

## Opis choroby \*

### Definicja

A rare, genetic, chromosomal anomaly syndrome resulting from partial duplication of the long arm of chromosome 2 characterized by congenital pendular nystagmus associated with bilateral cutaneous syndactyly between the third and fourth fingers.

### Dane

#### Klasyfikacja

Zespół wad wrodzonych Syndactyly-nystagmus syndrome due to

#### Synonimy

dup(2)(q31.1)

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Trisomia 2q31.1

Syndactyly-nystagmus syndrome due to trisomy  
2q31.1

#### Kod ORPHA

294026

#### Kod OMIM

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#### Kod ICD10

Q92.3

#### Kod ICD11

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#### \*Źródło

orphanet