

Open Peer Review on Qeios

Tyrosine Hydroxylase Deficiency

National Cancer Institute

Source

National Cancer Institute. <u>Tyrosine Hydroxylase Deficiency</u>. NCI Thesaurus. Code C157158

An autosomal recessive condition caused by mutation(s) in the TH gene, encoding tyrosine 3-monooxygenase. It is characterized by onset in infancy of dopa-responsive dystonia.

Qeios ID: YTY95U · https://doi.org/10.32388/YTY95U