Congenital hereditary endothelial dystrophy type II

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital hereditary endothelial dystrophy type II. ORPHA:293603

Congenital hereditary endothelial dystrophy II (CHED II) is a rare subtype of posterior corneal dystrophy (see this term) characterized by a diffuse ground-glass appearance of the corneas and marked corneal thickening from birth with nystagmus, and blurred vision.