Achromatopsia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Achromatopsia. ORPHA:49382

Achromatopsia (ACHM) is a rare autosomal recessive retinal disorder characterized by color blindness, nystagmus, photophobia, and severely reduced visual acuity due to the absence or impairment of cone function.