Trichorhinophalangeal syndrome type 2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Trichorhinophalangeal syndrome type 2. ORPHA:502

Langer-Giedon syndrome, also known as trichorhinophalangeal syndrome type 2, is a very rare, genetic, multiple congenital anomaly disorder characterized by bone abnormalities, distinctive facial features, multiple exostoses, and intellectual disability.