Severe early-childhood-onset retinal dystrophy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Severe early-childhood-onset retinal dystrophy. ORPHA:364055

Severe early childhood onset retinal dystrophy (SECORD) is an inherited retinal dystrophy characterized by a severe congenital night blindness, progressive retinal dystrophy and nystagmus. Best corrected visual acuity can reach 0.3 in the first decade of life and can pertain well into the second decade of life. Blindness is often complete by the age of 30 years.