Congenital microcoria

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital microcoria. ORPHA:566

Congenital microcoria is a rare autosomal dominant ophthalmological disease caused by maldevelopment of the dilator muscle of the pupil that is characterized by small pupils (<2 mm in diameter) from birth, peripheral iris hypopigmentation and transillumination defects leading to errors of refraction (myopia, astigmatism) and sometimes juvenile open angle glaucoma.