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Epidermolysis bullosa simplex due to plakophilin deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Epidermolysis bullosa simplex due to plakophilin deficiency. ORPHA:158668

Epidermolysis bullosa simplex due to plakophilin deficiency (EBS-PD) is a suprabasal subtype of epidermolysis bullosa simplex (EBS, see this term) characterized by generalized superficial erosions and less commonly blistering.

Qeios ID: 4BX58I · https://doi.org/10.32388/4BX58I