Familial congenital mirror movements

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial congenital mirror movements. ORPHA:238722

A rare, genetic, movement disorder characterized by involuntary movements on one side of the body that mirror intentional movements on the opposite side of the body, which are present in various first-degree members of a family, persist beyond the first decade of life, and have no associated comorbidities.