## Opis choroby *

Definicja

A rare X-linked syndromic intellectual disability characterized by intellectual deficit, choroideremia, horizontal nystagmus, severe myopia, acrokeratosis verruciformis-like skin abnormality, anhidrosis, and scapular winging. There have been no further descriptions in the literature since 1959.

Dane

**Klasyfikacja**
Zespół wad wrodzonych

**Kod ORPHA**
3417

**Kod OMIM**
314500

**Kod ICD10**
Q87.8

**Kod ICD11**
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*Źródło

orphanet