Hyposmia-nasal and ocular hypoplasia-hypogonadotropic hypogonadism syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hyposmia-nasal and ocular hypoplasia-hypogonadotropic hypogonadism syndrome. ORPHA:2250

This syndrome is characterized by the association of severe nasal hypoplasia, hypoplasia of the eyes, hyposmia, hypogeusia and hypogonadotropic hypogonadism.