

[Open Peer Review on Qeios](#)

# 6p22 microdeletion syndrome

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. 6p22 microdeletion syndrome. ORPHA:251046*

6p22 microdeletion syndrome is a newly described syndrome associated with a variable clinical phenotype including developmental delay, facial dysmorphism, short neck and diverse malformations.