Saethre-Chotzen syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Saethre-Chotzen syndrome. ORPHA:794

Saethre-Chotzen syndrome (SCS) is an inherited craniosynostosis syndrome characterized by unilateral or bilateral coronal synostosis, facial asymmetry, ptosis, strabismus and small ears with prominent crus, among other less common manifestations.