

## Opis choroby \*

### Definicja

A rare, genetic punctate palmoplantar keratoderma disease characterized by discrete, focal, punctate keratoderma on the palms and soles and/or slowly progressive spastic paralysis, predominantly affecting the lower limbs. Lesional histology reveals pronounced orthokeratosis, acanthosis, papillomatosis, and regular undulation to the surface keratin. There have been no further descriptions in the literature since 1983.

### Dane

Klasyfikacja	Synonimy
Choroba	Palmoplantar hyperkeratosis-spastic paralysis syndrome Hiperkeratoza dloni i stop-zespól porażenia spastycznego Zespół Powella, Venencie i Gordona Powell-Venencie-Gordon syndrome

Kod ORPHA	Kod OMIM	Kod ICD10
2201	148360	Q82.8

### Kod ICD11

-

---

### \*Źródło

orphanet