Open Peer Review on Qeios

Brachydactyly type B

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Brachydactyly type B. ORPHA:93383

Brachydactyly type B (BDB) is a very rare congenital malformation characterized by hypoplasia or aplasia of the terminal parts of fingers 2 to 5, with complete absence of the fingernails.