Autosomal recessive limb-girdle muscular dystrophy type 2N

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal recessive limb-girdle muscular dystrophy type 2N. ORPHA:206559

Autosomal recessive limb-girdle muscular dystrophy type 2N (LGMD2N) is a form of limb-girdle muscular dystrophy characterized by proximal weakness (manifesting as slowness in running) presenting in infancy, along with calf hypertrophy, mild lordosis, scapular winging and normal intelligence (or mild intellectual disability).