

Open Peer Review on Qeios

Congenital factor X deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Congenital</u> factor X deficiency. ORPHA:328

Congenital factor X deficiency is an inherited bleeding disorder with a decreased antigen and/or activity of factor X (FX) and characterized by mild to severe bleeding symptoms.

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