Open Peer Review on Qeios

Isochromosomy Yp

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Isochromosomy Yp. ORPHA:98797

Isochromosomy Yp is a rare gonosome anomaly characterized by various clinical presentations including normal healthy fertile males, male phenotype with infertility, and males with ambiguous genitalia or incomplete masculinization.