Open Peer Review on Qeios

Isolated Pierre Robin syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Isolated Pierre Robin syndrome. ORPHA:718

Pierre-Robin syndrome (or Pierre-Robin sequence) is characterised by triad of orofacial morphological anomalies consisting of retrognathism, glossoptosis and a posterior median velopalatal cleft.