Congenital muscular dystrophy, Fukuyama type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital muscular dystrophy, Fukuyama type. ORPHA:272

Fukuyama type muscular dystrophy (FCMD) is a congenital progressive muscular dystrophy characterized by brain malformation (cobblestone lissencephaly), dystrophic changes in skeletal muscle, severe intellectual deficit, epilepsy and motor impairment.