7q31 microdeletion syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. 7q31 microdeletion syndrome. ORPHA:251061

7q31 microdeletion syndrome is a rare chromosomal anomaly characterized by speech and language disorder, predominantly presenting as an apraxia of speech, sometimes associated with oral motor dyspraxia, dysarthria, receptive and expressive language disorder, and hearing loss. Individuals with larger deletions in this region have also been reported to display intellectual disability and autism.