Fundus albipunctatus

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Fundus albipunctatus. ORPHA:227796

Fundus albipunctatus is a rare, genetic retinal dystrophy disorder characterized by the presence of numerous small, round, yellowish-white retinal lesions that are distributed throughout the retina but spare the fovea. Patients present in childhood with non-progressive night blindness with prolonged cone and rod adaptation times. The macula may or may not be involved, which may result in a decrease of central visual acuity with age.