

Open Peer Review on Qeios

Rieger Syndrome Type 1

National Cancer Institute

Source

National Cancer Institute. <u>Rieger Syndrome Type 1</u>. NCI Thesaurus. Code C75015.

A rare autosomal dominant syndrome linked to mutations in the PITX2 gene. It is characterized by abnormalities in the anterior chamber of the eye and underdevelopment of the teeth.

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