Open Peer Review on Qeios

Hypergonadotropic hypogonadism-cataract syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hypergonadotropic hypogonadism-cataract syndrome. ORPHA:2410

This syndrome is characterized by the association of hypergonadotropic hypogonadism and cataracts with onset during adolescence. It has been described in three brothers from a consanguineous family.