Opis choroby *

Definicja

A rare, genetic, immune disease characterized by chronic neutrophilia, increase in the percentage of circulating CD34+ cells in peripheral blood, increase in granulocyte precursors in bone marrow and splenomegaly. Patients are predominantly asymptomatic, but may present with systemic inflammatory response syndrome with fever, dyspnea, tachycardia, pleural and pericardial effusion, or myelodysplastic syndrome.

Dane

Klasyfikacja
Choroba

Kod ORPHA
279943

Kod OMIM
162830

Kod ICD10
D72.8

Kod ICD11
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*Źródło

orphanet