Frontofacionasal dysplasia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Frontofacionasal dysplasia. ORPHA:1791

Fronto-facio-nasal dysostosis is characterized by multiple craniofacial anomalies (brachycephaly, blepharophimosis, ptosis, S-shaped palpebral fissures, coloboma, cleft lip and palate, deformed nostrils, encephalocele, hypertelorism, midface hypoplasia, malformed eyes, and absent inner eyelashes).