Open Peer Review on Qeios

Knobloch syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Knobloch</u> <u>syndrome</u>. ORPHA:1571

Knobloch syndrome (KS) is defined by vitreoretinal and macular degeneration, and occipital encephalocele.