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Ataxia-telangiectasia complicated with Hodgkin's lymphoma: A case report

Li XL *et al.* AT with Hodgkin's lymphoma

Xiao-Ling Li, Yi-Lin Wang

Abstract

BACKGROUND

Ataxia-telangiectasia (AT) is a rare, autosomal recessive, multisystem disorder. Because most clinicians have low awareness of the disease, only scarce reports of AT exist in the literature, especially of cases with lymphoma/leukemia.

CASE SUMMARY

A 7-year-old girl with history of recurrent respiratory tract infections was referred to our department because of unstable walking for 5 years and enlarged neck nodes for 2-mo duration. Physical examination revealed scleral telangiectasia and cerebellar ataxia. Elevated alpha-fetoprotein, decreased serum immunoglobulin and decreased T cell function were the major findings of laboratory examination. Histological analysis of cervical lymph node biopsy was suggestive of classical Hodgkin's lymphoma. Genetic examination showed heterozygous nucleotide variation of

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Hodgkin lymphoma in a young child contributing to a ...

https://link.springer.com/article/10.1007/s12308-010-0062-8

Jun 15, 2010 · We report the clinical, histopathologic, and molecular features of a 6-year-old child who presented with EBV-positive Hodgkin lymphoma (HL), which led to the diagnosis of ataxia telangiectasia. The diagnosis of HL at this unusually young age prompted further clinical, immunologic and cytogenetic evaluations, all of which supported a diagnosis ...

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Ataxia-Telangiectasia Complicated by Craniopharyngioma – A ...

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Ataxia-telangiectasia is a rare autosomal recessive neurodegenerative disorder. It is characterized by ocular telangiectasia, cerebellar degeneration, cellular and humoral immunodeficiency, and a high predisposition to cancer. Here, we report a case of a girl with ataxia-telangiectasia who developed craniopharyngioma and non-Hodgkin's lymphoma.

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Ataxia telangiectasia: A report of two cousins and review ...

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The initial clinical description of ataxia telangiectasia (A-T) was reported by Syllaba and Henner in 1926, and the syndrome was first signed to the term "AT" by Boder and Sedgwick in 1957. The prevalence of AT disease has estimated to be one case in 40,000 to one case in 100,1000.[1]

[PDF] Ataxia telangiectasia - A case report

www.e-mjm.org/1980/v35n2/ataxia-telangiectasia.pdf

ATAXIA TELANGIECTASIA - A Case Report A.KANAGANAYAGAM SUMMARY CASE REPORT The patient is a 17 year old Indian male who was first seen at the University Hospital in May 1973 at the age of 10 years. He presented then with a history of not being able to walk by himself for a period of one year. Apparently this difficulty in walking had been ...

Non-Hodgkin B-cell Lymphoma of the Ovary in a Child with ...

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We present the first case of a primary ovarian non-Hodgkin B-cell lymphoma, of Burkitt's type morphology, in a child with AT. Case An 11-year-old girl was admitted to the pediatrics ward at a tertiary care children's hospital with a 5-day history of diffuse abdominal pain, ...

Cited by: 3 Author: C.S. Danby, L. Allen, M.D. Moharir, S. Weitz... Publish Year: 2013

Ataxia Telangiectasia and Lymphoma: An Indication for ...

https://www.tandfonline.com/doi/abs/10.3109/08880019509029550

Ten of 18 children in a highly inbred Arab kindred suffered from either ataxia telangiectasia (AT) or a variant syndrome consisting of ataxia, microcephaly, and congenital cataract (AMC). Four of the nine afflicted children were treated in our unit when they developed lymphomas (both Hodgkin's and non-Hodgkin's including Burkitt's). They were given chemotherapy (either standard COMP or low ...

Cited by: 24 Author: M. Weyl Ben Arush, J. Rosenthal, J. Dale, Y... Publish Year: 1995

Ataxia Telangiectasia - NORD (National Organization for ...

https://rarediseases.org/rare-diseases/ataxia-telangiectasia

Signs and symptoms	Genetics	Purpose	Availability
<p>Ataxia telangiectasia (AT) is a complex genetic neurodegenerative disorder that may become apparent during infancy or early childhood. The disorder is characterized by progressively impaired coordination of voluntary movements (ataxia), the development of reddish lesions of the skin and mucous membranes due to permanent widening of groups of blood vessels (telangiectasia), and impaired functioning of the immune system (i.e., cellular and humoral immunodeficiency), resulting in increased susceptibility to up...</p> <p>See more on rarediseases.org</p>			

Ataxia-Telangiectasia - St. Jude Children's Research Hospital

https://www.stjude.org/disease/ataxia-telangiectasia.html

People with ataxia-telangiectasia should be managed by a doctor who knows the condition well. Children with ataxia-telangiectasia should have regular screenings to detect cancer as early as possible. The goal of screening is finding and treating cancer early to allow the best outcome for patients.

Ataxia Telangiectasia - National Cancer Institute

https://www.cancer.gov/.../genetics/ataxia-fact-sheet

Ataxia telangiectasia (A-T) is a primary immunodeficiency disease that affects a number of different organs in the body. An immunodeficiency disease is one that causes the immune system to break down, making the body susceptible to diseases. It is a rare, recessive genetic disorder of childhood...

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Ataxia-telangiectasia

Rare Neurodegenerative Autosomal Recessiv

Ataxia-telangiectasia, also referred to as ataxia-telangiectasia syndrome or Louis-Bar syndrome, is a rare, neurodegenerative, autosomal recessive disease causing severe disability. Ataxia refers to poor coordination and telangiectasia to small dilated blood vessels, both of which are hallmarks of the disease.



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To our knowledge, this is the first case of ataxia telangiectasia complicated by **craniopharyngioma** in the English literature. Introduction Ataxia-telangiectasia is a rare **autosomal recessive neurodegenerative disorder**.

Ataxia telangiectasia

Rare Neurodegenerative Autosomal Recessive Disease

Ataxia-telangiectasia, also referred to as ataxia-telangiectasia syndrome or Louis-Bar syndrome, is a rare, neurodegenerative, autosomal recessive disease causing severe disability. It refers to poor coordination and ataxia, telangiectasia to small dilated blood vessels, both of which are hallmarks of the disease.

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Ataxia-Telangiectasia Complicated by Craniopharyngioma – A ...

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