



June has been designated Worldwide LAM Awareness Month (WWLAM) by the Worldwide LAM Patient Coalition (WLPC).

 LAM Australia Research Alliance	 LAM Selbsthilfe Deutschland E.V.	 LAM Nederland	 LAM Action
 LAM Austria	 LAM Support Ireland	 New Zealand LAM Trust	 The LAM Foundation
 ALAMBRA: Associação dos Portadores de Linfangioleiomiomatose do Brasil	 Israel LAM Organization	 AELAM: Asociación Española de Linfangioleiomiomatosis	
 LAM China	 LAM Italia	 LAM Academy	
 FLAM: France Lymphangioliomyomatose	 J-LAM	 LAM Türkiye	

Find LAM Support in your country: thelamfoundation.org/wlpc

Lymphangioliomyomatosis (*lim-FAN-je-o-ll-o-MI-o-ma-TO-sis*) is a rare lung disease that usually strikes women during the prime of their lives. Known as LAM for short, this disease is characterized by an abnormal growth of smooth muscle cells, especially in the lungs, lymphatic system and kidneys. Although research is ongoing, there is currently no cure.