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Severe generalized recessive dystrophic epidermolysis bullosa

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base. Severe generalized recessive dystrophic epidermolysis bullosa. ORPHA:79408*

Severe generalized recessive dystrophic epidermolysis bullosa (RDEB-sev gen) is the most severe subtype of dystrophic epidermolysis bullosa (DEB, see this term), formerly known as the Hallopeau-Siemens type, and is characterized by generalized cutaneous and mucosal blistering and scarring associated with severe deformities and major extracutaneous involvement.