

Asociación Galega de Linfedema, Lipedema, Insuficiencia Venosa Crónica y otras Patologías Vasculares Periféricas (AGL) is a Galician non-profit association in the North-West Autonomous Region of Spain. It was established in 2004, and represents patients with Primary and Secondary Lymphedema, Lipedema and Chronic Venous Insufficiency. The AGL Association is a member of FEDEAL (Spanish Lymphedema Federation), FEGEREC (Galician Rare Diseases Federation) and COGAMI-COCEMFE (Galician Disability Federation). The focus of the Association is to give greater visibility and acceptance of lymphatic disorders to health professionals and society, and that all patients in Sapin have access to good quality health care and compression garments, subsidised by our national health system.