Open Peer Review on Qeios

Baller-Gerold syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Baller-Gerold syndrome. ORPHA:1225

Baller-Gerold syndrome is characterized by the association of coronal craniosynostosis with radial ray anomalies (oligodactyly, aplasia or hypoplasia of the thumb, aplasia or hypoplasia of the radius).