

Open Peer Review on Qeios

Chordoma

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Chordoma</u>. ORPHA:178

Chordomas are rare malignant tumors arising from embryonic remnants of the notochord in axial skeleton.

Qeios ID: CQK8U8 · https://doi.org/10.32388/CQK8U8