Autosomal recessive palmoplantar keratoderma and congenital alopecia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal recessive palmoplantar keratoderma and congenital alopecia. ORPHA:1366

Autosomal recessive palmoplantar hyperkeratosis and congenital alopecia (PPK-CA) is a rare genetic skin disorder characterized by congenital alopecia and palmoplantar hyperkeratosis. It is usually associated with cataracts, progressive sclerodactyly and pseudo-ainhum.