Congenital hereditary endothelial dystrophy type I

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital hereditary endothelial dystrophy type I. ORPHA:98975

Congenital hereditary endothelial dystrophy I (CHED I) 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 or infancy without nystagmus, with blurred vision.