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Duchenne muscular dystrophy

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [Duchenne muscular dystrophy](#). ORPHA:98896

Duchenne muscular dystrophy (DMD) is a neuromuscular disease characterized by rapidly progressive muscle weakness and wasting due to degeneration of skeletal, smooth and cardiac muscle.