Open Peer Review on Qeios

Keratosis palmaris et plantarisclinodactyly syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Keratosis</u> <u>palmaris et plantaris-clinodactyly syndrome</u>. ORPHA:86919

Keratosis palmaris et plantaris-clinodactyly syndrome is characterised by the association of palmoplantar keratosis with clinodactyly of the fifth finger. Less than 20 cases have been described in the literature so far, and the majority of reported patients were of Mexican origin. Transmission is autosomal dominant.