

## Opis choroby \*

### Definicja

A relatively severe form of brachyolmia, a group of rare genetic skeletal disorders, characterized by short-trunked short stature, platyspondyly and kyphoscoliosis. Degenerative joint disease (osteoarthropathy) in the spine, large joints and interphalangeal joints becomes manifest in adulthood.

### Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych	Brachyolmia type 3 Brachyolmia typu 3

Kod ORPHA	Kod OMIM	Kod ICD10
93304	113500	Q76.3

**Kod ICD11**  
LD24.5Y

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### \*Źródło

orphanet