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8
Adult-onset Still's disease evolving with multiple organ failure and death: case
report and literature review

Adult-onset Still's disease

Abstract

BACKGROUND

Adult-onset Still's Disease (AOSD) is a rare systemic inflammatory disease, which is characterized by daily fever and arthritis, with an evanescent rash and neutrophilic leukocytosis. To date, there is no definite laboratory or imaging test available for diagnosing AOSD; the diagnosis is one of exclusion, which can be very challenging. In particular, AOSD patients may experience different complications affecting their clinical picture, management and prognosis. The treatment of AOSD remains largely empirical.

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Adult Still's Disease

Medical Condition

A rare auto inflammatory condition characterized by fever, rash and joint pain.

Very rare (Fewer than 20,000 cases per year in US)

No vaccine available

Often requires lab test or imaging

Treatment from medical professional advised

Can last several months or years

The cause of adult still's disease is not clear and is thought to be caused by viral or bacterial infection. Fever, rashes, sore throat, muscle pain and painful swollen joints are the commonly noted symptoms. Treatment consists of anti-inflammatory agents.

Symptoms

Most commonly noted symptoms are:

- Fever
- Rashes
- Sore throat
- Achy and swollen joints
- Muscle pain

Treatments

Treatment aims at managing inflammation, preventing end-organ damage, and minimizing risk of adverse effects of therapy.

Medication

- Nonsteroidal anti-inflammatory drugs (NSAIDs): Used to reduce inflammation and relieve pain.