Hypopigmentation-punctate palmoplantar keratoderma syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hypopigmentation-punctate palmoplantar keratoderma syndrome. ORPHA:324561

A rare, genetic, epidermal disease characterized by punctate keratoderma on palms and soles associated with irregularly shaped hypopigmented macules (typically localized on the extremities). Ectopic calcification (e.g. early-onset calcific tendinopathy, calcinosis cutis) and pachyonychia may be occasionally associated.