

Name of Journal: *World Journal of Clinical Cases*

Manuscript NO: 58955

Manuscript Type: CASE REPORT

An atypical adult-onset Still's Disease with an initial and sole manifestation of liver injury: A case report and review of the literature

Liver injury secondary to adult-onset Still's Disease

Abstract

BACKGROUND

Adult-onset Still's disease (AOSD) typically presents with a high spiking fever, polyarthritis, transient maculopapular rash, neutrophilic leukocytosis, and hepatosplenomegaly. It has a wide clinical spectrum ranging from mild to severe, with extensive involvement of almost every organ. Although liver involvement in the form of increased hepatic enzymes and bilirubin is common, no AOSD case with liver involvement as the initial manifestation of AOSD have been reported.

Match Overview

1	Crossref 30 words Hao Jiang, Chong Jin, Jing-Gang Mo, Lie-Zhi Wang, Lei Ma ... Kun-Peng Wang. "Rare recurrent gallstone ileus: A case repor	1%
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Unusual presentation of adult-onset Still's Disease with liver injury



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An Unusual Presentation of Adult-Onset Still's Disease in ...

<https://www.sciencedirect.com/science/article/pii/S0002962920302834>

Jun 27, 2020 · Introduction. **Adult-Onset Still's Disease** (AOSD) is a very uncommon **disease** of unknown etiology that can be triggered by both genetic factors and a mosaic of infectious **diseases**.^{1, 2, 3} It commonly presents with fever, rash and musculoskeletal symptoms like arthralgias or myalgias. Diagnosis is usually clinical after excluding any other possible causes of ...

Author: Arsany Anis, Kok Hoe Chan **Publish Year:** 2020

Unusual presentation of more common disease/injury: Adult ...

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4545114>

Background. Juvenile arthritis (**Still's disease**) was not considered as an adult **disease** until 1971, when Bywaters¹ distinguished an adult variant of the same (**adult-onset Still's disease**, AOSD). Its incidence is estimated at 0.16/100 000 in France,² with a median age of 25 years at the onset.³ It may be present as a severe polysystemic **disease**^{4–8} with a very heterogeneous clinical ...

Cited by: 7 **Author:** Roderich Meckenstock, Audrey Therby, ...

Publish Year: 2012

An atypical presentation of adult-onset Still's disease ...

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4860549>

An atypical **presentation of adult-onset Still's disease** complicated by pulmonary hypertension and macrophage activation syndrome treated with immunosuppression: **a case-based review of the literature**. ... improved with amlodipine and completely resolved with the use of the IL-1 antagonist anakinra.³ The most recent **case report** is of a 38-year ...

Cited by: 12 **Author:** Mili V. Mehta, Daniel K. Manson, Evelyn ...

Publish Year: 2016

An Unusual Case of Adult-Onset Still's Disease with ...

<https://www.hindawi.com/journals/crirh/2014/128623> ▾

2. Case Report. A 34-year-old African American female was diagnosed with adult-onset Still's

An atypical adult-onset Still's Disease with an initial and sole manife



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An atypical presentation of adult-onset Still's disease ...

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4860549>

An atypical presentation of **adult-onset Still's disease** complicated by pulmonary hypertension and macrophage activation syndrome treated with immunosuppression: a **case-based review of the literature**.
... **Adult-onset Still's disease** (AOSD) is a rare multisystemic autoinflammatory disorder that affects young adults. ...

Cited by: 12

Author: Mili V. Mehta, Daniel K. Manson, Evelyn M. ...

Publish Year: 2016

An Atypical Presentation of Adult-Onset Still's Disease ...

<https://journals.sagepub.com/doi/full/10.1086/685112>

Mar 01, 2016 · An Atypical Presentation of **Adult-Onset Still's Disease** Complicated by Pulmonary Hypertension and Macrophage Activation Syndrome Treated with Immunosuppression: A **Case-based Review of the Literature** Mili V. Mehta, Daniel K. Manson, Evelyn M. Horn, and Jennifer Haythe

Cited by: 12

Author: Mili V. Mehta, Daniel K. Manson, Evelyn M. ...

Publish Year: 2016

A case report of Adult-onset Still's disease presenting ...

<https://journals.sagepub.com/doi/full/10.1177/2054270419894834>

Apr 02, 2020 · A study of 57 patients by Gerfaud-Valentin et al. in 2014 suggested a mean time to diagnosis of four months. 4 A **case** published in 2017 by Stella Pak and Cindy Pham shows similarities to our **case** where a 31-year-old man was diagnosed with **Adult-onset Still's disease** three years after his **initial** presentation. 5 This highlights the importance ...

Author: Robert Perry, Dimitrios Christidis, Andre...

Publish Year: 2020



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[An atypical presentation of adult-onset Still's disease ...](#)

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Adult-onset Still's disease (AOSD) is a rare multisystemic autoinflammatory disorder that affects **young adults**. It is characterized by major criteria of quotidian temperatures $>39^{\circ}\text{C}$, arthralgias/arthritis, evanescent salmon-pink rash, lymphadenopathy, pharyngitis, and hepatosplenomegaly.

Cited by: 12**Author:** Mili V. Mehta, Daniel K. Manson, Evelyn ...**Publish Year:** 2016

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Apr 02, 2020 · In conclusion, this **case** highlights the importance of considering **Adult-onset Still's disease** in patients with a presumed inflammatory disorder of unknown aetiology. We also highlight the rare association between **Adult-onset Still's disease** and acute fibrinous and organising pneumonia, and the fact that an associated collagen vascular ...

Author: Robert Perry, Dimitrios Christidis, And...**Publish Year:** 2020

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Adult onset Still's disease (AOSD) is a rare inflammatory disorder of unknown etiology that usually affects young adults. Very few patients older than 70-year-old have been reported.

[\(PDF\) Adult onset Still's disease: Clinical experience ...](#)