Transient erythroblastopenia of childhood

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Transient erythroblastopenia of childhood. ORPHA:98871

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.