DYRK1A-related intellectual disability syndrome due to 21q22.13q22.2 microdeletion

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. DYRK1A-related intellectual disability syndrome due to 21q22.13q22.2 microdeletion.
ORPHA:268261

A rare, syndromic intellectual disability characterized by global developmental delay including severely delayed or absent speech, moderate to severe intellectual disability, behavioral issues, stereotypic behavior, febrile seizures and epilepsy, abnormal gait, vision defects, and characteristic facial features. Intrauterine growth restriction and feeding difficulties are frequently present.