## Opis choroby *

Definicja

A rare, X-linked syndromic intellectual disability disorder characterized by severe intellectual disability, psychomotor developmental delay, generalized seizures, and psoriasis. Mild craniofacial dysmorphism, such as hypertelorism, broad nasal bridge, anteverted nares, macrostomia, highly arched palate and large ears, is also associated. There have been no further descriptions in the literature since 1988.

Dane

**Klasyfikacja**
Choroba

Synonimy
Tranebjaerg-Svejgaard syndrome
Zespół Tranebjaerga i Svejgaarda

**Kod ORPHA**
3052

**Kod OMIM**
309480

**Kod ICD10**
Q87.8

**Kod ICD11**
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*Źródło

orphanet