Autosomal dominant rhegmatogenous retinal detachment

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant rhegmatogenous retinal detachment. ORPHA:209867

Autosomal dominant rhegmatogenous retinal detachment is a rare, hereditary, non-syndromic form of vitreoretinopathy characterized by retinal tears due to abnormal vitreous, and commonly present refractive errors. No other signs or symptoms of Stickler syndrome is present.