

Open Peer Review on Qeios

Oliver syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Oliver</u> <u>syndrome</u>. ORPHA:2920

Oliver syndrome is a very rare syndrome characterized by intellectual deficit, postaxial polydactyly, and epilepsy.

Qeios ID: S759Z7 · https://doi.org/10.32388/S759Z7