Grant syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Grant syndrome. ORPHA:2097

Grant syndrome is a rare osteogenesis imperfecta-like disorder, described in two patients to date, characterized clinically by persistent wormian bones, blue sclera, mandibular hypoplasia, shallow glenoid fossa, and campomelia. There have been no further descriptions in the literature since 1986.