Huntington disease-like 3

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Huntington disease-like 3. ORPHA:157946

Huntington disease-like 3 is a rare Huntington disease-like syndrome characterized by childhood-onset progressive neurologic deterioration with pyramidal and extrapyramidal abnormalities, chorea, dystonia, ataxia, gait instability, spasticity, seizures, mutism, and (on brain MRI) progressive frontal cortical atrophy and bilateral caudate atrophy.