# Waardenburg syndrome type 2

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Waardenburg syndrome type 2. ORPHA:895*

Waardenburg syndrome type 2 (WS2) is an autosomal dominant subtype of Waardenburg syndrome (WS; see this term), characterized by varying degrees of deafness and pigmentation anomalies of eyes, hair and skin, but without dystopia canthorum.