Multiple epiphyseal dysplasia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Multiple epiphyseal dysplasia. ORPHA:251

Multiple epiphyseal dysplasias (MED/EDMs) are characterized by epiphyseal anomalies causing joint pain early in life, recurrent osteochondritis and early arthrosis. The EDMs are a heterogeneous group of diseases with variable expression classed as MED/EDMs 1-6.