Opis choroby *

Definicja

A rare genetic skin disease characterized by congenital generalized anhidrosis resulting in severe heat intolerance, due to functionally impaired eccrine sweat production. Skin biopsy reveals normal morphology and number of sweat glands. Dental, hair, nail, or other skin or extracutaneous anomalies are absent.

Dane

Klasyfikacja
Choroba

Kod ORPHA
468666

Kod OMIM
106190

Kod ICD10
L98.8

Kod ICD11
-

---

*Źródło

orphanet