Spastic tetraplegia-retinitis pigmentosa-intellectual disability syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Spastic tetraplegia-retinitis pigmentosa-intellectual disability syndrome. ORPHA:3011

Spastic tetraplegia-retinitis pigmentosa-intellectual disability syndrome is characterized by nonprogressive spastic paraplegia, retinitis pigmentosa, and intellectual deficit. It has been described in two brothers born to consanguineous parents.