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Systemic lupus erythematosus combined with primary hyperfibrinolysis and protein C and protein S deficiency: A case report

Liao YX *et al.* SLE combined with primary hyperfibrinolysis

Yi-Xuan Liao, Yan-Fei Guo, Yu-Xia Wang, Ai-Hua Liu, Chun-Li Zhang

Abstract

14 BACKGROUND

Systemic lupus erythematosus (SLE) is an autoimmune disease characterized by systemic involvement and multiple autoantibodies in the serum. Patients with protein C (PC) and protein S (PS) deficiency are prone to thrombosis. In contrast, patients with primary hyperfibrinolysis tend to bleed.

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Introduction: Associations between systemic lupus erythematosus (SLE) and primary immunodeficiencies (PIDs) were analyzed to gain insight into the physiopathology of SLE. Some PIDs have been consistently associated with SLE or lupus-like manifestations: (a) homozygous deficiencies of the early components of the classical complement pathway in the following decreasing order: in C1q, ... Cited by: 79 Author: Magda Carneiro-Sampaio, Bernadete Lourd... Publish Year: 2008

Systemic Lupus Erythematosus (SLE) Clinical Presentation ...

Mar 18, 2020 · Systemic lupus erythematosus (SLE) is a chronic inflammatory disease that has protean manifestations and follows a relapsing and remitting course. More than 90% of cases of SLE occur in women, frequently starting at childbearing age.

PEOPLE ALSO ASK

- Is systemic lupus erythematosus genetic?
- Is SLE an autoimmune disease?
- Is sle a genetic disorder?
- What is photosensitive lupus erythematosus?

Feedback

Systemic lupus erythematosus: MedlinePlus Genetics

Systemic lupus erythematosus (SLE) is a chronic disease that causes inflammation in connective tissues, such as cartilage and the lining of blood vessels, which provide strength and flexibility to structures throughout the body. Explore symptoms, inheritance, genetics of this condition.

Systemic Lupus Erythematosus (SLE) Genetics: Practice ...

May 05, 2020 · Systemic lupus erythematosus (SLE) is a chronic inflammatory autoimmune disorder associated with a wide range of symptoms and physical findings. The risk of developing SLE is, at least in part, genetic, but it is a complex genetic illness with no clear mendelian pattern of inheritance.

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Systemic Lupus Erythematosus

Medical Condition

An autoimmune disease, with systemic manifestations including skin rash, erosion of joints or even kidney failure.

- Common (More than 200,000 cases per year in US)
- Often requires lab test or imaging
- Treatments can help manage condition, no known cure
- Can be lifelong

Causes can be heredity, ultra violet exposure and certain drugs Most common symptom is butterfly rash on face. Medication to reduce inflammation.

Symptoms

Signs and symptoms may include:

- Butterfly rash on the face
- Appetite loss
- Hair loss
- Fever
- Fatigue
- Photosensitivity
- Raynaud's phenomenon
- Pleuritis
- Pericarditis
- Painful or swollen joints and muscle pain
- Unexplained fever
- Chest pain upon deep breathing
- Swelling (edema) in legs or around eyes
- Mouth ulcers
- Chorea

Treatments

The treatment to reduce and manage symptoms.

Medication

- Anti-inflammatory drugs: Given as oral medications, ointments, or intravenous injections. Ibuprofen · Naproxen · Diclofenac
- Steroid cream: Reduces the rashes. Triamcinolone · Fluocinolone
- Immuno suppressants: Reduces the rashes. Azathioprine · Methotrexate · Mycophenolate

Causes

Exact cause is not known, but it is believed that the following can cause the disease

Genetics:

Families who have the history of having this disease are more likely to develop

Environmental Factors:

- Exposure to ultra violet rays
- Viruses
- Medications or certain drugs
- Emotional stress
- Trauma

Hormones:

- Females are more commonly affected than men
- Women during pregnancy are high risk of developing this condition
- It is believed that estrogen, a sex hormone can be responsible for this

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Cited by: 79 Author: Magda Carneiro-Sampaio, Bernadete Lou...

Publish Year: 2008

Systemic Lupus Erythematosus (SLE) Differential Diagnoses

<https://emedicine.medscape.com/article/332244-differential>

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PEOPLE ALSO ASK

What is systemic lupus erythematosus? ▾

What is protein C deficiency? ▾

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Can propylthiouracil cause SLE? ▾

Feedback

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Study of Recombinant Human B ... - ClinicalTrials.gov

<https://clinicaltrials.gov/ct2/show/NCT04082416>

Sep 09, 2019 Primary Purpose: Treatment: Official Title: A Phase III, Placebo-Controlled, Multi-Center, Randomized, Double-Blind, Dose-exploring Trial of RC18, a Recombinant Human B Lymphocyte

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