Epidermolysis bullosa simplex, autosomal recessive K14

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Epidermolysis bullosa simplex, autosomal recessive K14. ORPHA:89838

Epidermolysis bullosa simplex, autosomal recessive K14 (EBS-AR KRT14) is a basal subtype of epidermolysis bullosa simplex (EBS) characterized by generalized or, less frequently, localized acral blistering.