Waardenburg syndrome type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Waardenburg syndrome type 1. ORPHA:894

Waardenburg syndrome type 1 (WS1) is a subtype of Waardenburg syndrome (WS; see this term), disorder characterized by congenital deafness, minor defects in structures arising from neural crest resulting in pigmentation anomalies of eyes, hair, and skin, in combination with dystopia canthorum.