

## Opis choroby \*

### Definicja

A rare syndromic trigonocephaly characterized by marked malformations of the head and face (essentially acrocephaly), broad depressed nasal bridge, narrow maxillae, abnormalities of the hands and feet (polydactyly, brachydactyly, syndactyly, clinodactyly, camptodactyly, ulnar deviation), obesity and congenital heart disease. This disease is considered a variant of Carpenter syndrome without intellectual disability. There have been no further descriptions in the literature since 1992.

### Dane

### Klasyfikacja

Zespół wad wrodzonych

#### Kod ORPHA

3210

#### Kod OMIM

272350

#### Kod ICD10

Q82.0

#### Kod ICD11

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### \*Źródło

orphanet