Retinal degeneration-nanophthalmos-glaucoma syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Retinal degeneration-nanophthalmos-glaucoma syndrome. ORPHA:1574

Retinal degeneration-nanophthalmos-glaucoma syndrome is characterized by progressive pigmentary retinal degeneration (with nyctalopia and visual field restriction), cystic macular degeneration and angle closure glaucoma. It has been described in seven members of one family. Patients also have hyperopia and nanophthalmos. The mode of transmission is autosomal recessive.