

Open Peer Review on Qeios

Gorlin-Chaudhry-Moss syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Gorlin-Chaudhry-Moss syndrome</u>. ORPHA:2095

Gorlin-Chaudhry-Moss (GCM) syndrome is a multiple congenital anomaly syndrome characterized by craniofacial dysostosis, facial dysmorphism, conductive hearing loss, generalized hypertrichosis, and extremity, ocular and dental anomalies.

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