

## Opis choroby \*

### Definicja

Severe achondroplasia-developmental delay-acanthosis nigricans syndrome is characterised by the association of severe achondroplasia with developmental delay and acanthosis nigricans. It has been described in four unrelated individuals. Structural central nervous system anomalies, seizures and hearing loss were also reported, together with bowing of the clavicle, femur, tibia and fibula in some cases. The syndrome is caused by a Lys650Met substitution in the kinase domain of fibroblast growth factor receptor 3 (encoded by the *FGFR3* gene; 4p16.3).

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SADDAN

SADDAN

#### Kod ORPHA

85165

#### Kod OMIM

616482

#### Kod ICD10

Q77.4

#### Kod ICD11

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#### \*Źródło

orphanet