DISEASE:
Kindler epidermolysis bullosa

NAME:	Kindler epidermolysis bullosa
DESCRIPTION:	A rare inherited epidermolysis bullosa (EB) characterized by skin fragility and blistering at birth followed by development of photosensitivity and progressive poikilodermatous skin changes.
ORPHACODE:	2908
SYNONYMS:	Congenital bullous poikiloderma Kindler syndrome Poikiloderma of Kindler
XREF(S):	Orphanet MeSH ICD-10
ANALYTE(S):	FERMT1
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