

Open Peer Review on Qeios

Familial Glucocorticoid Deficiency Type 1

National Cancer Institute

Source

National Cancer Institute. <u>Familial Glucocorticoid Deficiency Type 1</u>. NCI Thesaurus. Code C123727.

Familial glucocorticoid deficiency caused by mutation(s) in the MC2R gene encoding the adrenocorticotropin (ACTH) receptor, also known as the melanocortin-2 receptor.

Qeios ID: 1FRP6V · https://doi.org/10.32388/1FRP6V