

Open Peer Review on Qeios

Hereditary Coproporphyria

National Cancer Institute

Source

National Cancer Institute. <u>Hereditary Coproporphyria</u>. NCI Thesaurus. Code C84759.

An autosomal dominant inherited disorder of porphyrin metabolism caused by deficiency of the enzyme coproporphyrinogen oxidase. It results in neurologic damage and can include abdominal pain, constipation and psychiatric manifestations.

Qeios ID: VUUI83 · https://doi.org/10.32388/VUUI83