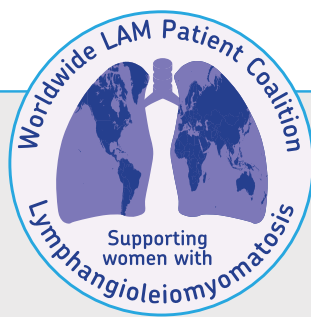


OUR GLOBAL CONNECTION

WORLD WIDE LAM AWARENESS MONTH • JUNE 2020

Lymphangiomyomatosis (lim-FAN-je-o-LI-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. June has been designated Worldwide LAM Awareness Month (WWLAM) by the Worldwide LAM Patient Coalition (WLPC). For more information about LAM, and to gain access to the support network in your country, please contact your local WLPC organization.



LAM AUSTRALIA RESEARCH ALLIANCE
www.lamaustralia.org.au



LAM AUSTRIA
<https://sites.google.com/site/lamselfsthilfe/>



ALAMBRA: ASSOCIAÇÃO DOS PORTADORES DE LINFANGIOLEIOMIOMATOSE DO BRASIL
www.alambrabrasil.org



LAM CHINA
www.lamchina.com.cn



THE EUROPE LAM FEDERATION
www.europelamfederation.org



FLAM: FRANCE LYMPHANGIOLÉIOMYOMATOSE
www.francelam.org



LAM SELBSTHILFE DEUTSCHLAND E.V.
www.lam-info.de



LAM SUPPORT IRELAND
www.lamsupportireland.com



ISRAEL LAM ORGANIZATION
www.lam-israel.org



LAM ITALIA
www.lam-italia.org



J-LAM
www.j-lam.net



LAM NEDERLAND
www.lam-nederland.nl



NEW ZEALAND LAM TRUST
www.lam.org.nz



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