

Open Peer Review on Qeios

6q25 microdeletion syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>6q25</u> <u>microdeletion syndrome</u>. ORPHA:251056

6q25 microdeletion syndrome is a recently described syndrome characterized by developmental delay, facial dysmorphism and hearing loss.

Qeios ID: TQ3GTJ · https://doi.org/10.32388/TQ3GTJ