

## Opis choroby \*

### Definicja

A rare, superficial fibromatosis characterized by non-malignant, locally invading, fibrosing tumour of differentiated fibroblasts, slowly growing subcutaneously, occurring predominantly distally on the extremities, especially the hands and feet. Histologic examination shows a multinodular pattern with large areas of calcification and fibrosis, and the presence of elongated spindle cells with hyperchromatic plump vesicular nuclei interspersed within fine bands of collagen.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Juvenile aponeurotic fibromatosis

Guz Keasby'ego

Młodzieńcza włókniakowatość rozciągną

Keasby tumor

#### Kod ORPHA

199260

#### Kod OMIM

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#### Kod ICD10

M72.8

#### Kod ICD11

EE61

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#### \*Źródło

orphanet