

## Opis choroby \*

### Definicja

A rare myeloproliferative neoplasm characterized by sustained peripheral blood neutrophilia, bone marrow hypercellularity due to neutrophilic granulocyte proliferation, and hepatosplenomegaly. Other organs may be infiltrated in addition. Microscopically, the bone marrow shows an increase in proportion of myelocytes and mature neutrophils, but no significant dysplasia in any of the cell lineages. Peripheral blood neutrophils are mostly segmented, although band forms may also be substantially increased. Cytogenetic abnormalities are absent in most cases. The disease is slowly progressive with progredient neutrophilia followed by anemia and thrombocytopenia. Transformation to acute myeloid leukemia may occur.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

86829

#### Kod OMIM

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#### Kod ICD10

D47.1

#### Kod ICD11

2A20.1

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### \*Źródło

orphanet