

## Opis choroby \*

### Definicja

A rare, malignant, neoplastic disease characterized by clonal proliferation of myeloid and/or lymphoid precursors harboring translocations or insertions involving the chromosome band 8p11 and the FGFR1 gene, in the blood, bone marrow and often other tissues as well (spleen, liver, lymph nodes, breast, etc.). It usually presents as myeloproliferative neoplasm with eosinophilia, T lymphoblastic lymphoma with eosinophilia or, less frequently, acute myeloid leukemia. The presenting signs and symptoms include eosinophilia, leukocytosis with leukemoid reaction, monocytosis, fatigue, sweating, weight loss, lymphadenopathy, splenomegaly and/or hepatomegaly. Extranodal involvement may include the tonsils, lungs and breasts.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

8p11 myeloproliferative syndrome  
Białaczka/chłoniak komórek macierzystych  
Zespół mieloproliferacyjny 8p11  
Stem cell leukemia/lymphoma

#### Kod ORPHA

168953

#### Kod OMIM

613523

#### Kod ICD10

D47.1

#### Kod ICD11

2A52

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#### [\\*Źródło](#)

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