

Nowotwory szpiku związane rearanżacją PDGFRB

Kod Orpha: 168950 Kod OMIM:

Opis choroby *

Definicja

A rare, malignant, neoplastic disease characterized by clonal proliferation of myeloid and/or lymphoid precursors harboring rearrangements in the PDGFRB gene, in the blood, bone marrow and often other tissues as well (spleen, lymph nodes, skin, etc.). It usually presents as chronic myelomonocytic leukemia with eosinophilia, chronic eosinophilic leukemia, atypical chronic myelogenous leukemia, juvenile myelomonocytic leukemia, myelodysplastic syndrome, acute myeloid leukemia or acute lymphoblastic leukemia. Patients usually present with anemia, leukocytosis, monocytosis, eosinophilia and/or splenomegaly, or systemic symptoms, such as fever, sweating and/or weight loss.

Dane

Klasyfikacja

Choroba

Kod ORPHA

168950

Kod OMIM

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Kod ICD10

D47.1

Kod ICD11

2A51

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

Dostępna na stronie www.orphanet.pl