Osteogenesis imperfecta type 4

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Osteogenesis imperfecta type 4. ORPHA:216820

Osteogenesis imperfecta type IV is a moderate type of osteogenesis imperfecta (OI; see this term), a genetic disorder characterized by increased bone fragility, low bone mass and susceptibility to bone fractures. Patients with type IV have moderately short stature, mild to moderate scoliosis, grayish or white sclera, and dentinogenesis imperfecta (DI; see this term).