Opis choroby *

Definicja

Hereditary palmoplantar keratoderma, Gamborg-Nielsen type is characterised by the presence of diffuse palmoplantar keratoderma without associated symptoms. The syndrome has been described in multiple families from the northernmost county of Sweden (Norrbotten). The palmoplantar keratoderma found in the Gamborg-Nielsen type disease is milder than that found in Mal de Meleda but more severe than that found in Thost-Unna palmoplantar keratoderma (see these terms). Transmission is autosomal recessive.

Dane

Klasyfikacja
Choroba

Synonimy
Hereditary palmoplantar hyperkeratosis, Gamborg-Nielsen type
Dziedziczna hiperkeratoza dłoniowo-podeszwowa typu Gamborga i Nielsena
PPK, typ Gamborga i Nielsena
PPK, Gamborg-Nielsen type

Kod ORPHA
86923

Kod OMIM
244850

Kod ICD10
Q82.8

Kod ICD11
EC20.3

---

*Źródło

orphanet