

Open Peer Review on Qeios

49,XXXXY syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>49,XXXXY</u> <u>syndrome</u>. ORPHA:96264

The 49,XXXXY syndrome represents a chromosomal anomaly of the aneuploidic type characterized by the presence of three extra X chromosomes in males.

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