

## Opis choroby \*

### Definicja

A rare inherited cancer-predisposing syndrome characterized by adult onset of hematologic malignancies mainly affecting the myeloid lineage (such as myelodysplastic syndrome and/or acute myeloid leukemia), less frequently lymphoid malignancies. Some patients have been reported to develop granulomatous or immune disorders (including sarcoidosis, systemic lupus erythematosus, asthma, eczema, or juvenile arthritis) before or in the absence of hematologic malignancies.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

488647

#### Kod OMIM

616871

#### Kod ICD10

C96.7

#### Kod ICD11

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### \*Źródło

orphanet