

## Opis choroby \*

### Definicja

A rare soft tissue sarcoma characterized by a high-grade lesion occurring almost exclusively in adults, composed of bizarre polygonal, round, and spindle cells with evidence of skeletal muscle differentiation. Patients usually present with a rapidly growing, painful mass located in the deep soft tissues of the extremities, but also other anatomic regions. Prognosis is generally poor.

### Dane

### Klasyfikacja

Podtyp kliniczny

#### Kod ORPHA

293199

#### Kod OMIM

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#### Kod ICD10

C49.9

#### Kod ICD11

XH4VB5

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### \*Źródło

orphanet