Monomelic amyotrophy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Monomelic amyotrophy. ORPHA:65684

Monomelic amyotrophy (MA) is a rare benign lower motor neuron disorder characterized by muscular weakness and wasting in the distal upper extremities during adolescence followed by a spontaneous halt in progression and a stabilization of symptoms.