greater exercise-induced hypoxemia, and lower survival than patients with predominantly nodular lesions.^{82,83} Findings on CT scans also correlate with lung function tests, gas exchange, and exercise performance,^{62,67-69} especially if the percentage of lung volume affected by cysts and texture changes in the vicinity of cysts are quantified.^{71,72}

Overall, pulmonary function testing is the most practical method of assessing the severity of lung disease in LAM.^{1,45} In patients with near normal flow rates and a reduced diffusion capacity, the severity of lung disease is best graded by a 6-minute walk test or cardiopulmonary exercise testing.^{61,62,70} Estimation of rates of FEV₁ and DL_{CO} declining over time may help in defining whether lung disease is rapidly or slowly progressing.^{45,80} A positive response to bronchodilators, which occurs in 25%–30% of LAM patients, may be a predictor of disease severity and rate of progression.^{83,84}

VEGF-D levels show a correlation with DL_{CO} and CT scan as measures of severity of lung disease.^{58,59} Analysis of data from the Multicenter International Lymphangi-

leiomyomatosis Efficacy and Safety of Sirolimus (MILES) trial showed that serum levels of VEGF-D correlated positively with use of oxygen, bronchodilator response, lower quality of life, decreased ability to perform daily living activities, lower diffusion capacity, and more severe airflow obstruction.⁶⁰

Treatment

Not long ago, LAM was defined as a fatal disease of young women, for which the only treatment option was lung transplantation. The finding that LAM is caused by mutations of the *TSC1* or *TSC2* genes, which encode hamartin and tuberin, two proteins with a major role in control of the mTOR signaling pathway, led to therapies targeting mTOR (Figure 2).^{8-10,36} mTOR is the catalytic subunit of two distinct complexes named mTOR complex 1 (mTORC1) and mTOR complex 2 (mTORC2) (Figure 2).⁸⁵ Sirolimus and everolimus, two mTORC1 inhibitors, have been shown to be effective in stabilizing lung function and reducing the size of chylous effusions, lymphangioliomyomas, and AMLs.⁸⁶⁻⁸⁸ Treatment with sirolimus or everolimus is now recognized as the standard therapy for patients with declining lung function, symptomatic chylous effusions, lymphangioliomyomas, or large AMLs.⁸⁶⁻⁸⁸ However, discontinuation of mTOR inhibitor therapy results in further decline in lung function or return of AMLs to pretherapy tumor size, suggesting that treatment has to be continued.^{86,88} Inhibition of mTOR results in increased autophagy and possibly enhanced LAM cell survival, reducing the beneficial effects of mTOR inhibitors (Figure 2).^{50,89} Inhibition of autophagy with chloroquine in combination with mTOR inhibition is now being examined as a possible treatment for LAM.⁸⁹ Deficiency of tuberin due to *TSC2* mutations results in increased RhoA GTPase activity and increased cell survival,⁹⁰ an effect mediated through mTORC2 signaling (Figure 2).⁹⁰ Because sirolimus and everolimus only inhibit the activity of mTORC1, it has been suggested that therapies targeting RhoA GTPases should be considered (Figure 2).^{90,91} Statins, namely simvastatin, inhibit Rho GTPases and promote apoptosis.^{56,92} In combination with sirolimus, simvastatin was reported to prevent lung destruction in a mouse model of LAM.⁵⁶ A Phase I clinical trial testing the effect of simvastatin combined with either sirolimus or everolimus in patients with LAM is underway.

Other treatments that are under investigation are estrogen receptor blockers and aromatase inhibitors.^{50,57,91} Because of the proposed role of lymphangiogenesis in the pathogenesis of LAM,⁵⁷ and evidence that increased levels of VEGF-D in

the serum of LAM patients correlate with disease severity and clinical course, blockade of VEGFRs or anti-VEGF therapies are also being considered.^{50,57} In an animal model of LAM, combination of VEGFR1, VEGFR2, and VEGFR3 inhibitor and sirolimus decreased tumor volume and increased survival more effectively than sirolimus alone.⁹³

LAM is considered to be a low-grade malignancy and, as in the case of cancer, treatment with drugs targeting signaling pathways considered important in the pathogenesis of the disease may eventually turn out to be effective. However, most ongoing clinical studies are in a preliminary stage, and there is yet no evidence that they will prove to be effective.

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Disclosure

The authors report no conflicts of interest in this work.

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