



9/16/25 Morning Report with @CPSolvers

"One life, so many dreams" Case Presenter: Akash Gupta Case Discussants: Andrew Sanchez (@ASanchez_PS) and Jeffrey Shen
<https://clinicalproblemsolving.com/present-a-case/>



Scribing (Gillian Z)

CC: 87 yo female **fever, fatigue, recent falls**

HPI: Stable for 7 yrs until last summer (on pred 1 mg 3 days/week)

Difficulty using hands (playing piano)

Worsening fatigue, weakness

CK 690 → CK 861

Falling in the past few years, more recently

Stopped steroids (concern for myopathy)

Stopped tocilizumab due to diverticulitis

Worsening fatigue, weakness → present to ED

ROS: no hA, no vision change, no oral/nasal ulcer, no joint swelling, no rashes, no belly pain/diarrhea, constipation.

PMH:

Giant cell arteritis: Diagnosed w/ GCA 2010, fevers 100 F, Fatigue, night sweats, jaw pain No headache or vision changes, Prednisone 70 mg monotherapy until 2017 → tocilizumab; pred slowly weaned over yrs until last summer BL temporal artery biopsy showed BL infiltrate w/ lymphs, plasma cell, neutrophils (no giant cells).

TAVR for aortic stenosis

Adrenal insufficiency (secondary chronic glucocorticoid use)

B12 def

HTN, prediabetes

BL hip prosthesis, cholecystectomy

Meds:

Amlodipine, metformine, aspirin, PoB12

Fam Hx:

None

Social Hx:

retired

Health-Related

Behaviors:

No tobacco, alcohol, drugs, supplement
Allergies:

Vitals: T: 98 (up to 99.6) HR: 71 BP: 125/65 RR: 18 Sat: 96 RA BMI:

Exam: Gen: no acute distress

HEENT: normal

CV, Pulm, Abd: nl

Musc: 5/5 strength in shoulder, triceps, biceps, 4/5 wrist flexion, extension, right grip strength 3/5, right knee flexion/extension 3/5, couldn't get off bed doing sit to stand

Notable Labs & Imaging:

Hematology:

WBC: 11.9 (nl) Hgb: 12 Plt: 319 MCV: nl

Chemistry:

Na: 131 K: 4.9 Cl: 97 HCO₃: 27 Cr: 0.74 (baseline) BUN: 16 Glucose: 106 AST: nl ALT: nl Alk-P: Bili: Albumin: 2.9 Total Protein: 8.3 hs trop: 190

ESR: 129 → >130 CRP: 22 (reference 5) LDH: CK: 690 → 861 (after stopping steroids) → 600s → 208 Aldolase: 8;

Low titer PL7: 13, Positive CN1A antibody, TSH: 5.68 FT4: 0.94

After muscle biopsy: p-ANCA +; MPO: 113

UA: Cloudy, Neg ketones, protein 30, large blood, small esterase, RBCs 20-50

Urine protein/creat ratio: 0.61

Imaging:

CXR: Coarse bl diffuse opacities

CTA Chest PE protocol: no PE, basilar and peripheral fibrosis and honeycombing w/o effusion or ptx

CT abdomen/pelvis: normal

EMG: RU & RL extremities show electrical evidence patchy myopathic process, muscle membrane irritability

MRI: T2 signal, fatty atrophy involving left quadriceps muscle

PET CT: nl

Muscle Biopsy: Marked evidence vasculitis, abundant chronic inflammation, fibrinoid necrosis, focal fatty replacement, perivascular lymph (CD3, CD4 pos, CD20 +)

Endomysial inflammation

No evidence of inclusion body myositis

Dx: ANCA vasculitis mimicking GCA and myositis

Problem Representation: 87yo F w/ PMH of GCS tx w/ steroids + recent d/c tocilizumab p/w fever and distal-predominant weakness. On exam, mixed distal > proximal weakness that on EMG confirmed myopathic process. Inflammatory markers markedly elevated and hematuria was noted on UA. Muscle biopsy showed vasculitis and p-ANCA came back positive.

Teaching Points (Vale):

Fever: IMADE. In our immunosuppressed px w/ high burden of rheum dz → Infection > Autoimmune > Drugs, Endocrinopathy > Malignancy.

FUO Approach:

1st pass: ESR, CBC, LFTs, BCx, UCx. Full-body imaging.

2nd pass: Echocardi, hepatitis, RPR, HIV, Quant, Bartonella, Brucella, Coxiella. ANA, Cryo, ANCA.

FUO + Rheum:

Connective tissue dz: Lupus, Sjogren, Scleroderma, Dermatomyositis, Mixed connective tissue dz +/- RA.

Vasculitis: Fever w/ subacute-chronic progression.

Crystals: Gout.

Autoinflammatory dz: Adult-onset Still's, periodic-fever syndromes.

💎 **ESR>100:** In the rheum category think ANCA-associated vasculitis, crystalline-arthritis.

💎 GCA is assoc w/ polymyalgia rheumatica, which causes bursitis and synovitis + normal CK levels.

Myopathy: Proximal > distal (inclusion-body p/w asymmetric mixed weakness).

💎 T-cell large granular lymphocytic leukemia can present w/ paraneoplastic rheumatologic dz (SLE, RA, Inclusion-body myositis).

Hematuria: 1st r/o mimics (ex. Period blood) → malignancy, glomerulonephritis. ? **FUO + Myositis + hematuria:** SLE.

ILD + Vascular involvement: MPA.

GC mimics: AL amyloidosis, MM, Waldenstrom's, Lymphoma.

💎 Cryoglobulinemia is a dx that lives in the intersection between infection, autoimmunity and malignancy.