Retinitis pigmentosa

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Retinitis pigmentosa. ORPHA:791

Retinitis pigmentosa (RP) is an inherited retinal dystrophy leading to progressive loss of the photoreceptors and retinal pigment epithelium and resulting in blindness usually after several decades.