WT limb-blood syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. WT limb-blood syndrome. ORPHA:3466

A rare constitutional aplastic anemia disorder characterized by severe hypo/aplastic anemia or pancytopenia associated with skeletal anomalies (such as radial/ulnar defects and hand/digit abnormalities) and an increased risk of leukemia. There have been no further descriptions in the literature since 1995.