

Open Peer Review on Qeios

Congenital nephrotic syndrome, Finnish type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Congenital</u> nephrotic syndrome, Finnish type. ORPHA:839

Congenital nephrotic syndrome, Finnish type is characterised by protein loss beginning during foetal life.

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