Epidermolysis bullosa simplex

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Epidermolysis bullosa simplex. ORPHA:304

Epidermolysis bullosa simplex (EBS) is a group of hereditary epidermolysis bullosa (HEB) disorders characterized by skin fragility resulting in intraepidermal blisters and erosions that occur either spontaneously or after physical trauma.