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 rozcięgna
Keasby tumor

Kod ORPHA
199260

Kod OMIM
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Kod ICD10
M72.8

Kod ICD11
EE61

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*Źródło

orphanet