

05/27/24 Morning Report with @CPSolvers

"One life, so many dreams" Case Presenter: Api Chewcharat (@Api_chew) Case Discussants: Raad Chowdhury (@raad_chowdhury)

CC: a 70 yo M BL lower extremity edema

HPI:

He started noticing PE* 1 month ago, has gained weight (10 kg) over the month.

No abdominal pain, nausea, vomiting, dysuria, hematuria.

Bubbly urine +ve, difficult to flush

Rash (MP) since 6 months ago and treated with Tacrolimus, subsided over 5 months.

PMH:

Hypertension

AML (Treated with induction Rx: Idarubicin, Cytarabine. Stem cell Transplant done 1yr ago, followed with Fludarabine, Melphalan)

Meds:

HCTZ, Tacrolimus, Bactrim omeprazole.

Vitals: T: HR:62 BP:130/70 RR: saO2: 100% Weight: 100kg

Exam:

Gen:

CV: No murmurs.

Pulm: clear, no added sounds

Abd: benign, non distended

Neuro: NFND

Extremities/skin: Edema up to his thigh. MP Rash on back and shoulders

Notable Labs & Imaging:

Hematology:

WBC: 6K Hgb: 13.5 HCT: Plt:180k (bl)

Chemistry:

Na: 139 K: 4.3Cl: 103 HCO3: 26 BUN: Cr:1.5 glucose: Ca:8.7 Mag: PO4: 8.3

AST: 20 ALT: 16 Alk-P:66 Albumin: 2.7 (2.8)

PS: no schistocyte.

Hepatitis panel: N, ANA: 1: 40 Ds DNA: N VIT D :18

TSH 8.82. FT4: 0.8

BM Biopsy: Normal.

GFR

Total chol: 284. LDL: 218

C3 120 C4 22

PLA2R: negative Phospholipase 2R: SPEP: N, no M spike K32.7 lambda 22.6, KL ratio: 1.45

Imaging:

ECHO EF: 65%. Normal ventricle. Normal Valves

CXR: clear. No consolidation. No evidence PE

Venous doppler: BL DVT in L femoral vein and common vein

USG ABD: kidneys normal. Liver normal

UA: clear. Specific gravity 1.020. No red cell. Protein 3+. 24 urine: protein:

13.5 (On FU after initiating Rx for Membranous Nephropathy: U alb: 1.8,

Total protein: 5 gm)

Kidney Biopsy: (on hep drip): LM: Double contouring GBM. Stain:

Craters, Spikes. IM: C3 deposits on papillary wall. EM: subepithelial deposits. Diffuse podocyte effacement. Anti PLA2R: negative

Dx: Membranous Nephropathy

Problem Representation: 70 M presented with BL edema, weight gain of 10 Kg *1 month. And bubbly urine. PMH: HTN, AML (Stem cell transplantation done 1 year ago on chemo) Inv showed Proteinuria, Kidney Bx:Membranous nephropathy

Teaching Points (Jia):

1. **Edema:** heart, kidney, liver, DVT-> medication, systemic review, pt population (eg. with cardiac disease-amlodipine, hemo condition -gemcitabine)

Foaming urine: not sensitive or specific in diagnosis

2. **AML-myeloid disorder:** pre-renal, intrinsic renal (infiltration condition)

Tacrolimus: AKI, ATI, TMA (Bp drops down)

3. **Post-BMT kidney injury:** timeline is helpful in ddx

Acute: catastrophic AKI (occlusion in vessels, like hepatic sinusoidal obstruction syndrome and thrombotic microangiopathy), acute engraftment syndrome

Post 100 days: ->**Intrinsic:** ATN, interstitial disease (bactrim, penicillin), glomerular disease, BK nephropathy; ->**GN:** membranous nephropathy, minimal change disease, thrombotic microangiopathy; ->**Myeloid specific etiology:** lysozyme induced nephrology: lymphoma- 60-70% involve kidney, especially CLL

4. **GVHD:** no renal criteria; the timeline relationship of renal symptom and appearance in other systems is important for renal GVHD diagnosis.

5. **Renal TMA:** may not have systemic signs like schistocyte

6. **Complements in nephrology:** narrow down the ddx;

Classical pathway (C4+C3 low): lupus;

Alternative pathway (C3 low): PNH, TMA;

CAMEL mnemonic: C-cryo, A-atheroembolic, M-myeloma (C3 nephropathy), E-endocarditis, L-lupus, Infection GN

7. **Biopsy interpretation:** Double contour: GBM changes can be seen in TMA; membranous nephropathy associated with hematopoietic cells transplantation: **FAT1-Ab against podocytes**

8. **Dense deposit disease:** alternative pathway disorder with low complement; MN and MCD: autoimmune disorder.