Atelosteogenesis type I

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Atelosteogenesis type I. ORPHA:1190

Atelosteogenesis I is a perinatally lethal skeletal dysplasia characterized by severe short-limbed dwarfism, joint dislocations, club feet along with distinctive facies and radiographic findings.