

01/19/24 Morning Report with @CPSolvers



"One life, so many dreams" Case Presenter: Yale IM Residency Case Discussants: Rabih (@rabihmqeha) and Reza (@DxRxEdu)

CC: A 26 yo M presents with persistent right sided headache since last week.

HPI: 1 week of right sided headache and left sided weakness intermittently. Felt well prior to presentation. Headache: No prodromal symptoms, No triggers, Resolved by itself, Weakness not always associated with the headache, No numbness or tingling, No additional factors aggravating or relieving headache were noted

Clinical course: Worsening of HA, new L sided weakness, L facial droop, slurred speech, L arm plegia & multiple stroke codes

ROS: Otherwise negative.

PMH: Splenectomy @ 3yrs for unknown reasons, TB lymphadenitis, recent URTI

Meds: None for

now. In the

past: RIPE for

TB, Augmentin

Fam Hx: None Soc Hx: Moved from India in 2019, software

engineer

Health-Relate d Behaviors:

Allergies: NA

Vitals: T: afeb HR: nl BP:nl RR: nl

HEENT: 2cm b/l anterior cervical lymphadenopathy
CV: nl, Pulm: nl, Abd: soft, non tender, non distended, scar

from splenectomy

Neuro: 5/5 sensation, normal cranial nerve exam, 5/5 strength in UE & LE

Extremities/skin: 2 cm bl firm inguinal lymphadenopathy

Notable Labs & Imaging:

Hematology: WBC: nl with nl differential Hgb:nl Plt:703k

Chemistry: Na: nl K: nl Cl: nl HCO3: nl BUN: nl Cr: nl glucose: nl Ca: nl, LFT: nl, PT: nl, PTT: 56, CRP: 7.3

Imaging:

CT w/o contrast head: nl, CT Sinus: Acute on chronic sinusitis CXR:abnormal-> CT Chest: Mass (4*5 cm) left lower lobe of lung with calcification CT AP: Inguinal lymphadenopathy

Imaging after worsening of course:

inflammation & epithelioid cell

TTE: nl, CTA: Large right ICA thrombus

MRI brain: acute infarcts over the left cerebral hemisphere with enlarged parotid glands

AFB: Neg 3x, TB PCR:neg, HIV & Histo: neg, C3 & C4: nl, ACE level:nl, ANA: 1:320 & 1:640 homogenous, Anti SSA/B: nl, Anti Smith: nl, Anti ds DNA: nl, Anti cardiolipin: neg, Anti beta2 GP:nl

Lupus anticoagulant: Positive -> Prior records from 2 yrs ago: Lupus anticoagulant and ANA found to be +ve Bronchoscopy: Neg for TB, **Biopsy** (lung mass): Acute

Dx: Antiphospholipid syndrome and Rosai Dorfman disease

Problem Representation: 26 yr old M patient presents with persistent right sided headache and intermittent left sided weakness was found to have progressive lymphadenopathy, a calcified lung mass and positive anticoagulant antibodies

Teaching Points (Anmolpreet):

Il Headache: associated red flag: intermittent weakness/focal neurologic deficit makes us focus on secondary headache more.: Approach: Primary headache syndrome (migraine, tension, cluster) OR Secondary headache syndrome (SNOOP(presence of red flag signs)}- causes include: intrinsic(meninges) or extrinsic (eye, sinus, dental, neck) or systemic(inflammatory process)

Important to note is diurnal variation, headache worse in morning can indicate high ICP(concern for SOL)

II] Weakness:- Headache helps us narrow the differential and localise the cause of weakness to brain.(left sided weakness-right sided headache)

Intermittent weakness for a week: subacute CNS syndrome almost always indicates a space occupying lesions, CT imaging a priority: suspect (a) Demyelinating ds, (b) venous sinus thrombosis

III] Splenectomy indicates immunocompromised state, hence broadens our differentials to consider infection from encapsulated organisms; imp to know immunisation history;

B/L parotid swelling: unlikely to be cancer and stone

IV Prior h/o TB: makes us think about recurrence or tuberculoma

<u>VI</u> Localised cervical LAD: <u>Anterior</u>: closer to oropharynx; 2/2 local oropharyngeal process/ infection; Posterior: most likely systemic:- eg Infectious Mononucleosis

VII Diffuse LAD: 2 important tests: HIV and Syphilis

VIII LAD: we have to consider infection and cancer. Absence of fever and significant weight loss in this case with disproportionate LAD is something we need to evaluate.

 $\underline{\textbf{VIII]}} \textbf{Thrombocytosis:} \ \underline{\textbf{Reactive}} : \textbf{to inflammation;} \ \textbf{not a primary hematologic issue;} \ \textbf{accompanying anemia of chronic ds } \& \ \textbf{increased inflammatory markers} \ \textbf{are expected}$

<u>IX1</u> elevated PTT: if can be corrected with mixing study: def of one of the factors in the pathway

XI MR Venography to understand if CNS is involved, which cannot be ruled out with CT without contrast, imp because ot came with neuro sx

XI] Calcifications: indicate granulomatous diseases like TB

XIII Mass in CT Chest: could have been lung cancer which has a predilection to go to brain and Lymphoma., GPA, Sarcoidosis

XIII] In situ thrombosis: atherosclerosis, (blood vessel wall)vasculitis, dissection, hypercoagulable

condition.(APLS- idiopathic, 2/2 lupus- APLS may precede any symptoms of SLE)

XIV] Rosai-Dorfman Disease: sinus histiocytosis with massive LAD