Opis choroby *

Definicja

A rare aggressive B-cell non-Hodgkin lymphoma characterized by a rearrangement in MYC and BCL2 and/or BCL6 (so-called double-hit or triple-hit lymphoma). The category includes double-hit cases with features intermediate between diffuse large B-cell lymphoma (DLBCL) and Burkitt lymphoma, blastoid cases with a double-hit, and cases with a DLBCL, not otherwise specified, morphology with a double-hit. It refers only to <i>de novo</i> cases, not to lymphomas with a history of pre-existing or coexistent indolent lymphoma. Patients typically present with widespread disease, including involvement of lymph nodes, bone marrow, and central nervous system.

Dane

Klasyfikacja

Choroba

Kod ORPHA Kod OMIM 480541

Kod ICD10 C85.7

Kod ICD11

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*Źródło

orphanet