# Syndactyly type 5

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Syndactyly type 5. ORPHA:93406*

Syndactyly type 5 (SD5) is a very rare congenital limb malformation characterized by postaxial syndactyly of hands and feet, associated with metacarpal and metatarsal fusion of fourth and fifth digits.