Acral dystrophic epidermolysis bullosa

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Acral dystrophic epidermolysis bullosa. ORPHA:158673

Acral dystrophic epidermolysis bullosa is a very rare subtype of dystrophic epidermolysis bullosa (DEB, see this term) characterized by blistering confined primarily to the hands and feet.