X-linked cone dysfunction syndrome with myopia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. X-linked cone dysfunction syndrome with myopia. ORPHA:90001

X-linked cone dysfunction syndrome with myopia is characterised by moderate to high myopia associated with astigmatism and deuteranopia. Less than 10 families have been described so far. Transmission is X-linked recessive and the locus has been mapped to Xq28.