Terminal osseous dysplasia-pigmentary defects syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Terminal osseous dysplasia-pigmentary defects syndrome. ORPHA:88630

Terminal osseous dysplasia-pigmentary defects syndrome is characterised by malformation of the hands and feet, pigmentary skin lesions on the face and scalp and digital fibromatosis.