Opis choroby *

Definicja

A rare, aggressive, malignant, neoplastic disease characterized by a usually ill-defined, solid, multilobulated mass, frequently having necrosis, located on any site of the body (except the central nervous system), composed of small, round, poorly differentiated cells, with or without Homer-Wright rosettes, showing varying degrees of neuroectodermal differentiation. Manifestations are variable depending on location, with osteolytic destruction being common when arising from bone.

Dane

Klasyfikacja
Choroba

Synonimy
PPNET
Obwodowy PNET
PPNET
Peripheral PNET
Peripheral neuroepithelioma

Kod ORPHA
370348

Kod OMIM
612219

Kod ICD10
C71.9

Kod ICD11
XH6P76

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*Źródło

orphanet