DISEASE:
Epidermolysis bullosa simplex with mottled pigmentation

NAME:	Epidermolysis bullosa simplex with mottled pigmentation
DESCRIPTION:	A rare, inherited, epidermolysis bullosa simplex characterized by neonatal or infantile onset of generalized blistering with mottled or reticulate brown pigmentation developing later. Blistering is often accompanied by mild nail dystrophy and focal palmoplantar keratoderma, and rarely by milia and mostly affects the limbs and trunk.
ORPHACODE:	79397
SYNONYMS:	EBS with mottled pigmentation EBS-MP
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	KRT14 KRT5
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