Cranioectodermal dysplasia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Cranioectodermal dysplasia. ORPHA:1515

Cranioectodermal dysplasia (CED) is a rare developmental disorder characterized by congenital skeletal and ectodermal defects associated with dysmorphic features, nephronophthisis, hepatic fibrosis and ocular anomalies (mainly retinitis pigmentosa).