

Open Peer Review on Qeios

48,XXXY syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>48,XXXY</u> <u>syndrome</u>. ORPHA:96263

The 48,XXXY syndrome represents a chromosomal anomaly of the aneuploidic type characterized by the presence of two extra X chromosomes in males.

Qeios ID: XFB1CM · https://doi.org/10.32388/XFB1CM