

Open Peer Review on Qeios

Trisomy X

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Trisomy X.</u>
ORPHA:3375

Trisomy X is a sex chromosome anomaly with a variable phenotype caused by the presence of an extra X chromosome in females (47,XXX instead of 46,XX).

Qeios ID: MD0HJP · https://doi.org/10.32388/MD0HJP