

Open Peer Review on Qeios

Noonan syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Noonan</u> <u>syndrome</u>. ORPHA:648

Noonan Syndrome (NS) is characterised by short stature, typical facial dysmorphism and congenital heart defects.

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