## **Opis choroby \***

## Definicja

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.

Q81.0

Dane

<b>Klasyfikacja</b> Choroba	Synonimy EBS with mottled pigmentation EBS-MP EBS with mottled pigmentation EBS-MP	
<b>Kod ORPHA</b> 79397	<b>Kod OMIM</b> 131960	<b>Kod ICD10</b>

Kod ICD11 EC30

## <u>\*Źródło</u>

orphanet