Arthrogryposis-severe scoliosis syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Arthrogryposis-severe scoliosis syndrome. ORPHA:65720

Distal arthrogryposis type 4 is an inherited developmental defect syndrome characterized by multiple congenital contractures of limbs, without primary neurologic and/or muscle disease that affects limb function, and a mild to severe scoliosis. Intelligence is normal.