Orofaciodigital syndrome type 4

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Orofaciodigital syndrome type 4. ORPHA:2753

Oral-facial-digital syndrome, type 4 is characterized by lingual hamartoma, postaxial polysyndactyly of hands and feet, and mesomelic shortening of the legs with supinate equinovarus feet.