Punctate palmoplantar keratoderma type 2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Punctate palmoplantar keratoderma type 2. ORPHA:79502

Punctate palmoplantar keratoderma type 2 is a type of isolated, punctate, hereditary palmoplantar keratoderma characterized by multiple, asymptomatic, 1 to 2 mm-long, firm, hyperkeratotic projections ("spiny keratosis") on the palms, soles and digits (typically confined to their volar and/or lateral aspects). Histopathologically, compact columnar parakeratosis over hypo- or agranular epidermis is observed.