Familial scaphocephaly syndrome, McGillivray type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial scaphocephaly syndrome, McGillivray type. ORPHA:168624

Familial scaphocephaly syndrome, McGillivray type is a rare newly described craniosynostosis (see this term) syndrome characterized by scaphocephaly, macrocephaly, severe maxillary retrusion, and mild intellectual disability.