



05/27/24 Morning Report with @CPSolvers

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



<p>CC: a 70 yo M BL lower extremity edema</p> <p>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</p> <p>Rash (MP) since 6 months ago and treated with Tacro, subsided over 5 months.</p>		<p>Vitals: T: HR:62 BP:130/70 RR: saO2: 100% Weight: 100kg</p> <p>Exam:</p> <p>Gen:</p> <p>CV: No murmurs.</p> <p>Pulm: clear, no added sounds</p> <p>Abd: benign, non distended</p> <p>Neuro: NFND</p> <p>Extremities/skin: Edema up to his thigh. MP Rash on back and shoulders</p>	<p>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</p>
<p>PMH: Hypertension AML (Treated with induction Rx: Idarubicin, Cytarabine. Stem cell Transplant done 1yr ago, followed with Fludarabine, Melphalan</p> <p>Meds: HCTZ , Tacrolimus, Bactrim omeprazole.</p>	<p>Fam Hx: Nothing significant</p> <p>Soc Hx: No alcohol, former smoker, quit 20 years back</p> <p>Health-Related Behaviors:</p> <p>Allergies: None</p>	<p>Notable Labs & Imaging:</p> <p>Hematology: WBC: 6K Hgb: 13.5 HCT: Plt:180k (bl)</p> <p>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</p> <p>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.0.08. 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)</p> <p>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</p> <p>Dx: Membranous Nephropathy</p>	<p>Teaching Points (Jia):</p> <ol style="list-style-type: none">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 diagnosisAML-myeloid disorder: pre-renal, intrinsic renal (infiltration condition) Tacrolimus: AKI, ATI, TMA (Bp drops down)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 CLLGVHD: no renal criteria; the timeline relationship of renal symptom and appearance in other systems is important for renal GVHD diagnosis.Renal TMA: may not have systemic signs like schistocyteComplements 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 GNBiopsy interpretation: Double contour: GBM changes can be seen in TMA; membranous nephropathy associated with hematopoietic cells transplantation: FAT1-Ab against podocytesDense deposit disease: alternative pathway disorder with low complement; MN and MCD: autoimmune disorder.