Deafness-hypogonadism syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Deafness-hypogonadism syndrome. ORPHA:90646

This syndrome is characterized by the association of congenital mixed hearing loss with perilymphatic gusher (Gusher syndrome or DFN3; see this term), hypogonadism and abnormal behavior.