

# Amioplazja wrodzona

**Kod Orpha: 488586 Kod OMIM:**

## Opis choroby \*

### Definicja

A rare sporadic arthrogryposis syndrome characterized by multiple congenital contractures presenting in a very specific pattern. It is typically symmetric, involving all four limbs, with internally rotated shoulders, fully extended and fixed elbows, the wrists fixed in flexion, partially flexed fingers, hips fixed in flexion or extension, adducted or abducted, and sometimes dislocated. The knees may be fixed in extension or flexion, and the feet are usually in severe equinovarus position. The jaw and trunk are relatively spared. Normal limb muscle tissue is replaced by fatty, fibrous tissue.

### Dane

#### Klasyfikacja

Zespół wad wrodzonych

#### Synonimy

Amyoplasia congenita  
Wrodzona amioplazja

#### Kod ORPHA

488586

#### Kod OMIM

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#### Kod ICD10

Q68.8

#### Kod ICD11

LD26.41

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[\\*Źródło](#)

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## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

Dostępna na stronie [www.orphanet.pl](http://www.orphanet.pl)