Osteogenesis imperfecta type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Osteogenesis imperfecta type 1. ORPHA:216796

Osteogenesis imperfecta type I is a mild type of osteogenesis imperfecta (OI; see this term), a genetic disorder characterized by increased bone fragility, low bone mass and susceptibility to bone fractures.