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Hemophagocytic **lymphohistiocytosis** (HLH) is an unusual syndrome characterized by fever, splenomegaly, jaundice, and the pathologic finding of hemophagocytosis (phagocytosis by macrophages of erythrocytes, leukocytes, platelets, and their precursors) in bone marrow and other tissues. HLH may be diagnosed in association

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Retrospective Study

Efficient management of secondary haemophagocytic lymphohistiocytosis with intravenous steroids and γ -immunoglobulin infusions

Georgiadou S *et al.* γ -immunoglobulin infusions for sHLH

Sarah Georgiadou, Nikolaos K Gatselis, Aggelos Stefos, Kalliopi Zachou, Konstantinos Makaritsis, Eirini I Rigopoulou, George N Dalekos

Abstract

BACKGROUND

Secondary haemophagocytic lymphohistiocytosis (sHLH) is a rare life-threatening condition mainly associated with underlying infections, malignancies, and autoimmune or immune-mediated diseases.

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