

Open Peer Review on Qeios

Congenital Disorder of Glycosylation Type la

National Cancer Institute

Source

National Cancer Institute. <u>Congenital Disorder of Glycosylation Type Ia</u>. NCI Thesaurus. Code C126868.

A congenital disorder of glycosylation sub-type caused by mutation(s) in the PMM2 gene, encoding phosphomannomutase 2.

Qeios ID: YTF31V · https://doi.org/10.32388/YTF31V