Kindler syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Kindler syndrome. ORPHA:2908

Kindler syndrome (KS) is the fourth major type of epidermolysis bullosa (EB), besides simplex, junctional and dystrophic forms, and is characterized by skin fragility and blistering at birth followed by development of photosensitivity and progressive poikilodermatous skin changes.