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Pulmonary Langerhans cell histiocytosis in adults: A case report

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Pulmonary Langerhans cell histiocytosis is an **uncommon diffuse cystic lung disease** in adults. In rare cases, it can involve extrapulmonary organs and lead to endocrine abnormalities such as central diabetes insipidus. A 42-year-old man presented with polyphagia and polydipsia, as well as a dry cough and dyspnea on exertion.

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Langerhans cell histiocytosis

Rare Disease

Langerhans cell histiocytosis is a rare disease involving clonal proliferation of Langerhans cells, abnormal cells deriving from bone marrow and capable of migrating from skin to lymph nodes. Clinically, its manifestations range from isolated bone lesions to multisystem disease. LCH is part of a group of clinical 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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Pulmonary Langerhans cell histiocytosis in adults: A case report

Feng-Feng Wang, Ya-Shuang Liu, Wei-Bo Zhu, Yan-Dong Liu, Yao Chen

Abstract

BACKGROUND

Langerhans cell histiocytosis (LCH) is a rare disease of unknown aetiology. While it may affect any organ of the body, few cases of solitary lung involvement are published

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Oct 01, 2015 · **Pulmonary Langerhans cell histiocytosis** (**LCH**) is a rare **interstitial lung disease** in **adults** that is a part of diseases caused by infiltration of **Langerhans cell** across multiple organs, such as the lungs, bones, skin, **pituitary gland**, and **lymph** nodes1. Among **cases** of multi-system LCH, simultaneous invasion of the lung and the **pituitary gland** is uncommon.

Cited by: 1 **Author:** Yeun Seoung Choi, Jung Soo Lim, Woo...
Publish Year: 2015

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Author: Mohamad Jihad Mansour, Elias Mokb... **Publish Year:** 2017

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Langerhans cell histiocytosis
Rare Cancer

Langerhans cell histiocytosis is a rare cancer involving clonal proliferation of Langerhans cells, abnormal cells deriving from bone marrow and capable of migrating from skin to lymph nodes. Clinically, its manifestations range from isolated bone lesions to multisystem disease. LCH is part of a group of clinical 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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