Open Peer Review on Qeios

Gyrate Atrophy

National Cancer Institute

Source

National Cancer Institute. <u>Gyrate Atrophy</u>. NCI Thesaurus. Code C84744.

A rare autosomal recessive inherited disorder caused by mutations in the OAT gene. It is characterized by progressive atrophy of the retina and choroid, leading to loss of vision and blindness.