Friedreich ataxia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Friedreich ataxia. ORPHA:95

Friedreich ataxia (FRDA) is an inherited neurodegenerative disorder classically characterized by progressive gait and limb ataxia, dysarthria, dysphagia, oculomotor dysfunction, loss of deep tendon reflexes, pyramidal tract signs, scoliosis, and in some, cardiomyopathy, diabetes mellitus, visual loss and defective hearing.