COMMENT

Progress in the treatment of lymphangioleiomyomatosis: From bench to bedside

Progressos no tratamento da linfangioleiomiomatose: da investigação à clínica

A.M. Taveira-DaSilva*, J. Moss

Cardiovascular and Pulmonary Branch, National Heart, Lung, and Blood Institute, National Institutes of Health, Bethesda, MD, USA

KEYWORDS
Lymphangioleiomyomatosis; Sirolimus therapy; mTOR

PALAVRAS-CHAVE
Linfangioleiomiomatose; Terapia com rapamicina; mTOR

Lymphangioleiomyomatosis is a multisystem disease affecting women, which is characterized by cystic lung destruction, lymphatic abnormalities (e.g., lymphangioleiomyomas), and abdominal tumors which are found primarily in the kidneys, and consist of smooth muscle cells, vascular structures and adipocytes, termed angiomyolipomas.1,2 Lymphangioleiomyomatosis occurs sporadically and in about one-third of women with tuberous sclerosis complex (TSC), an autosomal dominant disorder with variable penetrance.1,2 In either case, LAM is caused by mutations of TSC1 or TSC2, two genes that encode, respectively, hamartin and tuberin, two proteins that regulate cell growth.3,4 Deficiency of these proteins causes activation of mammalian target of rapamycin (mTOR) downstream elements, leading to increased cell proliferation and size (Fig. 1).

Sporadic lymphangioleiomyomatosis is a rare disease occurring in circa 2-5/1,000,000 persons.1,2 Yet, for those afflicted, LAM is devastating. Progressive lung destruction leads to respiratory failure, lung transplantation or death.1,2 Lymphatic involvement may cause chylous pleural effusions, ascites and lymphangioleiomyomas, abdominal tumors that may mimic lymphomas and sarcomas.3 Renal angiomyolipomas may be associated with abdominal hemorrhage, requiring partial or total nephrectomy or embolization of arterial feeding vessels.6

In the past, patients with LAM were advised to undergo bilateral oophorectomy and hysterectomy and/or be treated with anti-estrogen agents such as progesterone or gonadotropin-releasing hormone analogues. No controlled studies showing that these therapies are beneficial to the patients have ever been carried out.7,8

Several studies have demonstrated the important role of the TSC1/TSC2 complex in cell cycle progression and in cell size and proliferation.9,10 Tuberin binds a cyclin-dependent kinase inhibitor, thus stabilizing it, resulting in inhibition of cell cycle progression. Tuberin also has a GTPase activating protein activity, which converts Ras homolog enriched in brain (Rheb)-GTP to Rheb-GDP, resulting in inactivation of Rheb (Fig. 1) and reduction in activity of mTOR, a kinase that controls translation through the phosphorylation of 4E-BP1 and S6K1 (Fig. 1). Protein kinase B (Akt), when activated by growth factors, phosphorylates and inhibits tuberin, resulting in increased cell size and proliferation.9,10

* Corresponding author.
E-mail address: dasilvaa@nhlbi.nih.gov (A.M. Taveira-DaSilva).

0873-2159/$ - see front matter Published by Elsevier España, S.L. doi:10.1016/j.rppneu.2012.02.005
Sirolimus (Rapamycin), in the presence of FKBP12 (FK506-binding protein), inhibits mTOR. Animal models with a functionally null germline mutation of Tsc2 spontaneously develop renal cell carcinomas which decrease in size following administration of sirolimus.\textsuperscript{11,12} In patients with angiomyolipomas, tumor size decreased by half after 1 year of sirolimus therapy, while lung function improved in some patients but not in others.\textsuperscript{13} Other studies have shown that sirolimus therapy leads to resolution of chylous effusions in post-transplant LAM patients,\textsuperscript{14} decreased the size of giant cell astrocytomas and reduced the frequency of seizures in TSC patients.\textsuperscript{15} Results of a multicenter placebo-controlled randomized trial just completed (MILES trial), showed that sirolimus stabilized lung function, and improved symptoms and quality of life in patients with LAM.\textsuperscript{16} There was a highly significant difference in the change in lung function over 1 year of therapy between sirolimus-treated and placebo-treated patients.\textsuperscript{16} In another study, sirolimus therapy improved chylous effusions and/or lymphangioleiomyomas.\textsuperscript{17} By 3–6 months of therapy, effusions start to resolve and complete resolution may be observed, pre-empting the need for repeated thoracentesis, chest tube drainage or pleurodesis.\textsuperscript{17}

These observations provide the rationale for the use of sirolimus in the treatment of selected LAM patients. Sirolimus therapy may be considered for patients that fulfill the inclusion/exclusion criteria for MILES, those with rapidly progressing lung disease, those with chylous effusions that compromise the lung function and require frequent drainage or pleurodesis, and patients with large symptomatic thoraco-abdominal lymphangioleiomyomas. The starting dose of sirolimus is 2 mg/day. Sirolimus levels must be monitored and dosage adjusted to attain serum trough levels between 5 and 15 ng/ml, a range thought to be therapeutic for patients with renal transplant. The occurrence of adverse events associated with sirolimus therapy, such as, oral mucosa ulcers, hypertension, hyperlipidaemia, proteinuria, increased serum creatinine, infections, and sirolimus-related interstitial pneumonitis, mandate close patient monitoring, and may require withholding of the drug or reduction in dose. In addition, given the limited experience with sirolimus in the treatment of LAM, it is not known whether treatment may have to be continued for life, when it should be started, the optimal drug levels, or if sirolimus resistance will eventually develop.

Figure 1  Simplified schematic model of TSC1 and TSC2 pathways. The TSC1/TSC2 complex has roles in cell cycle progression and in cell size and proliferation. This complex stimulates the conversion of active Rheb-GTP to inactive Rheb-GDP, resulting in inactive Rheb. Rheb controls mTOR, a kinase that controls translation through phosphorylation of S6K1. Sirolimus inhibits mTOR. Abbreviations: PI3K: phosphoinositide 3-kinase; Akt: protein kinase B; mTOR: mammalian target of rapamycin; Rheb GAP: GTPase-activating protein for Ras homolog enriched in brain; S6K1: S6 kinase 1 (modified from Ref. 10).

Funding

Research funded by the Intramural Research Program, National Institutes of Health, National Heart, Lung, and Blood Institute.

Conflict of interest

The authors have no conflicts of interest to declare.

References

10. Taveira-Dasilva AM, Steagall WK, Moss J. Therapeutic options for lymphangioleiomyomatosis (LAM): where we are, where we are going. Med Rep. 2009;1 [pii: 93, PMID: 20948684].