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Acral dystrophic epidermolysis bullosa

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Acral</u> <u>dystrophic epidermolysis bullosa</u>. 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.

Qeios ID: 2DUFVB · https://doi.org/10.32388/2DUFVB