

Open Peer Review on Qeios

Robinow syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Robinow</u> <u>syndrome</u>. ORPHA:97360

Robinow syndrome (RS) is a rare genetic syndrome characterized by limb shortening and abnormalities of the head, face and external genitalia.

Qeios ID: LL38HD · https://doi.org/10.32388/LL38HD