Opis choroby *

Definicja

A rare neoplastic disease characterized by infantile to childhood onset of evidence of bone marrow insufficiency/failure associated with increased risk for myelodysplastic syndrome or acute myeloid leukemia. Most patients present with petechiae, easy bruising, or anemia. Rapid progression is common, and prognosis is generally poor.

Dane

Klasyfikacja
Choroba

Kod ORPHA
495930

Kod OMIM
252270

Kod ICD10
D46.7

Kod ICD11
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*Źródło

orphanet