Isolated congenital ectropion

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Isolated congenital ectropion. ORPHA:99171

Isolated congenital ectropion is a rare ocular disease characterized by congenital, unilateral or bilateral, lower or upper eyelid malposition with eversion of the margin due to a vertical shortage of skin, leading to exposure of the conjunctiva and sometimes the cornea. Chronic epiphora and exposure keratitis may be observed in severe cases.