Cogan-Reese syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Cogan-Reese syndrome. ORPHA:98980

Cogan-Reese syndrome is a clinical variant of iridocorneal endothelial (ICE) syndrome (see this term) characterized by variable iris atrophy, pigmented and pedunculated nodules on the iris and corneal abonormalities. Secondary glaucoma is also a common complication of the disease.