Odontotrichomelic syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Odontotrichomelic syndrome. ORPHA:2723

Odontotrichomelic syndrome is characterised by malformations of all four extremities, hypoplastic nails, ear anomalies, hypotrichosis, abnormal dentition, hyperhidrosis and nasolacrimal duct obstruction. So far, it has been described in less than 10 patients. Transmission is autosomal recessive.