Open Peer Review on Qeios

Jacobsen syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Jacobsen</u> <u>syndrome</u>. ORPHA:2308

Jacobsen syndrome is a multiple congenital anomaly/mental retardation (MCA/MR) contiguous gene syndrome caused by partial deletion of the long arm of chromosome 11.