

Open Peer Review on Qeios

## Congenital factor II deficiency

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Congenital</u> factor II deficiency. ORPHA:325

Congenital factor II deficiency is an inherited bleeding disorder due to reduced activity of factor II (FII, prothrombin) and characterized by mucocutaneous bleeding symptoms.

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