Qazi-Markouizos syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Qazi-Markouizos syndrome. ORPHA:3010

Qazi-Markouizos syndrome is characterised principally by non-progressive central hypotonia, chronic constipation, severe psychomotor retardation, abnormal dermatoglyphics, dysharmonic skeletal maturation and disproportionate muscle fibres. Seizures or an abnormal electroencephalograph were also reported. To date, the syndrome has been reported in three unrelated Puerto Rican boys.