Thanatophoric dysplasia type 2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Thanatophoric dysplasia type 2. ORPHA:93274

Thanatophoric dysplasia, type 2 (TD2) is a form of TD (see this term) characterized by micromelia, straight long-bones, macrocephaly, brachydactyly, shortened ribs and a clover-leaf skull (kleeblattschaedel).