

## 04/05/24 Morning Report with @CPSolvers



"One life, so many dreams" Case Presenter: Nidhi Patel (@nidhipat19) Case Discussants: Rabih (@rabihmqeha) and Reza (@DxRxEdu)

**CC**: 20 female presented to the ED for worsening dyspnea, orthopnea, emesis.

**HPI**: Patient diagnosed with HF 3 months ago from a non-ischemic cause. She has been experiencing emesis which started earlier today, she looks exhausted, has difficulty breathing. She was given PO lasix. She has more fluid buildup over the last few weeks, and she feels very tired.

afebrile, tachy, Pro-BNP: 15,000, normal CBC & CMP, Trops normal. ESR 94 CRP 123. Left heart catch was normal, initial ECG showed atrial flutter, no conduction system disease, Echo showed 10-15% and Grade 3 diastolic dysfunction, and a thrombus. Aortic lymph node with mild uptake, cardiac MRI showed enhancement in different places concerning of infiltrative disease. Started on steroids, after leaving hospital,

she felt better and after taking steroids she felt great, and was

able to focus in school. Since then, she had atrial flutter

Previously - 3 months ago, she came in with volume overload,

PMH: Bone marrow transplant at age 10 for Sickle cell disease, Mucocutaneous graft vs host disease, Hypertension (with recurrent HTN urgency) diagnosed 2 years ago, HfrEF 10-15%

Meds: B-blocker, ACE inhibitor, PO

ablation, ESR and CRP returned to normal.

lasix, PO steroids given after HF diagnosed (stopped 1 month ago), blood thinner for thrombus in Left ventricle HF

Soc Hx: currently in school, doesn't drink alcohol

Fam Hx: sister was

no family history of

donor for transplant.

alcohol Health-Related Behaviors: Allergies: Vitals: T: 100.5 HR: 110 BP: 80/74 RR: SpO2:

Exam:

Gen: very fatigued, still responding

CV: heart sounds normal, JVD bilaterally to the ears, 2+ edema, cold on exam in proximal and distal extremities

Pulm: crackles in both lungs

Notable Labs & Imaging:

Hematology:

WBC: leukocytes 11.5, no neutrophils, no eosinophils Hgb: Plt:

Chemistry:

fibrosis.

Na: K: Cl: HCO3: BUN: Cr: glucose: Ca: Mag: AST: 47 ALT: 114 Alk-P: wnl Albumin: Trops: normal

Viral panel: negative

Blood cultures, urine culture: negative

RHC: Cardiogenic shock, started on meds (lasix), sent to CCU.

Repeat ECHO was similar to before, CTPE: negative for PE, no other systemic findings of sarcoidosis.

No changes over the next one week and half. Later started on

inotropes, possible concerns of patchy inflammation. Thought process - might eventually need a heart transplant.

Repeat FDG PET Scan showed areas of patchy inflammation from previously migrated to an alternative area. Aortic lymph node also enhanced, could not biopsy. Repeat cardiac MRI showed

Rheu - agreed it might not be cardiac sarcoid.

Endomyocardial biopsy: lymphocytic infiltration, no fibrosis.

Biopsy sent for a second opinion - showed giant cells

increased delayed enhancement in left and right ventricles, and

**Dx: Giant Cell Myocarditis** 

Problem Representation: 20 y.o F w/PMH of BM transplant for SCD, GVHD, HTN, HFrEF came to ED % worsening dyspnea, orthopnea and emesis. Patient was febrile, and in cardiogenic shock with dampened QRS complex with evidence of migrating patchy inflammation of PET Scan and lymphocytic infiltration of endomyocardium.

## Teaching Points (Ibrahim):

 $\underline{\mathsf{HF}}$  (CAD [HFrEF], valvular, HTN, toxin [alcohol, cocaine], arrhythmia [can be genetic: arrhythmogenic right ventricular dysplasia], idiopathic) hx  $\rightarrow$   $\underline{\mathsf{HFREF}}$  (obesity) vs.  $\underline{\mathsf{HFPEF}}$ 

- Many causes for non-ischemic cardiomyopathy exist, make sure to distinguish one from the other
   Rapid progression HF: (3Ts: Takotsubo, Thyroid, Thiamine).
- Rapid progression HF: (3Ts: Takotsubo, Thyroid, Thiamine peripartum, myocarditis, large ischemia

HF + troponin elevation = don't miss out **myocarditis** 

- Clue → troponin persistently elevated
- MRI w/ gadolinium → delayed enhancement, patchy (myocarditis) vs coronary artery distribution (CAD)

Echo in HF → structural (e.g. filling pressure), EF, structural dx (e.g. VSD, shunting, valvular disease = diastolic dysfunction → infiltration [intracellular [HOCM] vs extracellular])

OCMJ vs extracellularJ)

 ECG → low-voltage = think extracellular material accumulation: 1.
 Amyloid, 2. Myxedema (GAG), 3. Cancer, 4. Fibrosis → histology for dx/considering base-rate [conduction disease on previous ECG] [overlap w/ myocarditis exists]

Nausea and heart aren't mutually exclusive

Persistent shock in pt. on steroids → consider hydrocort. [adrenal insufficiency]

Restrictive cardiomyopathy w/ <u>overwhelming</u> systemic presentation → EXPOSURES [e.g.

transplant source, infections: Chagas, genetic, GVHD, meds: tacrolimus]

Associated lymphadenopathy → may point towards immune → GVHD, sarcoid, cell diff.

[including giant cells]

Lymphocytic and giant-cell myocarditis (easily missed because of patchy distribution, can happen in any age) → both present w/ lymphocytic infiltrate