

Open Peer Review on Qeios

Scott syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Scott</u> <u>syndrome</u>. ORPHA:806

Scott syndrome is an extremely rare congenital hemorrhagic disorder characterized by hemorrhagic episodes due to impaired platelet coagulant activity.

Qeios ID: 3O4LOB · https://doi.org/10.32388/3O4LOB