Allan-Herndon-Dudley syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Allan-Herndon-Dudley syndrome. ORPHA:59

Allan-Herndon-Dudley syndrome (AHDS) is an X-linked intellectual disability syndrome with neuromuscular involvement characterized by infantile hypotonia, muscular hypoplasia, spastic paraparesis with dystonic/athetoic movements, and severe cognitive deficiency.