

Open Peer Review on Qeios

## Autosomal dominant keratitis

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Autosomal dominant keratitis</u>. ORPHA:2334

Hereditary keratitis is characterised by opacification and vascularisation of the cornea, often associated with macula hypoplasia.

Qeios ID: GVRU52 · https://doi.org/10.32388/GVRU52