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Hepatic langerhans cell histiocytosis: A review

Hepatic LCH

Zhiyan Fu, Hua Li, Mustafa Erdem Arslan, Peter F. Ells, Hwajeong Lee

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Author: Adelaine Wong, Clara L. Ortiz-Neira, Wali...

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Jun 01, 2015 · Langerhans cell histiocytosis: a comprehensive review DINA EL DEMELLAWY^{1,2}, JAMES LEE YOUNG¹, JOSEPH DE NANASSY^{1,2}, ELIZAVETA CHERNETSOVA³ AND AHMED NASR⁴ ¹Department of Pediatric Pathology, University of Ottawa, ²Department of Pediatric Pathology, Children's Hospital of Eastern Ontario, ³Department of Innovations in Medical Education ...

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Author: Hua Li, Peter Ells, Mustafa Erdem Ar...

Publish Year: 2020

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Jan 23, 2021 · Langerhans cell histiocytosis (LCH) is **the most common histiocytic disorder**. Liver involvement is seen in 10.1%-19.8% of patients with LCH and can lead to secondary sclerosing cholangitis requiring liver transplantation (LT). We aimed to describe the characteristics and outcomes of patients undergoing LT for LCH.

Author: Ioannis A. Ziogas, Christos D. Kakos, W... **Publish Year:** 2021

Langerhans cell histiocytosis

Langerhans cell histiocytosis is an abnormal clonal proliferation of Langerhans cells, abnormal cells deriving from bone marrow and capable of migrating from skin to lymph nodes. Symptoms range from isolated bone lesions to multisystem disease. LCH is part of a group of syndromes called histiocytoses, which are characterized by an abnormal proliferation of histiocytes. These diseases are related to other forms of abnormal proliferation of white blood cells, such as leukemias and lymphomas.



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