Steinert myotonic dystrophy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Steinert myotonic dystrophy. ORPHA:273

Steinert disease, also known as myotonic dystrophy type 1, is a muscle disease characterized by myotonia and by multiorgan damage that combines various degrees of muscle weakness, arrhythmia and/or cardiac conduction disorders, cataract, endocrine damage, sleep disorders and baldness.