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16q24.3 microdeletion syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. [16q24.3 microdeletion syndrome](#). ORPHA:261250

16q24.3 microdeletion syndrome is a recently described syndrome associated with variable developmental delay, facial dysmorphism, seizures and autistic spectrum disorder.