X-linked complicated corpus callosum dysgenesis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. X-linked complicated corpus callosum dysgenesis. ORPHA:1497

X-linked complicated corpus callosum dysgenesis is a historical term used to describe a phenotype now considered to be part of the L1 clinical spectrum (L1 syndrome, see this term). The disorder is characterized by variable spastic paraplegia, mild to moderate intellectual deficit, and dysplasia, hypoplasia or aplasia of the corpus callosum.