

LETTER TO THE EDITOR

ADULT-ONSET STILL'S DISEASE EVOLVING WITH MULTIPLE ORGAN FAILURE: CASE REPORT AND LITERATURE REVIEW

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INTRODUCTION

Adult-onset Still's disease (AOSD) is a disorder similar to systemic onset juvenile idiopathic arthritis (Still's disease), affecting young adults worldwide.¹⁻² Rarely, AOSD can be life-threatening.³⁻⁵ From Brazil, where recurrent forms are prevalent,⁶ we report on a case of AOSD presenting with multiple organ failure.

CASE DESCRIPTION

A 49-year-old white woman with a 3-month fever and weight loss history presented to our hospital. She had symmetric arthritis of hands, facial and thoracic skin rash, high morning fever, nausea, myalgia, and sore throat. The hemoglobin level was 11.4 mg/dL, white blood cell count was 16900/mm³ (bands, 3600/mm³), platelet count was 386000/ μ L, aspartate and alanine aminotransferases were 94 and 86 U/L, respectively, and erythrocyte sedimentation rate (ESR) was 45 mm/h. Autoimmune and infective serological tests (eg, viral hepatitis tests, anti-immunodeficiency virus antibody, antinuclear antibody rheumatoid factor and mononucleosis antibodies) were negative; urinalysis and chest X-ray were normal. *Staphylococcus aureus* was present in 1 of 5 samples of blood cultures, and oxacyllin plus gentamicin therapy was started, but a transesophageal echocardiogram showed no vegetations. Abdominal computed tomography revealed nothing but splenomegaly.

After an initial improvement, the high fever and skin rash returned, and cervical lymphadenopathy was detected. Lymph node, bone marrow, and rash cutaneous biopsies were performed. Multiple organ failure (cardiovascular, pulmonary, renal, and hematologic) motivated intensive care unit admission, where management of septic shock, including hydrocortisone, succeeded. Lymph node biopsy revealed

chronic inflammatory infiltrate. Cultures were negative. A bone marrow specimen showed diffuse hypercellularity. The tuberculin test reading was impaired due to rash. A high serum ferritin level (> 2000 ng/mL) was detected, and cutaneous biopsy showed vacuolar interface dermatitis, interstitial mucinosis, and little superficial chronic inflammatory infiltrate, unspecific findings of autoimmune diseases. The patient gradually improved and was discharged asymptomatic from the hospital with oral prednisone after completing antibiotic treatment. In outpatient follow-up, slow tapering of prednisone is in course without evidence of recurrence.

DISCUSSION

Adult-onset Still's disease was first described by Bywaters in 1971.¹ It is an uncommon disorder (incidence of 0.16 cases/100000 persons/year), presenting mostly with arthritis, spiking daily fever, a salmon-colored rash, lymphadenopathy, leukocytosis, and elevated ESR.^{2,7} High-sensitivity classification criteria have been proposed, since there is no single test to establish the diagnosis (Table 1).^{8,9} In Brazil, disease onset is around 30 years.^{6,10} Fever, arthritis, and skin rash are the most common clinical signs, and elevated ESR, leukocytosis, absent antinuclear antibody, and absent rheumatoid factor are the most frequent laboratory findings.^{6,10-11}

Table 1 - Classification criteria for adult-onset Still's disease proposed by Yamaguchi et al

Major criteria	Minor criteria
Temperature of > 39°C for > 1 wk	Sore throat
Leukocytosis > 10 000/mm ³	Lymph node enlargement
Typical rash	Splenomegaly
Arthralgias > 2 wk	High transaminases
	Negative ANA, RF

After excluding infections, malignancies, and other rheumatic diseases, adult Still's should be considered if 5 criteria (2 of which being major ones) are met. ANA = antinuclear antibody; RF = rheumatoid factor

By the installation of multiple organ failure, the diagnosis had not yet been confirmed. Large spectrum antibiotics

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Table 2 - Previous case reports of adult-onset Still's disease evolving with organ failure

Gender / Age	Organ failure	Antibiotic use	Years since first symptoms	Previous immuno-suppressive therapy	Follow-up	Year / Country
F / 49	R; H; CV; P	Yes	0	No	OMT	2007 / Brazil
M / 47	H, C, P	Yes	3	Yes	Death	2001 / France
F / 29	H, CV, P	Yes	20	No	OMT	1999 / USA
F / 23	H, R, C, P	Yes	2	No	OMT	1991 / Canada

F = female; M = male; OMT = Outpatient maintenance therapy; R = renal; H = hematological; CV = cardiovascular; P = pulmonary, C = central nervous system

were mandatory, since for diagnosis of AOSD, a fever not related to infection is always one criterion of inclusion. After innumerable negative cultures, steroid response, and the fulfilling of classification criteria (Table 1), the diagnosis could be established. Lack of hemophagocytosis and hypofibrinogenemia with elevated ESR allowed the exclusion of macrophage activation syndrome.¹²⁻¹³

Only 3 cases of AOSD evolving with shock and multi-

ple organ failure have been reported (Table 2), and all of them had previous medical data facilitating early diagnosis.⁵⁻⁷ Treatment of severe cases requires glucocorticoids, sometimes immunosuppressants (eg, methotrexate), and biological agents (anti-tumor necrosis factor-alpha, anti-interleukin-1, anti-interleukin-6 agents, and most recently anti-CD20-expressing B-cell antibodies).¹³⁻¹⁴ Due to its rarity, prognosis of the severe onset form is still unknown.¹⁵

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