IVIC syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. IVIC syndrome. ORPHA:2307

IVIC syndrome is a very rare genetic malformation syndrome characterized by upper limb anomalies (radial ray defects, carpal bone fusion), extraocular motor disturbances, and congenital bilateral non-progressive mixed hearing loss.