Familial benign flecked retina

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial benign flecked retina. ORPHA:363989

Familial benign flecked retina is a rare retinal dystrophy characterized by diffuse bilateral white-yellow fleck-like lessions extending to the far periphery of the retina but sparing the foveal region, with asymptomatic clinical phenotype and absence of electrophysiologic deficits.