Muenke syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Muenke syndrome. ORPHA:53271

Muenke syndrome is a syndromic craniosynostosis with significant phenotypic variability, usually characterized by coronal synostosis, midfacial retrusion, strabismus, hearing loss and developmental delay.