## Opis choroby *

Definicja

Type 1 lissencephaly due to doublecortin (DCX) gene mutations is a semi-dominant X-linked disease characterised by intellectual deficiency and seizures that are more severe in male patients.

Dane

**Klasyfikacja**
Choroba

Synonimy
X-linked lissencephaly type 1
Lizencefalia typu 1 sprzężona z chromosomem X

**Kod ORPHA**
2148

**Kod OMIM**
300067

**Kod ICD10**
Q04.3

**Kod ICD11**
LD20.1

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*Źródło

orphanet