Open Peer Review on Qeios

Neurogenic arthrogryposis multiplex congenita

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Neurogenic arthrogryposis multiplex congenita. ORPHA:1143

Neurogenic arthrogryposis multiplex congenita is a form of arthrogryposis multiplex congenita characterized by congenital immobility of the limbs with fixation of multiple joints and muscle wasting. This condition is secondary to neurogenic muscular atrophy.