

Open Peer Review on Qeios

Hexokinase Deficiency

National Cancer Institute

Source

National Cancer Institute. <u>Hexokinase Deficiency</u>. NCI Thesaurus. Code C131645.

A rare, autosomal recessive, inherited disorder caused by mutation of the HK1 gene. It is characterized by the early-onset of severe, non-spherocytic hemolytic anemia.

Qeios ID: FMQMN5 · https://doi.org/10.32388/FMQMN5