

Opis choroby *

Definicja

A rare genetic bone development disorder characterized by pre- and postnatal growth retardation, skeletal anomalies such as arachnodactyly and bilateral talipes equinovarus, joint contractures with camptodactyly, dysmorphic facial features (including midface hypoplasia or micrognathia), and abnormalities of the anterior segment of the eye. Skeletal imaging may show diffuse osteopenia and multiple fractures. The syndrome is lethal within the first year of life.

Dane

Klasyfikacja

Zespół wad wrodzonych Al Gazali-Al Talabani syndrome

Zespół Al Gazali i Al Talabani

Zespół Al Gazali i Lytle

Al Gazali-Lytle syndrome

Kod ORPHA

2725

Kod OMIM

609465

Kod ICD10

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Kod ICD11

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*[Źródło](#)

orphanet