

Open Peer Review on Qeios

Wilson disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Wilson</u> <u>disease</u>. ORPHA:905

Wilson disease is a very rare inherited multisystemic disease presenting non-specific neurological, hepatic, psychiatric or osseo-muscular manifestations due to excessive copper deposition in the body.

Qeios ID: 5DQV40 · https://doi.org/10.32388/5DQV40