# 2q37 microdeletion syndrome

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. 2q37 microdeletion syndrome. ORPHA:1001*

Deletion 2q37 or monosomy 2q37 is a chromosomal anomaly involving deletion of chromosome band 2q37 and manifests as three major clinical findings: developmental delay, skeletal malformations and facial dysmorphism.