

Open Peer Review on Qeios

Oguchi disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Oguchi</u> <u>disease</u>. ORPHA:75382

Oguchi disease is an autosomal recessive retinal disorder characterized by congenital stationary night blindness (see this term) and the Mizuo-Nakamura phenomenon.

Qeios ID: N1YD4T · https://doi.org/10.32388/N1YD4T