CASE REPORT

Fever, sore throat and myalgia

P Ive, FCP (SA); M Mendelson, PhD, FRCP; S Dlamini, FCP (SA), Cert ID (SA) Phys

Division of Infectious Diseases and HIV Medicine, Department of Medicine, Faculty of Health Sciences, University of Cape Town and Groote Schuur Hospital, Cape Town, South Africa

Corresponding author: P Ive (prue.ive@gmail.com)

A 20-year-old man presented with a severely sore throat and myalgia, which were unresponsive to antibiotics. He was admitted to a regional hospital with an ongoing painful throat, generalised myalgia, fever and a transient, recurring, salmon-pink rash on his hands and trunk. He did not respond to ceftriaxone and had a continual significant fever daily.

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A 20-year-old man presented with a severely sore throat and myalgia, which were unresponsive to antibiotics. Ten days into his illness he

was admitted to a regional hospital with an ongoing painful throat, generalised myalgia, fever (38.5°C) and a transient, recurring, salmon-pink rash on his hands and trunk. He did not respond to ceftriaxone and had

a continual significant fever daily. Shotty cervical lymphadenopathy was noted. The relevant laboratory investigations are shown in Table 1.

Table 1. Relevant investigations during the patient's illness Day 8 Day 11 Day 16 Day 23 Day 25 Haemoglobin (g/dL) 13.5 12.6 12.3 10.3 10.0 White blood cell count (× 109/L) 28.1 33.2 23.0 28.1 28.8 Polymorphonuclear neutrophil 21.3 26.58 $(\times 10^{9}/L)$ Alanine aminotransferase (U/L) 55 66 303 C-reactive protein (mg/L) Ferritin (µg/L) 9 405 14 448 Creatine kinase (U/L) 38 Negative Blood culture Negative Antinuclear antibody, rheumatoid Negative factor Antistreptolysin Negative

On admission to tertiary care, an aetiological differential diagnosis for the pyrexia of unknown origin (PUO) $^{[1]}$ (Table 2) included retropharyngeal abscess, Lemierre's syndrome, HIV seroconversion, and adultonset Still's disease (AOSD). A slightly enlarged left adenoid was noted that exuded a small volume of pus on biopsy, while histology revealed lymphoid hyperplasia. A fourth-generation HIV enzyme-linked

immunosorbent assay (ELISA) was negative. A computed tomography (CT) scan and Doppler test excluded retropharyngeal abscess and Lemierre's syndrome. The patient fulfilled the Yamaguchi criteria^[2] (Table 3) for AOSD,[3] with a good clinical and laboratory response to prednisone

When the aetiological diagnosis eludes the clinician, patients with PUO are often diagnosed as having AOSD. As steroids suppress fever and

80 mg once daily.

Table 2. Leading causes of classic pyrexia of unknown origin				
Infection	Neoplasms	Connective tissue	Other (geographical)	
Tuberculosis	Lymphoma	Still's disease	Familial Mediterranean fever	
Occult bacterial abscess	Renal carcinoma	Variants of rheumatoid arthritis	Kikuchi-Fujimoto disease	
Endocarditis	Atrial myxoma	Systemic lupus erythematosus	Melioidosis	
Brucellosis		Temporal arteritis		
		Polymyalgia rheumatica		
Pyrexia of unknown origin defined as temperature $>38.3^{\circ}\text{C}$ for >3 weeks, with >2 outpatient visits or 3 days' inpatient investigations.				

CONTINUING MEDICAL EDUCATION

Major	Minor	
Fever of at least 39°C lasting at least 1 week	Sore throat	
Arthralgias and arthritis lasting ≥2 weeks	Lymphadenopathy	
Non-pruritic macular or maculopapular skin rash, salmon-coloured, usually over trunk and extremities during febrile episodes	Abnormal liver function tests (particularly elevations in aspartate and alanine aminotransferase)	
Leucocytosis ($\geq 10~000/\mu L$), with at least 80% granulocytes	Negative tests for antinuclear antibody and rheumatoid factor	
	Hepatomegaly or splenomegaly	

inflammation, idiopathic PUO would respond to this treatment because its natural history is to abate. However, the key findings in this case of unrelenting quotidian fever, evanescent rash, marked neutrophilia, and ferritin >10 000 $\mu g/L$ are highly suggestive of AOSD. A low percentage of glycosylated fraction of ferritin is another pointer to the diagnosis, but it cannot be determined in most laboratories in South Africa.^[4]

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