Opis choroby *

Definicja

A rare, benign, red cell aplasia of young children or infants characterized by a normocytic normochromic anaemia with severe reticulocytopenia in otherwise normocellular bone marrow, and a complete spontaneous recovery within 1-2 months after diagnosis. Neutropenia and thrombocytosis may be associated findings at diagnosis, and a history of a preceding viral illness is frequent. No organomegaly is observed.

Dane

Klasyfikacja
Choroba

Synonimy
Transient acquired pure red cell aplasia
Przejściowa nabyta aplazja czysto czerwonokrwinkowa

Kod ORPHA
98871

Kod OMIM
227050

Kod ICD10
D60.1

Kod ICD11
3A61.0

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*Źródło

orphanet