



# 6/19/20 Morning Report with @CPSolvers



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<p><b>CC:</b> chest pain</p> <p><b>HPI:</b> 29 M presented with chest pain for 2 days, describes it as sharp pain, anterior chest, does not radiate. No exacerbating or relieving factors. Some episodes of tachycardia, SOB, x1 episode syncope 1 day prior  <b>ROS:</b> intermittent fevers, chills, drenching night sweats, loss of appetite, weight loss 4-5 kg, and swollen LN in neck over 3 mo, denies sore throat/oral ulcers, intermittent diarrhea, denies GU Sx, denies abdominal pain</p>	<p><b>Vitals:</b> T: 37 HR: 118 BP: 133/86 RR: 18 SpO<sub>2</sub>: 99</p> <p><b>Exam:</b>  <b>Gen:</b> NAD, alert  <b>Lymph:</b> palpable cervical and inguinal LN  <b>CV:</b> no PAD, no m/r/g  <b>Pulm:</b> CABL  <b>Abd:</b> soft, non tender, non distended, no organomegaly  <b>Neuro:</b> nl  <b>Extremities/Skin:</b> seborrheic dermatitis on face, hyperpigmented patches over chest/back (new)</p>	<p><b>Problem Representation:</b> 29yo man presenting with several months of constitutional symptoms, weight loss, chest pain, generalized LAD, and new skin findings with anemia and protein gap. The patient was ultimately diagnosed with HIV-associated, HHV8-associated, multicentric Castleman's disease</p>
<p><b>PMH:</b> none</p> <p><b>Meds:</b></p>	<p><b>Notable Labs &amp; Imaging:</b>  <b>Hematology:</b>  WBC: 9.1 (46% seg neut, 37% lymph, 17% mono), Hgb: 8.3 Plt: 350  Peripheral smear- rare smudge cells, slight anisocytosis, rare monocytes, rare stomatocytes</p> <p><b>Chemistry:</b>  Na: 133 K: 5 Cl: 99 CO<sub>2</sub>: BUN: 10 Cr: 1 glucose: 121  Ca: Phos: Mag:  AST: 22 ALT: 15 Alk-P: T. Bili: 0.6 Albumin: 2.4 TP: 8.7  D-dimer: 43.44 Trop: &lt;0.01  HIV ab - pos ; HIV 1 ab - pos ; HIV 2 ab- non reactive  Flow: no evidence of a lymphoproliferative disorder or leukemia</p> <p><b>Imaging:</b>  EKG: sinus tachycardia with no ST changes  CXR: no acute infiltrates, normal heart size  CTA chest: no evidence of PE; lungs clear; no infiltrates. B/L mediastinal axillary/hilar LAD  <b>Core Bx of R inguinal LN-</b> atrophic germinal centers + strong plasma cells  <b>Special stains:</b> CD 23 w/ residual atretic germinal centers; BCL-2 outlining atretic germinal centers; CD 34- strong vascular background, CD 138- numerous plasma cells; Kappa/ Lambda light chains- numerous plasma cells  HHV 8- focally positive in spindled areas</p> <p><b>Dx:</b> HIV and HHV-8 associated multicentric Castleman's disease; outpatient Ritux/ HIV tx</p>	<p><b>Teaching Points (Anand):</b>  <b>Chest Pain:</b> outside-in approach → skin, musculoskeletal, lungs, heart, GI, and other  <b>Lymphadenopathy w/fevers, sweats (code as inflammation):</b> infection, malignancy, or autoimmune buckets are the most common; also remember drugs, DVTs, and endocrinopathies  <b>Syphilis pearl:</b> untreated syphilis can become latent, and then progress to secondary syphilis  <b>Inflammation workup:</b> are there localized sites we can hone in on from the physical exam?  <b>Lymph node red flags:</b> firm/matted, non-tender; Generalized LAD → higher concern for malignancy  <b>Anemia pearls:</b> anemia in a young patient → hard to attribute it to a chronic inflammatory condition; is there blood loss? Is there a hemolytic process? Is there a primary bone marrow process? Smudge cells on a smear → think CLL or mononucleosis  <b>Protein Gap:</b> There is excess paraprotein (from a polyclonal process like inflammation or a monoclonal process like myeloma)  <b>HIV pearls:</b> HIV+ patients are not just at risk of concurrent infection, but also at risk of malignancy (lymphoproliferative diseases (lymphoma, Castleman's), kaposi sarcoma)  <b>When plasma cells and B-lymphocytes mix</b> → think lymphoplasmacytic lymphoma or Castleman's disease  <b>Castleman's Disease pearls:</b> can be unicentric (one lymph node region) or multicentric (generalized LAD, associated with HIV and HHV8)</p>