Distal monosomy 7p

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Distal monosomy 7p. ORPHA:96126

Distal monosomy 7p is a partial autosomal monosomy characterized by developmental delay and intellectual disability, digital anomalies, congenital heart and urogenital anomalies, and specific craniofacial features, commonly including craniosynostosis.