Open Peer Review on Qeios

Stargardt Disease

National Cancer Institute

Source

National Cancer Institute. <u>Stargardt Disease</u>. NCI Thesaurus. Code C85078.

An autosomal recessive and rarely autosomal dominant inherited disorder caused by mutations in the ABCA4 or ELOVL4 genes respectively. It is characterized by macular degeneration that begins in late childhood resulting in progressive loss of vision.