Focal facial dermal dysplasia type I

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Focal facial dermal dysplasia type I. ORPHA:79133

Focal facial dermal dysplasia type I (FFDD1), also known as Brauer syndrome, is a focal facial dysplasia (FFDD; see this term) characterized by congenital bitemporal cutis aplasia.