Opis choroby *

Definicja

A rare mature B-cell neoplasm characterized by clonal proliferation of B-cell prolymphocytes, with prolymphocytes constituting more than 55% of lymphoid cells in peripheral blood. IG genes are clonally rearranged. Neoplastic cells are present in the bone marrow, peripheral blood, and spleen. Patients usually present with B symptoms, massive splenomegaly but absent or minimal lymphadenopathy, rapidly increasing lymphocyte count, anemia, and thrombocytopenia. Therapy response is poor.

Dane

Klasyfikacja
Choroba

Synonimy
B-PLL
B-plL

Kod ORPHA
86852

Kod OMIM
-

Kod ICD10
C91.3

Kod ICD11
2A82.1

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*Źródło

orphanet