

## Opis choroby \*

### Definicja

A rare genetic disease characterized by thickening of the skin on palms and soles restricted to areas of weight bearing and/or friction (focal, non-epidermolytic palmoplantar keratoderma) and oral and esophageal leukokeratosis, associated with a very high lifetime risk of developing squamous cell carcinoma of the esophagus. The skin lesions appear in childhood and can be complicated by fissuring and infection.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Bennion-Patterson syndrome  
Hiperkeratoza dłoni i stóp - rak przełyku  
Keratosis palmoplantaris-esophageal carcinoma syndrome  
Zespół Benniona i Pattersona  
Zespół hiperkeratozy dłoni i stóp oraz nowotworu przełyku  
Zespół Howella i Evansa  
Howell-Evans syndrome  
Keratosis palmoplantaris-esophageal carcinoma syndrome  
Palmoplantar hyperkeratosis-esophageal carcinoma syndrome  
Tylosis-oesophageal carcinoma syndrome

#### Kod ORPHA

2198

#### Kod OMIM

148500

#### Kod ICD10

Q82.8

#### Kod ICD11

EC20.31

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#### [\\*Źródło](#)

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