

Open Peer Review on Qeios

6q terminal deletion syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>6q terminal</u> <u>deletion syndrome</u>. ORPHA:75857

6q terminal deletion syndrome is marked by a characteristic facial dysmorphism, short neck and psychomotor retardation, generally associated with a range of non-specific malformations.

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