Opis choroby *

Definicja

A rare myeloproliferative neoplasm characterized by stem-cell derived clonal over proliferation of mature myeloid lineages, such as erythrocytes, leukocytes, and megakaryocytes, with variable degrees of megakaryocyte atypia, associated with reticulin and/or collagen bone marrow fibrosis, osteosclerosis, ineffective erythropoiesis, angiogenesis, extramedullary hematopoiesis, and abnormal cytokine expression.

Dane

Klasyfikacja
Choroba

Synonimy
Agnogenic myeloid metaplasia
Agnogenic myeloid metaplasia
Mielofibroza idiopatyczna
Mielofibroza pierwotna
Mieloskleroza z metaplazją szpiku
Idiopathic myelofibrosis
Myelofibrosis with myeloid metaplasia
Osteomyelofibrosis

Kod ORPHA
824

Kod OMIM
254450

Kod ICD10
D47.4

Kod ICD11
2A20.2

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*Źródło

orphanet