Holoprosencephaly

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Holoprosencephaly. ORPHA:2162

Holoprosencephaly (HPE) is a complex brain malformation resulting from incomplete cleavage of the prosencephalon, occurring between the 18th and 28th day of gestation, and affecting both the forebrain and face, which results in neurological manifestations and facial anomalies of variable severity.