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# 14q11.2 microdeletion syndrome

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. [14q11.2 microdeletion syndrome](#). ORPHA:261120*

14q11.2 microdeletion syndrome is a recently described syndrome characterized by developmental delay, hypotonia and facial dysmorphism.