Autosomal dominant palmoplantar keratoderma and congenital alopecia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant palmoplantar keratoderma and congenital alopecia. ORPHA:1010

Autosomal dominant palmoplantar keratoderma with congenital alopecia (PPK-CA) is a rare genetic skin disorder characterized by absence of scalp and body hair and palmoplantar keratoderma, without other hand complications.