Limb-Girdle Muscular Dystrophy Type 2D

National Cancer Institute

Source

National Cancer Institute. Limb-Girdle Muscular Dystrophy Type 2D. NCI Thesaurus. Code C142081.

An autosomal recessive condition caused by mutation(s) in the SGCA gene, encoding alpha-sarcoglycan. It is characterized by progressive muscular dystrophy, primarily affecting the proximal muscles, resulting in difficulty walking.