Atypical Rett syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Atypical Rett syndrome. ORPHA:3095

Atypical Rett syndrome (atypical RTT) is a neurodevelopmental disorder that is diagnosed when a child presents with a Rett-like syndrome but does not fulfill all the diagnostic criteria for typical Rett syndrome (classic/typical RTT; see this term).