FATCO syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. FATCO syndrome. ORPHA:2492

A rare, genetic, congenital limb malformation syndrome characterized by unilateral or bilateral fibular aplasia/hypoplasia, tibial campomelia, and lower limb oligosyndactyly involving the lateral rays. Upper limb oligosyndactyly and cleft lip/palate may also be associated.