

[Open Peer Review on Qeios](#)

Gitelman syndrome

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [Gitelman syndrome](#). ORPHA:358

A rare genetic renal salt-losing tubular disease, that is characterized by hypokalemic metabolic alkalosis with hypomagnesemia and hypocalciuria.