

## 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

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Al Gazali-Al Talabani syndrome Zespół Al Gazali i Al Talabani Zespół Al Gazali i Lytle Al Gazali-Lytle syndrome

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
2725	609465	-

**Kod ICD11**  
-

---

### \*Źródło

orphanet