Brachydactyly type A2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Brachydactyly type A2. ORPHA:93396

Brachydactyly type A2 (BDA2) is a congenital malformation characterized by shortening (hypoplasia or aplasia) of the middle phalanges of the index finger and, sometimes, of the little finger.