

Opis choroby *

Definicja

A very severe type of RAEB characterized by cytopenias and the following hematological parameters: uni- or multilineage dysplasia, 10% to 19% blasts in bone marrow or 5% to 19% in peripheral blood, variable presence of Auer rods (abnormal, needle-shaped or round inclusions in the cytoplasm of myeloblasts and promyelocytes). Median survival has been reported to be 18 months.

Dane

Klasyfikacja	Synonimy
Podtyp kliniczny	RAEB-2 RAEB-2

Kod ORPHA	Kod OMIM	Kod ICD10
100020	-	D46.2

Kod ICD11
2A35

*Źródło

orphanet