

Open Peer Review on Qeios

Kleefstra syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Kleefstra syndrome</u>. ORPHA:261494

Kleefstra syndrome (KS) is a genetic disorder characterized by intellectual disability, childhood hypotonia, severe expressive speech delay and a distinctive facial appearance with a spectrum of additional clinical features.

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