IRVAN syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. IRVAN syndrome. ORPHA:209943

A rare retinal vasculopathy disease characterized by idiopathic retinal vasculitis (IRV), aneurysmal dilations (A) at arteriolar bifurcations, and neuroretinitis (N), which if untreated progresses to peripheral capillary non-perfusion, retinal neovascularization, and macular exudation, leading to severe, bilateral vision loss.