Goldmann-Favre syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Goldmann-Favre syndrome. ORPHA:53540

Goldmann-Favre syndrome (GFS) is a vitreoretinal dystrophy characterized by early onset of night blindness, reduced bilateral visual acuity, and typical fundus findings (progressive pigmentary degenerative changes, macular edema, retinoschisis).