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Duplication/inversion 15q11

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base.*

[Duplication/inversion 15q11. ORPHA:3306](#)

The duplication/inversion 15q11 or isodicentric 15 chromosome (inv dup(15) or idic(15)) syndrome is a chromosomal disorder with distinctive clinical findings characterized by early central hypotonia, developmental delay and intellectual deficit, epilepsy, and autistic behavior.