Van der Woude syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Van der Woude syndrome. ORPHA:888

Van der Woude syndrome (VWS) is a rare congenital genetic dysmorphic syndrome characterized by paramedian lower-lip fistulae, cleft lip with or without cleft palate, or isolated cleft palate.