Junctional epidermolysis bullosa

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Junctional epidermolysis bullosa. ORPHA:305

Junctional epidermolysis bullosa (JEB) is a form of inherited epidermolysis bullosa (see this term) characterized by involvement of the skin and mucous membranes, and is defined by the formation of blistering lesions between the epidermis and the dermis at the lamina lucida level of the cutaneous basement membrane zone and by healing of lesions with atrophy and/or exuberant granulation tissue formation.