Orofaciodigital syndrome type 5

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Orofaciodigital syndrome type 5. ORPHA:2919

Oral-facial-digital syndrome, type 5 is characterized by median cleft of the upper lip, postaxial polydactyly of hands and feet, and oral manifestations (duplicated frenulum).