

## Opis choroby \*

### Definicja

A severe type of RAEB characterized by cytopenias and the following hematological parameters: uni- or multilineage dysplasia, 5% to 9% blasts in bone marrow or 2% to 4% in peripheral blood, and no 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

<b>Klasyfikacja</b>	<b>Synonimy</b>
Podtyp kliniczny	RAEB-1 RAEB-1

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
100019	-	D46.2

**Kod ICD11**  
2A35

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### \*Źródło

orphanet