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::step2CK Aug24

Renal system

1. Young male child, Cv suggestive of VHL syndrome—what to screen for? – retinal hemangioblastoma
2. 12-13 yrs male patient with flank pain and hematuria, CT IVA done showing stone in renal pelvis. How to prevent? Increase fluid intake >2 l/day
3. Patient taking cephalosporin group drug, present with `wbc cast—to dx AIN
4. CV of orthostatic proteinuria in children--- reassurance
5. Spinal stenosis dx in cv—laminectomy done—on postoperative period, patient develop incontinence , neurological examination was normal, anal sphincter tone is normal, PVRV= 20 ml (which is normal), cause for this symptoms asked I guess. Option I forgot
6. Case of HTN, prescribed linsopril, rise of serum creatinine on follow up- dx RAS
7. Normal adolescent person goes for hiking. He used to have sedentary life style, now comes with severe pain in thigh. Serum CK level increased in > 1000. What to do next? Advise to continue exercise
8. Sudden respiratory distress in child, chest xray normal, NBS- bronchoscopy

Respiratory system

1. Updown arrow in khyposis
2. Cv of laryngomalacia in infant- reassure
3. CV of recurrent laryngeal papillomatosis with HPV6 positive in mother- MX: resection
4. Adolescent male with neck swelling with signs of inflammation, undergone supportive management, later developed chest pain and SOB, CT chest done showed pneumomediastinum with fluid collection. After starting antibiotics, NBS- debridement
5. Lung abscess to diagnose, xray shows airfluid level
6. Asbestosis to be diagnose, x- ray given
7. History suggestive of silicosis—what to look for? Pleura, carcinoma
8. Obese female, liposuction done--- develop respiratory distress, petechie---- fat embolism
9. MVA—respiratory distress suggestive of tension pneumothorax—NBS: tube thoracostomy
10. CV of tension pneumothorax, what will be the physiological change... dec right ventricle venous return
11. History of recurrent pneumonia, HRCT showing bronchiectasis, sweat chloride test is normal, no history of chronic diarrhea, nasal polyp... mx- IVIG thinking of CVID , no antibiotics in option I guess (primary ciliary dyskinesia can be ruled out from cv)
12. Female presented with leg pain, sudden onset tachypnea suggestive of PE, CT angiogram done, reports awaited.. nbs: LMWH (all form of anticoagulant given)

Cardiovascular system

1. HTN related question—I forgot

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2. Subacute onset of fever, abdominal pain, abdominal distension with firm mass palpable over left lower quadrant of abdomen... what to look for? Splenic abscess
3. Cv of cholesterol emboli
4. ASCVD risk below 7, LDL high but not above 190, no history of DM, what next--- exercise
5. Ecg of inferior wall MI, Pericarditis
6. Heart sound: AS, VSD, PDA
7. Male patient with respiratory distress, hemodynamically unstable, normal JVP mention, lung auscultation normal, however, cath lab values were given showing increased with equal diastolic pressure in RA, RV--- next best step: pericardiocentesis
8. Patient is in shock, no history to think for adrenal insufficiency, fluid bolus given, dopa/ dobuta stated @ 10mcg/kg/min, still BP in lower side... NBS: increase dose
9. Some surgical procedure going on in OT, blood transfusion done--- suddenly develop respiratory distress... BP normal (r/o TACO), ABG not given, NBS? ...
10. Cv describing orthostatic hypotension: ACE inhibitors

Endocrine:

1. Operated for pituitary adenoma, presented with polyuria, sp. Gravity of urine is decrease--- DI to diagnosed
2. Multiple question on hypothyroidism and hyperthyroidism
3. CV of conns disease showing unilateral increased Aldosterone/ renin in adrenal venous sampling: adenoma
4. DKA CV: nbs: NS
5. 42 yr male, everything normal, what to screen for? FBS
6. Carcinoid tumor origin, diagnosed in biopsy: small intestine
7. Cv of hypoglycemia with slightly raised c-peptide: insulinoma
8. Patient in shock, managed with fluid.. NBS: add adrenaline
9. CCf with EF < 40%, HBA1C 8.1%, NBS: add STLG1

Immunology:

1. Recurrent staph aureus infection: oxidative burst
2. CVID to diagnose
3. BT done, feature of volume overload with increased BP: TACO
4. Ceftriaxone given, developed rash, fever, arthralgia--- serum like sickness
5. Pregnancy: Tdap to be given
6. 17 yrs old male: meningococcal vaccine

Neurology:

1. Developmental milestone related 2 question—easy one
2. CSF picture suggestive of TB to diagnose
3. Constitutional macrocephaly: reassure
4. Restless leg syndrome, what to look further: serum ferritin

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5. Essential tremor: primidone
6. Case of non exertional heat exhaustion, Temp around 104F, MX: supportive
7. ALS to diagnose
8. Bell's palsy , MX: supportive
9. PCKD CV given, future risk: SAH
10. Female around 32-34 yrs with feature of normal pressure hydrocephalus, NBS: MRI brain
11. REM sleep disorder with parkinsonian feature, NBS: MRI brain, polysomnography
12. CT scan showing orbital fracture (medial side): problem??
13. CV of meniere's disease
14. CMV retinitis funduscopy picture

Others

1. Question about pt with bloody diarrhoea describing right lower quadrant pain and on colonoscopy you see cobblestone appearance in ascending colon as well as terminal ileum. Asking for dx – chrons disease
2. In second question asking for treatment- steroids
3. 3- old pt with abdominal pain, last stool was 5 days ago, picture of sigmoid volvulus given, NBS- proctosigmoidoscopy
4. Old pt with some abdominal pain, hemodynamically stable, no fever or leukocytosis, some vague abdominal xray is given, looks like small bowel obstruction NBS- NG tube
5. 5. Some middle aged pt with history of hernia repair 10 year ago now have mass in right lower quadrant, xray given looks normal to me, asking for dx
6. Pt with 32 weeks of pregnancy comes to you with cervix dilated and effaced, intact membrane -preterm labor
7. Pt with some terminal disease and doctor discussed all the treatment option but pt dont want any treatment asking for the principle- autonomy
8. pt with some disease and doctor refuse some procedure which will harm the patient asking the principal- nonmaleficence
9. Biostats forest blot shown and asking for relative risk question
10. HIV pt with CD4 count 150, against which organism you will give prophylaxis- P.jerovci
11. Pt went on hiking develop itchy rash on his neck that spread into chest and trunk, picture of neck rash given, also pt completed 10 day course of TMP-SMX for UTI, asking for dx- Sunburn or allergic reaction
12. Pt with gynaecomastia and small testis asking NBS- all option were hormones
13. Pt with some disease hospitalized and started on antibiotics now have diarrhoea asking the cause - C diff infection
14. Picture of tenia capitis asking for TX- oral fluconazole
15. Pt with some cat bite NBS- give antibiotics
16. Child with vesicles on soft palate and throat NBS- Reassure
17. GVHD biopsy

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18. Pt with lymphoma undergoing chemotherapy what will you find in labs - increase phosphorus
19. Couple come for pregrancy counselling, both of them were more than 35y year old, what disorder will you counsel them - chromosomal trisomy
20. pt with migraine with aura and history of heavy menstrual bleeding for contraceptive what will you give - levonogestral IUD
21. Pregnant Pt with history of MDD discontinue SSRI NBS- Start SSRI again safe during pregnancy
22. 16y old with personality changes, irritability, picture of eye given asking for tx- penicillamine
23. Pt with mass in abdomen and question stem describing hemihypertrophy of calves asking for dx- nephroblastoma
24. Child with systemic system like fever malaise and a bone mass asking for dx- Ewing sarcoma, because ewing sarcoma
25. Child with bone mass asking for dx- osteosarcoma.



1. c. difficile- vancomycin

Tinea capitis is a fungal infection of the scalp that causes pruritic patchy, fine, white scales that may resemble SD. However, it does not involve the eyebrows or nasolabial folds and is uncommon in the first year.

Tinea capitis	
Epidemiology	• Most common in African American children • Transmission via direct contact or from fomite (eg, shared combs)
Clinical features	• Scaly, erythematous patch with hair loss on scalp

2. Tinea capitis

Clinical features	• Scaly, erythematous patch with hair loss on scalp • ± Black dots in affected area • ± Tender lymphadenopathy
Management	• Oral griseofulvin or terbinafine

TC is often diagnosed clinically but can be confirmed with potassium hydroxide (KOH) examination of hair stubs. Oral griseofulvin or terbinafine is preferred in most patients. As dermatophyte carriers can be asymptomatic, many experts recommend that household contacts be treated with selenium sulfide or ketoconazole shampoo.

3. CML imatinib

28) Diagnose CML(granulocytosis too much WBC) and histology is given - BCR-ABL / t(9;22)

Diagnosis: BCR-ABL / t(9;22)

Management: Imatinib (tyrosine kinase inhibitor)

Other notes:

- CBC findings: Eosinophilia
- Presentation: 10-15 yr old with fatigue + weight loss + splenomegaly
- Think of "t(9;22)" for "t(9;22)" - presents in middle-aged etc.!
- T(9;22) is BCR-ABL fusion gene
- BCR-ABL fusion gene is the cause of CML
- Imatinib is the treatment

4. Anchoring and availability bias

5. Root cause analysis

6. Hemothorax- thoracotomy

Penetrating trauma accompanied by shock (eg, severe hypotension) is attributed to hemorrhage until proven otherwise. Although **tube thoracostomy** is often sufficient to manage hemothorax, some patients (up to 15%) require **emergent thoracotomy** for extreme bleeding, including those with:

- Initial bloody output >1,500 mL (>20 mL/kg)
- Persistent hemorrhage: >200 mL/hr for >2 hours, or continuous need for blood transfusion to maintain hemodynamic stability

7. Splenectomy done- which vaccine to give pneumococcal and h influenza both were in option

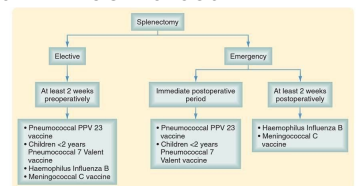


FIGURE 57-7 Splenectomy immunoprophylaxis flow chart. (From Harji DP, Jaunoo SS, Misty P, Nesaigkar PN: Immunoprophylaxis in asplenic patients. Int J Surg 7:421-423, 2009.)

8. Another splenectomy case- meningococcal vaccine

9. Asthma treatment

10. Interstitial cystitis question treatment

Apnea of prematurity caffeine dini

Avoid caffeine..

11. Ctg given - looked normal nbe in management

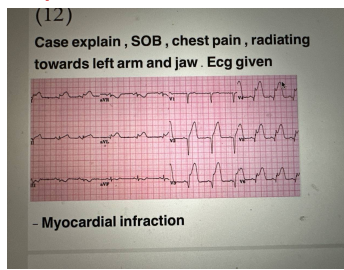
VEAL CHOP MINE Nursing Mnemonic

Find Pattern (VEAL)	Cause (CHOP)	Management (MINE)
V Variable deceleration	C Cord compression	M Maternal repositioning
E Early deceleration	H Head compression	I Identify labor progress
A Acceleration	O Occlus	N No interventions
L Late deceleration	P Placental insufficiency	E Early interventions

Head compression - early deceleration
Variable deceleration - cord compression
Late deceleration - uteroplacenta insufficiency

Paila tauko niskinxa, ani cord ani placenta , early , variable and late

12. Ecg on MI and Wpw



13. Up and down arrows on copd and xiphosis fvc **recoil** elasticity and something more

Dlco- dec

14. Had to diagnose crohns and then treatment asked (sequential question)

15. Case of pertusis treatment **azithro as t/t and prophylaxis**

Pertussis postexposure prophylaxis	
Indications (regardless of vaccination history)	<ul style="list-style-type: none"> Close contact (eg, household members, direct contact with secretions) with symptomatic patient High-risk patients, even with limited exposure (eg, pregnant, infant, immunodeficient)
Treatment	<ul style="list-style-type: none"> Age <1 month: azithromycin Age ≥1 month: azithromycin, clarithromycin, or erythromycin

16. Ewing sarcoma

Ewing sarcoma	
Epidemiology	<ul style="list-style-type: none"> Second most common pediatric bone malignancy (after osteosarcoma) Peak incidence age 10-15
Clinical features	<ul style="list-style-type: none"> Chronic, localized pain & swelling ± soft tissue mass Involves long bone diaphyses, pelvis, axial skeleton ± Systemic findings (fever, leukocytosis, ↑ inflammatory markers) Early metastases common (eg lung, bone)
Evaluation	<ul style="list-style-type: none"> X-ray: central lytic lesion with moth-eaten appearance, lamellated periosteal reaction ("onion skinning") Biopsy: small, uniform, round blue cells

Abdominal ultrasound every 3 months until 8 years of age
Alpha-fetoprotein levels every 3 months until 4 years of age

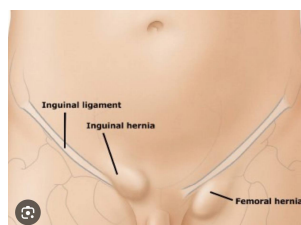
17. Rash in diaper region- did not look like Candida to me more like irritant contact dermatitis treatment asked

18. Asbestosis question what is the patient more likely at risk for it's been 10-20 years since he was exposed pleural plaques or intraparenchymal tumour

19. Lichen sclerosis is child, she had stair ia and the examination findings were describing i

adolescents ma sidhai clobetasol, adults ma suruma biopsy then clobetasol

20. Femoral hernia



21. Empyema nbs

Empyema like features = low ph , low glucose , inc LDH , ALSO SEEN IN Rheumatic pleurisy

22. OCD treatment

SIGECAP

23. MDD to diagnose

24. Social anxiety vs phobia

Social anxiety disorder (social phobia)	
Diagnosis	<ul style="list-style-type: none"> Marked anxiety about 2+ social situations for 6 months Fear of scrutiny by others, humiliation, embarrassment Social situations avoided or endured with intense distress Marked impairment (social, academic, occupational) Subtype specifier: performance only
Treatment	<ul style="list-style-type: none"> SSRI/SNRI Cognitive-behavioral therapy Beta blocker or benzodiazepine for performance-only subtype

SNRI = serotonin-norepinephrine reuptake inhibitor; SSRI = selective serotonin reuptake inhibitor

A 22-year-old woman comes to student health services because of increasing **anxiety** about being called on in class for the past 6 weeks. Yesterday, she was so worried about being called on that she was shaking and spilled her coffee. Today, she was so frightened that she was unable to attend class. She says that she has almost found it difficult to be the center of attention and has always avoided situations in which she would have to perform or speak in public. Physical examination shows no abnormalities. On mental status examination, she is cooperative and well dressed. She is anxious and fearful, her thought processes is organized. There is no evidence of suicidal ideation. Which of the following is the most appropriate next step in management?

A. Cognitive behavioral psychotherapy
 B. Imipramine psychotherapy
 C. Buspirone therapy
 D. Fluoxetine therapy

Correct answer: A
 I% answered correctly

Explanation:
 Correct Answer: A

Social **(anxiety)** disorder refers to an excessive fear of scrutiny, embarrassment, or rejection in social and/or **(performance)** situations, leading to significant distress and/or impaired functioning (eg, class or work avoidance). Patients with social **(anxiety)** disorder commonly exhibit **anxiety and performance (anxiety)**. **Performance (anxiety)** refers to the fear of negative evaluation in situations such as public speaking, job, being called on in class, test-taking, and sex. Cognitive behavioral therapy (CBT) and selective serotonin reuptake inhibitors are the first-line treatments for social **(anxiety)** disorder. Some patients with social **(anxiety)** disorder experience **anxiety** that is specific to **(performance)** situations only. In the **(performance)** only subtype of social **(anxiety)**, CBT is recommended as the first-line treatment, though as-needed **(anxiety)** medications (eg, propranolol, benzodiazepines) are frequently used as well. CBT for social **(anxiety)** disorder focuses on gradual exposure to feared social situations.

25. Pericarditis case

26. AS heart sound I think tried to pick it from the scenario

27. Pic of horse shoe kidney- ureteropelvic junction obstruction

- Horseshoe kidney:** Fusion of the left and right inferior renal poles
 - Normal ascent interrupted as fused kidney gets caught on the inferior mesenteric artery (IMA).
 - Increased incidence in patients with chromosomal aneuploidy (e.g., trisomies 13, 18, 21, and Turner syndrome)
 - Usually asymptomatic; typically diagnosed incidentally on abdominal imaging for unrelated conditions
 - Rarely requires treatment
 - Increased risk of renal stones, ureteropelvic junction obstruction, hydronephrosis, infections, and renal cancer

Horseshoe kidney

A congenital disorder characterized by the fusion of both kidneys into a horseshoe shape. Kidney function in itself is not impaired, but the abnormal anatomy increases the risk of ureteropelvic junction obstruction and kidney stones with subsequent inflammation.

Horseshoe kidney Horseshoe kidney

28. A lot of MVA questions

29. Pic of leg with Bullae and happened after he injured himself with a nail or something c perfringens

Necrotizing fasciitis (gluteal region)

Necrotizing fasciitis

Daptomycin/Linezolid/Vancomycin
 +
 Carbapenem/Tazo-pip/FQ-Metron/Xone-Metron
 +
 Clindamycin (if linezolid not taken on 1)

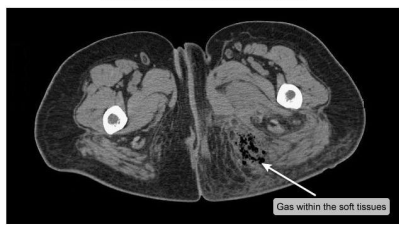
Cellulitis--

Mild - Cephalexin
 Penicillin VK
 Dicloxacillin
 If Penicillin allergic: Clindamycin

Mod - Cefazolin
 Ceftriaxone
 Narciilin/Oxacillin If Penicillin allergic:
 Clindamycin

Severe - Vancomycin +/- Piperacillin/Tazobactam
 Meropenem/Imipenem
 Linezolid

**Vanco+
 meropenem
 thyo last
 pool ma**



30. Got a question about floppy baby and decrease tone was confused between **spinal muscular atrophy** and muscular dystrophy

Spinal Muscular Atrophy (SMA)

- Pathogenesis:**
 - Impaired of cholinesterase synthesis (e.g. **SMN2 gene mutation**)
 - Spores release the immature germinative tract & produce toxin
 - Toxin inhibits presynaptic acetylcholine release
- Clinical presentation:**
 - Age: **18 months**
 - Progressive muscle weakness**
 - Respiratory failure**
 - Intellectual disability**
- Diagnosis:**
 - Clinical, supported by **EMG/Genetic testing**
 - Confirmation by **Genetic testing**
- Treatment:**
 - SMN2 gene therapy**

Myotonic Dystrophy (DM)

- Pathogenesis:**
 - Expansion of CTG repeats in the 3' UTR of the DMPK gene
 - Expansion of CTG repeats in the 5' UTR of the ZNF9 gene
- Clinical presentation:**
 - Age: **15-20 years**
 - Myotonia**
 - Proximal muscle weakness**
 - Cardiac conduction system abnormalities**
 - Connective tissue abnormalities**
 - Intellectual disability**
- Diagnosis:**
 - Clinical, supported by **EMG/Genetic testing**
 - Confirmation by **Genetic testing**
- Treatment:**
 - Cardiac monitoring**
 - Respiratory support**

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 - Confirmation by **Genetic testing**
- Treatment:**
 - Cardiac monitoring**
 - Respiratory support**

31. Female comes for prenatal counselling, on carbamazepine nbs increase dose of **folic acid**

Neural tube defects	
Types	<ul style="list-style-type: none"> Anencephaly Encephalocele Spina bifida, myelomeningocele
Risk factors	<ul style="list-style-type: none"> Low folic acid intake Methotrexate, antiepileptics Diabetes mellitus Prior pregnancy with neural tube defect
Prenatal screening	<ul style="list-style-type: none"> 2nd trimester ultrasound Maternal serum alpha-fetoprotein
Prevention	<ul style="list-style-type: none"> Average risk: 0.4 mg folic acid daily High risk: 4 mg folic acid daily

32. She has increased dose nbs usg, karyotype or chronic villous sampling

Aba usg garni

Test	Time point	Prenatal testing	
		Advantages	Disadvantages
First trimester combined test	11-13	Early screening	Not diagnostic
Cell-free DNA	10-13	High accuracy & specificity for aneuploidy	Not diagnostic
Chorionic villus sampling	10-13	Definitive karyotype diagnosis	Invasive, risk of spontaneous abortion
Second trimester quadruple screen	15-20	Screening for neural tube defects & aneuploidy	Not diagnostic
Amniocentesis	15-20	Definitive karyotype diagnosis	Invasive, risk of miscarriage, foetal loss & pregnancy loss
Second trimester ultrasound	15-20	Structural fetal growth, available fetal anatomy, maternal placental position	Change quality of amniotic fluid, foetal loss are of uncertain significance

33. S1Q3T3 on ecg nbs

34. Blood on urethral meatus nbs retrograde urethrography

35. Intention tremor propranolol and primidone both were in option

**Incase of Copd and asthma - premidone
 Natra propranolol**

36. Femoral neck fracture repaired has buttock pain limited range of motion due to pain osteonecrosis or osteoarthritis

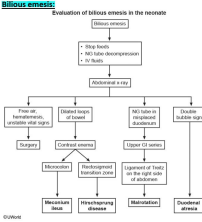
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37. Chagas' disease no symptoms nbs

**Trypanosoma cruzi - Megalo organs
 Large kinetoplast ,ecg or echo**

**Trypanosoma bruze - African sleeping
 sickness
 Small kinetoplast**

Chagas disease	
Epidemiology	<ul style="list-style-type: none"> Caused by protozoan Trypanosoma cruzi Most common in Mexico, Central & South America
Cardiac manifestations	<ul style="list-style-type: none"> Biventricular heart failure (right > left) with cardiomegaly Ventricular apical aneurysm Mural thrombosis with embolic complications Fibrosis leading to conduction abnormalities (heart block & ventricular tachycardia)
Gastrointestinal manifestations	<ul style="list-style-type: none"> Progressive dilation of esophagus & colon



i think in NBME 11 mid gud volvulus huncha CV but Xray pani gareko hunna , i chose Abd xray first but ans was UGI series...

Funda , NBS — XRay abdomen , tya chai diagnostic sodheko raicha so altho X ray comes before UGI series , diagnosis lai UGI series jane raicha

Pyloric stenosis - USG
Intususception - USG
Necrotising Enterocolitis - X ray
Meckels - 99Tc
Hirschsprung - Contrast enema (Diagnostic - Biopsy)
Diverticulitis - CT Abd
Malrotation - GI series
Volvulus (adult) - C

38. X-ray what looked like malrotation I marked upper GI series

39. A couple of questions on thyroid nodules and nbs

40. Kaposi sarcoma

41. Aspiration pneumonia treatment

aspiration -ampi sulbactam, clinda amoxi clav

Aspiration pneumonia

Rx

Out patient: Amoxicillin+ Clavulanic acid (Clavam)

In patient: Ampicillin+ Sulbactam (PQ)

ICU: Piperacillin+ Tazobactam (Durataz)

Bacterial aspiration pneumonia	
Pathophysiology	<ul style="list-style-type: none"> Oropharyngeal/gastric microbes aspirated into lungs → host defenses overwhelmed due to large inoculum size
Major risk factors	<ul style="list-style-type: none"> Reduced consciousness (eg, alcoholism) Dysphagia (eg, neuromuscular disorders) Impaired gastric closure (eg, distal duodenal obstruction) Florid dental hygiene Poor dental hygiene
Clinical features	<ul style="list-style-type: none"> Fever & cough & foul-smelling sputum Infiltrate in dependent portions of lung Abscess & empyema pathogens on sputum studies
Management	<ul style="list-style-type: none"> No empyema or lung abscess: treat for community-acquired pneumonia Empyema or lung abscess present: extend coverage to include anaerobes (eg, ampicillin/sulbactam)

42. Case on ttp nbs plasmapheresis



Thrombotic thrombocytopenic purpura	
Pathophysiology	<ul style="list-style-type: none"> ADAMTS13 level → uncoupled vWF multimers → platelet trapping & activation Acquired (autoantibody) or hereditary
Clinical features	<ul style="list-style-type: none"> Hemolytic anemia (↓LDH, ↑hepatoptin) with schistocytes Thrombocytopenia (↓bleeding time, normal PTT) Sometimes with: <ul style="list-style-type: none"> Renal failure Neurologic manifestations Fever
Management	<ul style="list-style-type: none"> Plasma exchange Glucocorticoids Rituximab

LDH = lactate dehydrogenase, vWF = von Willebrand factor.

plasma pheresis ...gluco... rituximab

Aspiration- Amp/ Sulbactam.. ma yesari yaad garchu

43. Pic of Wilson disease nbs I think the option that made sense in my question of urinary copper

Wilson disease	
Pathogenesis	<ul style="list-style-type: none"> Autosomal recessive mutation of ATP7B → hepatic copper accumulation → leak from damaged hepatocytes → deposits in tissues (eg, basal ganglia, cornea)
Clinical findings	<ul style="list-style-type: none"> Hepatic: acute liver failure, chronic hepatitis, cirrhosis Neurologic: parkinsonism, gait disturbance, dystonia Psychiatric: depression, personality changes, psychosis
Diagnosis	<ul style="list-style-type: none"> ↑Ceruloplasmin & ↑urinary copper excretion Kayser-Fleischer rings on slit-lamp examination ↑Copper content on liver biopsy
Treatment	<ul style="list-style-type: none"> Chelators (eg, D-penicillamine, trientine) Zinc (statensin with copper absorption)

cerulo dec , urinary copper badne

44. Digeorge syndrome asking what will be less hypocalcemia

Di George
If catch22 and hypoCa already mentioned in vignette then go with - HypoMg

45. Pic of a lesion on hand I guess with histology pic I marked kaposi because I could rule out cmv and ebv I may be wrong

46. Very obese female wants to get pregnant nbs weight loss

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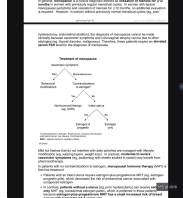
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- All patients: Encourage lifestyle interventions and address modifiable factors.
- Adjuvant therapies (e.g., lipase inhibitors, bariatric surgery): Individualize based on BMI and comorbidities.^{[13][16]}
- BMI ≥ 27 kg/m² PLUS obesity-related comorbidities: Consider weight loss drugs as an adjunctive treatment.
- BMI ≥ 30 kg/m²
 - With no additional comorbidities: Consider weight loss drugs as an adjunctive treatment. ☑
 - Patients with severe comorbidities (e.g., diabetes, metabolic syndrome): Consider bariatric surgery. ☑^{[4][19]}
- BMI ≥ 35 kg/m² PLUS obesity-related comorbidities OR BMI ≥ 40 kg/m²: Bariatric surgery is indicated.

18. Lifestyle modifications, the primary treatment for metabolic syndrome and obesity, can lead to weight reduction, increased insulin sensitivity, and reduction of cardiovascular risk factors.^[15]

19. Bariatric surgery is a valid option if sufficient weight loss cannot be achieved through lifestyle modifications with or without pharmacological intervention.^[20]

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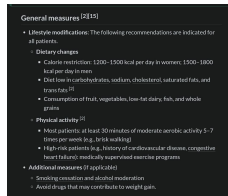
47. Menopause symptoms nbs she crossed her age of menopause so I marked **do nothing**

45 cut off Menopause 40 bhnda kam POF 50 tra vasomotor symptoms HRT

regular chha ki iregular cycle chha ma depend huncha haina ?

Diagnosis hola menopause ko, if regular periods thiyo bhney no investigation, irregular bhaye

48. Person wants to loose weight, what type of **diet reduced calorie diet**, low fat, low protein , diet with no sweets or desserts



49. Case on adhd weird options like eeg, **lead levels** none made sense tbh

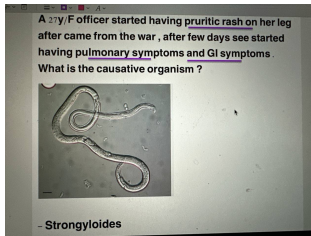
50. X-ray what looked **like detrocardia** I marked immotile sperm

kartageners

51. Post streptococcal glomerulonephritis- type of hypersensitivity **Type 3 . Good pasture 2**

psgn , t-3 HSR , dec complement
IgA nephropathy — no dec in complement

52. Strongyloides with pulmonary symptoms



53. Angular chelosis

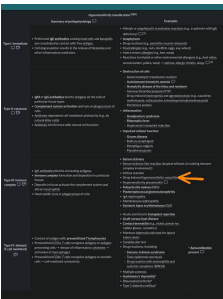
Vitaminnion b 2 Riboflavin and pyridoxine

54. Initial management of transposition of great arteries

“Endomethacin” ends the PDA.
Prostaglandins E1 and E2 KEEp PDA open.

55. Complication of asymptomatic bacteruria in pregnancy

Bacteruria ko Pyelo hunxa pyelo ko preterm Pyelonephritis risk



pyelonephritis and complications (eg, **preterm delivery, low birth weight**) associated with ASB. This is likely related to progesterone-induced smooth muscle relaxation (ie, ureteral dilation, valve laxity) that allows bacteria to ascend to the upper urinary tract. Because **ASB during pregnancy** increases the risk of **acute pyelonephritis**, all patients require **urine culture screening at the initial prenatal visit** and treatment as indicated. **A repeat urine culture (ie, test of cure)** is performed after antibiotic completion to ensure clearance of the bacteriuria. Patients with a **negative repeat urine culture** may resume routine prenatal care. In contrast, those with another positive urine culture require retreatment and another test of cure. **Daily antibiotic suppression** may be indicated in patients with persistent bacteriuria on repeat urine culture or acute pyelonephritis during pregnancy.

- 56. Megaloblastic anemia cbc finding

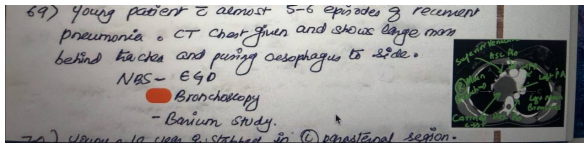
- 57. Couple of ethics question

- 58. Couple of cps/eps

- 59. Biostats questions- did not get any calculations

- 60. New abstracts don't remember

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A 55-year-old man comes to the emergency department due to 2 days of fever, chills, and productive cough. Over the past 4 months, he has had 2 episodes of pneumonia, both of which resolved completely with antibiotics. He smoked a pack of cigarettes per day for 30 years but quit following the second bout of pneumonia. The patient's only other medical problem is hypertension. Temperature is 38.9 C (102 F), blood pressure is 130/86 mm Hg, pulse is 98/min, and respirations are 18/min. Oxygen saturation is 94% on room air. Crackles and increased breath sounds are present over the right lower lung field. Leukocyte count is 14,200/mm³ and blood urea nitrogen is 12 mg/dL. Chest x-ray reveals right lower lobe consolidation. Antibiotics are administered. On follow-up 5 days later, the patient describes improvement in his cough and fever. Crackles are still present at the right lung base. Review of chest x-rays obtained during the previous episodes of pneumonia reveals a consolidation in the right lower lung segment, same as found in the current episode. Which of the following is the most appropriate next step in management of this patient?

- A. Barium swallow study
- B. Bronchoscopy
- C. CT scan of the chest
- D. Repeat chest x-ray in 6 weeks
- E. Serum quantitative immunoglobulin assessment

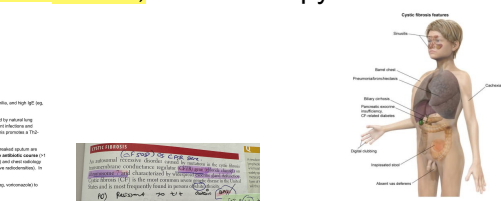


1. CV of cystic fibrosis explained. Persistent pneumonia explained. (Not recurrent pneumonia).

NBS? CT chest, bronchoscopy

Explanation:

- Alveolar bronchopulmonary aspergillosis**
 - Distal allergic bronchopulmonary aspergillosis (ABPA) (HBM)
 - Fungal hypersensitivity reaction - not blood immunobility - allergic inflammation
- Risk factors & pathogenesis**
 - Distal allergic bronchopulmonary aspergillosis (ABPA) (HBM)
 - Fungal hypersensitivity reaction - not blood immunobility - allergic inflammation
- Other features & diagnosis**
 - Distal allergic bronchopulmonary aspergillosis (ABPA) (HBM)
 - Fungal hypersensitivity reaction - not blood immunobility - allergic inflammation
- Treatment**
 - Systemic glucocorticoids - allergic inflammation
 - Antifungal drugs (itraconazole) - acute bronchitis
 - Treatment of underlying asthma (eg, bronchodilators)



1. Young girl with cystic fibrosis. Is unable to perform rapid alternating movements of his hands. Labs show:
 FEV1: 50
 MCV: 90
 BUN: 10
 BUN: 10 (slightly above the upper limit they gave - can't remember exact value)
 Abnormalities: Vitamin was deficient - iron, Vit E, B12, and a couple others.

Recurrent pneumonia is defined as having two or more episodes of pneumonia in a year, or three or more episodes in a lifetime. The episodes must be separated by a month of being asymptomatic or by clear chest X-rays.

Single — mechanical obstruction — suspect malignancy

Same location - mass, different location - immunocompromised

2. Female Dr goes to examine male patient. He says wow what a wonderful Dr. What is best response by Dr. Forgot option but had concept like this question

Direct approach gamru hudaina wala answer auxa ...

76-year-old man with chronic obstructive pulmonary disease is admitted overnight to the hospital due to worsening shortness of breath over the past 3 days. The following morning, a resident physician on the primary team introduces herself and asks the patient how he has been feeling. The patient whistles in response and says, "Wow, a doctor who's beautiful and smart. I'm the luckiest patient in the world." The resident feels uncomfortable but smiles and asks him again how he is feeling today. The patient shares how he has been and the symptoms he is experiencing. During the physical examination, as the resident leans over the bed to auscultate the lungs, the patient reaches up and hugs her. He says, "I just had to give you a hug. I couldn't stop myself." The resident finishes the examination and leaves the room. She immediately reports the incident to her supervising physician. Which of the following is the most appropriate next step for the supervising physician to take?

- A. Advise the resident to tell the patient that his behaviors are unacceptable and ask if she feels comfortable continuing to care for him.
 - B. Discuss with the resident that inappropriate patient behaviors can occur but should not interfere with providing unbiased patient care.
 - C. Document the patient's behavior in the chart and assure the resident that a chaperone will accompany her for future evaluations.
 - D. Share that the resident should have left the patient's room when she first felt uncomfortable instead of staying to complete the evaluation.
 - E. Tell the resident that the patient will be informed of the hospital code of conduct and switched to another physician's care.
- Correct answer is E

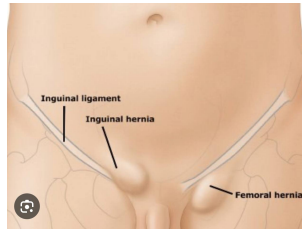
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3. A female kiddo being treated by a physician. The kiddo send fren request to Dr on social media. Next day she comes with mother to office. They don't talk about the request on social media. Best response by Dr.... Forgot option
P.s but had it been by the mother, I'd have accepted 😊)

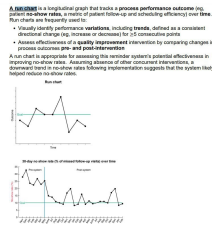
Report him?
 -A child's mom had sent a friend request to a physician on a social media account, now they came to their child's visit. What do you do? Ignore it if she doesn't bring it up or discuss boundaries about social media use etc?
 -A physician gets psychiatric manias->put him in a psychiatric hold?
 -Diverticulitis sequential question get a CT->2nd one was perforation of the

Appropriate response: Thank you for understanding, but I keep my social media strictly personal to maintain professional boundaries

4. Femoral hernia to diagnose



5. Run chart- scenarios given...



6. Effective care

Dimensions of health care quality	
Dimension	Description
Safety	<ul style="list-style-type: none"> Minimizes preventable errors Avoids harms from care
Effectiveness	<ul style="list-style-type: none"> Adheres to scientific guidelines/evidence Avoids undertreatment & overtreatment
Patient-centeredness	<ul style="list-style-type: none"> Identifies patient values, goals & preferences Tailors care delivery to expressed patient values
Timeliness	<ul style="list-style-type: none"> Avoids delays in care, reduces wait times
Efficiency	<ul style="list-style-type: none"> Avoids wasting or overusing resources
Equity	<ul style="list-style-type: none"> Provides quality care to all individuals regardless of demographic attributes (eg, ethnicity, age, gender)

Effective - not (under treatment and over treatment)
 Efficient - use resources that is available in system

Effective vs efficient care
 Scenario: Hospital wants to improve patient treatment by taking
 -Vitals immediately, decrease stay in hospital, fast lab result

Efficient vs effective scenario
 Hospital planned for sepsis prevention / early UT for which they made protocol which included things like:

- when pt arrives at ER take vitals within 5 min
- lab workup within 30 min
- 2 other points were also there similar to further diagnosis in UT

Kunal machine bi overuse garjo vish kono efficiency ghabas i.e
 Avoid wasting or overusing resources

7. Randomisation - for both known and unknown confounder?

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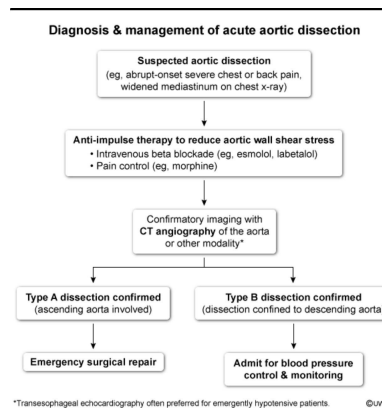
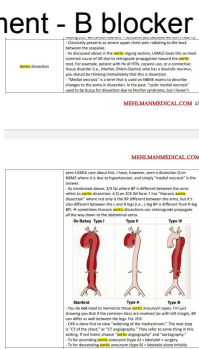
8. Digorge syndrome explained - option 22q deletion

Di George
If catch22 and hypoCa already mentioned in vignette then go with - HypoMg

9. Aortic dissection. CT descending. Treatment - B blocker

Stanford type A - emergency surgery.
Stanford type B (stable) - B-blocker therapy.
Stanford type B (unstable) - surgery.

Dissections involving the ascending aorta can cause aortic regurgitation.



b blocker + nitroprusside if pressure control vayana vane

10. Indication of endometrial biopsy

Endometrial biopsy indications	
Age ≥45	<ul style="list-style-type: none"> Abnormal uterine bleeding Postmenopausal bleeding
Age <45	Abnormal uterine bleeding PLUS: <ul style="list-style-type: none"> Unopposed estrogen (obesity, anovulation) Failed medical management Lynch syndrome (hereditary nonpolyposis colorectal cancer)
Age ≥35	<ul style="list-style-type: none"> Atypical glandular cells on Pap test

11. Husband abuses wife. Wife says the husband doesn't beat the children but they observe it. After safety counseling, NBS? A. Contact CPS B. police c. Do nothing

12. Female has trichomoniasis. Male have no symptoms. Treatment to male? Metronidazole

ceftriaxone+doxy in cervicitis

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	Recommended vaccines for asplenic adult (age >19) patients
Pneumococcus	<ul style="list-style-type: none"> • PCV13 first followed by PPSV23 >8 weeks later • Revaccination with PPSV23 5 years later & at age 65
Haemophilus influenzae	<ul style="list-style-type: none"> • 1 dose Hib vaccine
Meningococcus	<ul style="list-style-type: none"> • Meningococcal quadrivalent vaccine • Revaccinate every 5 years
Influenza	<ul style="list-style-type: none"> • Inactivated influenza vaccine annually
Other vaccines	<ul style="list-style-type: none"> • HAV • HBV • Tdap (tetanus-diphtheria-acellular pertussis) once as substitute for Td (tetanus-diphtheria toxoid booster), then Td every 10 years

13. RTA. 8 yr child. Spleen injury- splenectomy done. What vaccine to give? Td (no strep pneumonia n Hemophilus in option)

Pcv13 Natra
If h/o splenectomy of few days and also h/o of abrasion then TT

14. Splenectomy done. What antibiotics to give. I think amoxicillin/penicillin not in option. Study all antibiotics that can be given after splenecy

Standard regimen ¹	Cephalosporin allergy or intolerance ²
Vancomycin: Administer an initial loading dose (25 to 30 mg/kg, rounded to the nearest 250 mg increment) then give 15 to 20 mg/kg IV every 8 to 12 hours (maximum 2 g per dose) or a total daily dose of 40 mg/kg intravenously. plus either: Ceftriaxone: 2 g IV every 12 hours or Cefepime: 2 g IV every 4 to 6 hours	Vancomycin: Administer an initial loading dose (25 to 30 mg/kg, rounded to the nearest 250 mg increment) then give 15 to 20 mg/kg IV every 8 to 12 hours (maximum 2 g per dose) or a total daily dose of 40 mg/kg intravenously. plus either: Moxifloxacin: 400 mg IV every 24 hours ³ or Mergemont: 2 g IV every 8 hours ⁴
Vancomycin: 15 mg/kg IV every 6 hours (maximum 2 g per dose, intrathecally) plus either: Ceftriaxone: 50 mg/kg IV every 12 hours (maximum 2 g per dose) or Cefepime: 75 mg/kg IV every 6 hours or 50 mg/kg IV every 4 hours (maximum 2 g per dose and 1 g per dose)	Vancomycin: 15 mg/kg per IV every 6 hours (maximum 2 g per dose, intrathecally) plus either: Levofloxacin⁵: 250 mg IV qd and 500 mg IV qd every 12 hours (maximum 750 mg per day) or Mergemont: 40 mg/kg IV every 8 hours (maximum 2 g per dose)

Vaccine	Primary source	Revaccination (boosters)
Pneumococcal vaccine⁶	16 to 20 (13 to 16 for adults)	1 year after primary dose
Haemophilus influenzae type b⁷	1 dose ⁷	1 year after primary dose
Meningococcal conjugate vaccine⁸	1 dose ⁸	1 year after primary dose
Influenza vaccine⁹	1 dose annually	1 year after primary dose

Post splenectomy
Daily Amoxicillin or penicillin for 5 years BD
And
If infection at home - emergency antibiotics : amoxiclav or cefuroxime or fluoroquinolone .
And
After reaching hospital with infection: IV vanco + ceftriaxone if allergic moxifloxacin

Primary	Revised options	Alternative options
Amoxicillin	• Amoxicillin 500 mg qd	• Amoxicillin 500 mg qd
Penicillin V	• Penicillin V 250 mg qd	• Penicillin V 250 mg qd
Ceftriaxone	• Ceftriaxone 1 g qd	• Ceftriaxone 1 g qd
Vancomycin	• Vancomycin 15 mg/kg qd	• Vancomycin 15 mg/kg qd
Moxifloxacin	• Moxifloxacin 400 mg qd	• Moxifloxacin 400 mg qd
Mergemont	• Mergemont 2 g qd	• Mergemont 2 g qd

Prevention of infection in patients with impaired splenic function
Topic: Prophylaxis (13)
Daily antibiotic prophylaxis – We typically determine the need for daily prophylaxis and its duration on an individual patient basis. Factors that we consider in making this determination include patient age, immune status, history of infections with encapsulated organisms, potential antibiotic side effects, local prevalence of antibiotic-resistant organisms, and patient values and preferences. Generally, we favor providing daily antibiotic prophylaxis to asplenic or hyposplenic patients who are at higher risk for severe infections based on young age, concurrent immunocompromising conditions, or history of sepsis caused by encapsulated bacteria. Penicillin and amoxicillin are the preferred agents for daily prophylaxis. Cephalosporins, fluoroquinolones, and macrolides are alternatives to penicillins (see table 5 and 6) (see Antibiotic selection) (below)

15. Student scared to do presentation in school. She is so scared that she is planning to leave school but she goes out freely with friends. Dx ? Agoraphobia? Social anxiety disorder

Agoraphobia

An inordinate fear or anxiety of being in situations that are perceived as difficult to escape from and/or situations in which it might be difficult to seek help. Symptoms must occur over a period of ≥ 6 months in ≥ 2 of the following situations: 1) when using public transportation, 2) when in open spaces, 3) when in enclosed places, 4) when in line or a crowd, 5) when outside of the home alone.

Social phobia	an answer, where person is correct. • Fear of being judged, negatively evaluated, or rejected in a social or performance scenario. Presents on USMLE as fear of public speaking. • USMLE wants beta blocker (propranolol) or atenolol as 1 st line Tx. • If G gives you an anxiety patient, choose beta. They make this distinction on NBME. CBT for longer-term management.
Specific phobia	• Fear of a specific object or situation leading to significant distress or functional impairment. High-yield example on USMLE is fear of flying. • USMLE wants beta for acute relief, CBT for longer-term.
Agoraphobia	• Fear of being in places or situations from which escape might be difficult or embarrassing, or where help might not be available if one were to have a panic attack. This usually refers to crowded, open spaces. • Excessive distress with separation from home or major attachment figure. • Peaks at 12-18 months of age and usually subsides by 2-3 years, but children can still get this age into their teenage years. • USMLE can give a vignette of a child going to summer camp or school who gets
Separation anxiety disorder	

Individuals with **performance-only social anxiety disorder** do not fear nonperformance social situations and are not socially avoidant in general.

Social anxiety disorder (social phobia)	
Diagnosis	<ul style="list-style-type: none"> • Marked anxiety about ≥1 social situations for ≥6 months • Fear of scrutiny by others, humiliation, embarrassment • Social situations avoided or endured with intense distress • Marked impairment (social, academic, occupational) • Subtype specifier: performance only
Treatment	<ul style="list-style-type: none"> • SSRI/SNRI • Cognitive-behavioral therapy • Beta blocker or benzodiazepine for performance-only subtype

SNRI = serotonin norepinephrine reuptake inhibitor; SSRI = selective serotonin reuptake inhibitor.

A 23-year-old woman comes to student health services because of increasing anxiety about being called on in class for the past 6 weeks. Yesterday, she was worried about being called on in class and called her mother. Today, she has to be in a room for a lecture and she has to avoid situations in which she would have to perform or speak in public. Physical examination reveals no abnormalities. On mental status examination, she is cooperative and well dressed. She is anxious and tearful. Her thought process is organized. There is no evidence of suicidal ideation. Which of the following is the most appropriate next step in management?

A. Cognitive behavioral psychotherapy
 B. High-dose benzodiazepine
 C. Beta-blocker therapy
 D. Buspirone therapy
 E. Dose-increased paroxetine

Correct Answer: A.
 1% incorrect answer.

Explanation:
 Social anxiety disorder refers to an excessive fear of scrutiny, embarrassment, or rejection in social and/or performance situations, leading to significant distress and/or impaired functioning (eg, class or work avoidance). Patients with social anxiety disorder commonly exhibit distress and performance anxiety. Performance anxiety refers to the fear of negative evaluation in situations such as public speaking (eg, being called on in class), test-taking, and sex. Cognitive behavioral therapy (CBT) and selective serotonin reuptake inhibitors are the first-line treatments for social anxiety disorder. Some patients with social anxiety disorder experience anxiety that is specific to performance situations only. In the performance-only subtype of social anxiety disorder, CBT is recommended as the first-line treatment, though as-needed (prn) medications (eg, propranolol, benzodiazepines) are frequently used as well. CBT for social anxiety disorder focuses on gradual exposure to feared social situations.

16. aortic aneurysm. Size increase > 0.5 cm in 6 months - sx

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17. 1st trimester. **Usg anechoic**. Doesn't want surgery. Best t/t? **Misoprostol**, mtx, ergot

18. Neonate. High tsh, low T4, lowfT4. **Dx- primary hypothyroidism**

A 34-year-old woman comes to the hospital after a day of sudden onset fever, chest tightness, dyspnea, and dry cough. The patient has been hospitalized twice in the past 3 months with similar symptoms. Both times she received antibiotic treatment for pneumonia and the symptoms resolved within 1-2 days. She does not use tobacco, alcohol, or illicit drugs. She has no known drug allergies. Temperature is 37.9 C (100 F), blood pressure is 128/80 mm Hg, pulse is 92/min, and respirations are 20/min. Pulse oximetry is 90% on room air. BMI is 29 kg/m². The patient is in mild respiratory distress. Examination shows normal jugular venous pressure, no lymphadenopathy, and normal heart sounds. Diffuse fine crackles are heard throughout both lung fields. Leukocytes are 11,200/mm³. CT scan of the chest reveals a bilateral micronodular interstitial pattern. Blood cultures are negative. Which of the following is the most likely diagnosis?

- A. Complement deficiency [10%]
- B. Granulomatosis with polyangiitis [5%]
- C. **Hypersensitivity pneumonitis** [45%]
- D. Idiopathic pulmonary fibrosis [32%]
- E. Influenza pneumonia [4%]

19. CV of hypersensitivity pneumonitis. Asked about investigation

Work place farm gayo Ki pneumonia Ko feature aunxa and rest ma ,house ma huda chai thik hunxa , yestai k thyo question last pool ma

dui tin din aghi chai purai serious feature dini , imaging ma ni nana bhaati bhanne bhare 2-3 din ma sancho

Photos

20. Tinea capitis- treatment - griseo

21 Kaposi sarcoma- CV explained n image given to diagnose





22. Erythema toxicum neonatorum- diagnose

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23. Portwine stain- complication asked- seizure

24. Candidal diaper dermatitis

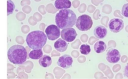
	Diaper dermatitis	
	Candidal dermatitis	Diaper dermatitis
Appearance	Well-demarcated, bright red, satellite lesions	Generalized, non-satellite lesions
Examination		
Treatment	Topical antifungal therapy (e.g., nystatin, clotrimazole)	Topical corticosteroids

- CML
 - o Translocation? t(9;22)
 - o Dx? FISH

Di Podcast Main Document 144

25. PBS and CV explained - asked to diagnose cml

28) Diagnose CML (granulocytosis too much WBC) and histology is given - BCR-ABL / t(9;22)



- o CBC findings? Basophilia
- o Presentation? 45 yo M with fatigue + weight loss + splenomegaly
 - Think of "ML" in CML for "mid-life" - presents in middle-aged pts.)
- o Tx? Imatinib (tyrosine kinase inhibitor)
- o Differentiating b/w CML vs leukemoid reaction
 - LAP high -> leukemoid rxn
 - LAP low -> CML

CML vignette ma Basophil + splenomegaly , LAP score LOW , hit word

26. Ecg- normal- anxiety feature explained in CV Other- probably hocm

Cornel criteria in HOCM MVP chai panic attack jastai hx hunxa - Mehlman


Mitral valve prolapse

- Most common murmur.
- Described as mid-systolic click.
- "Mitral regurgitation" is a busy term that refers to connective tissue degeneration causing MVP in Marfan and Ehlers-Danlos.
- Almost always asymptomatic. On 2D forms, they want you to know about "mitral valve prolapse syndrome," which is symptomatic MVP that presents as repeated episodes of "flashing chest pain" on the left side in an otherwise healthy patient 20s-30s. They might say there is hx of MI in the family, but this is MVP, not MI. Answer on surgery form is "no treatment necessary."
- USMLE loves using MVP as a distractor in panic disorder questions, particularly on the 2d Psych C/S forms. They will give long paragraphs about panic attack/disorder - also mention there's a mid-systolic click, but the MVP isn't the cause of the patient's presentation -> answer = panic disorder, not MVP -> student is confused because they say mid-systolic click, but the MVP is usually incidental, benign, and asymptomatic.
- MVP does not progress to mitral regurg almost always. So don't think that MVP and MR are the same.

Hypertrophic Cardiomyopathy (HCM) Characteristics

Typical ECG findings may include:

- Deep and narrow Q waves in the lateral and inferior leads
- General signs of left bundle branch repolarization abnormalities
- Deeply inverted T waves in the precordial leads are seen in the apical variant.



S wave in V1 + R wave in V5 or V6 > 35

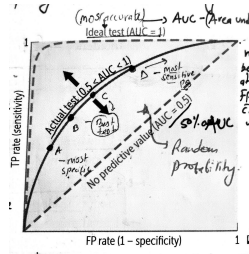
Mitral valve prolapse, albeit a common condition, is associated with ventricular arrhythmias and sudden cardiac death. Electrocardiographic features include ST-segment depression, T wave inversion or biphasic T waves in the inferior leads, QT prolongation and premature ventricular complexes

- ECG findings may include [6]:
- Prominent abnormal Q waves, particularly in the inferior (II, III, and aVF) and lateral leads (I, aVL, and V4-V6). These changes reflect septal depolarization of the hypertrophied myocardial tissue. (See "Pathogenesis and diagnosis of Q waves on the electrocardiogram".)
 - P wave abnormalities, reflecting left atrial (LA) or biatrial enlargement. The combination of LVH with right atrial enlargement is strongly suggestive of HCM.
 - Left axis deviation.
 - Deeply inverted T waves (so-called "giant negative T waves") may be seen in the mid-precordial leads (V2 through V4) in patients with the apical variant of HCM. (See "Hypertrophic cardiomyopathy: Morphologic variants and the pathophysiology of left ventricular outflow tract obstruction", section on 'Apical HCM'.)

27. CV of post herpetic neuralgia. NBS for dx. Nothing other option biopsy

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28. Roc curve- best of sensitivity



29. Kaplan Meir curve

30. Sjogren- SSA, ana

Anti ssa, anti ro
Anti ssb, anti la

Similar question from nbme/ free 120

A 5-day-old boy is brought to the office for an initial well-child examination. He was born at 40 weeks' gestation and discharged at 60 hours of life. On newborn screening, hemoglobin electrophoresis showed an FS pattern. He is at the 50th percentile for length and weight. Temperature is 37.0°C (98.6°F), pulse is 136/min, and respirations are 34/min. He appears well. Examination shows no abnormalities. Which of the following is the most appropriate next step in management? (A) Deferoxamine therapy (B) Hydroxyurea therapy (C) Iron supplementation (D) Monthly blood transfusions (E) Penicillin prophylaxis (answer). (F) Vitamin B12 (cyanocobalamin) supplementation

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A 42-year-old woman comes to the physician because of a lump in her left thigh since she fell on the ice while playing hockey 4 months ago. At that time, she noticed bruising on her left thigh that has resolved, but the lump has increased in size. She has not had fever, night sweats, or weight loss. She has no history of serious illness and takes no medications. Examination shows a 5 × 6-cm, painless, firm, soft tissue mass on the anterolateral aspect of the left thigh. The overlying skin is intact, and there is no discoloration. There is no inguinal lymphadenopathy. Distal pulses are normal, and sensation is intact. Range of motion of the hips and knees is full, and muscle strength is normal. X-rays of the left femur and knee show no abnormalities. An MRI of the left femur shows a deep soft tissue mass with inhomogeneous signal on T1- and T2-weighted images. Which of the following is the most appropriate next step in diagnosis?

- A.
CT scan of the thigh
 - B.
Core-needle biopsy Fibrosarcoma.
 - C.
Ultrasonography
 - D.
Wide resection of the mass
 - E.
Observation and reevaluation in 1 month
- Correct answer is B

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Digeorge - fisch of ch 22

Di George

If catch22 and hypoCa already mentioned in vignette then go with - HypoMg

-Pul emb- to diagnose. ECG was also given, pt took 8 hrs flight. Fligh duration implies immobility

-ITP- bacha, mild infection month ago, fever 3-4 days back and rash. Platelet size normal Kei nagarm until symptomatic Life threatening bleeding if present.. platelets infusion only if needed

-Circumcission k baad bleeding, Platelet count normal. What to check? Thrombin time, factor 8, platelet antibody etc options. Vwf factor 8

-Girl, factor 8 mildly decreased- type 1 vwd sibling , father aru lai pani bleeding manifestation deko huna sakcha , aut dominant

-GFR 40/min, hypokalemia/hyperkalemia (asked arrows about renin, aldo etc)

-RAS- investigation asked to diagnose(renal doppler)

-yoga abstract. 1 question was to calculate NNT

- Transient synovitis treatment? Nsaids? No reassurance

- Kawasaki treatment. High dose aspirin + echo 2 weeks

- Bacha ataxia, bilateral papilledema— intracranial tumor Meduloblastoma

- Hemangioma, mathay pe lump flesh coloured enlarge ho raha. 18 months old.

- Nigerian aurat, delivery karni thi, labia majora cut, grade 4 laceration hui v jo theek kar di, family says pehli wali halat mein kar k do. What principle? Autonomy, nanmalificence

- Sharp ulcers in mouth, sore throat, female, cervical + inguinal lymphadeno. dx? HIV

- Barbar, cutting karti tou baazu mein dard. Dx? SVC?(why not subclavian steal

- Hematopoitic transplant, uske baad diarhea, rash—gvhd

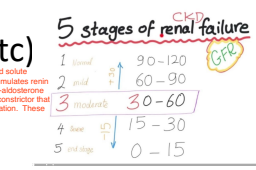
- Pregnant, ast alt 500, Hep E?

- 16yrs girl, 12 yrs pe UC hogya but no flare ever,ab jaundice, ast alt raised, inc. Direct bilirubin—Primary sclerosing. No PBC in options. AMA=female,PBC

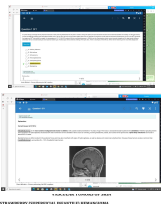
- Bone tumors, osteochondroma, what to do? Biopsy or reassurance?

Table with 2 columns: Clinical features, Laboratory findings. Rows include: Type 1: Partial deficiency of vWD factor, Less deranged aPTT; Type 2: Complete deficiency of vWD factor, More deranged aPTT; Type 3: Defect in vWD factor, aPTT Normal.

Table with 2 columns: Clinical features, Management. Rows include: Transient synovitis (Age 3-8, preceding viral illness, Able to bear weight), Diagnosis (Normal to mildly elevated WBC count, ESR, CRP), Management (Conservative, Nonsteroidal anti-inflammatory medications), Prognosis (Full recovery within 1-2 weeks, Recurrence uncommon (<15%)).



sometimes 8 year old ma perthe's ma jhukaucha, tay long history , 1-2 months , transient synovitis , short history



nanmalificence: Hence doctor should not provide treatment because of NORMALMALIFICENCE

HIV: Patient demanding test doc not doing... Justice. Elderly psychosidistellium wanting to discharge—doc stops... Beneficence. Doctor didn't provide same information to both patient for minimal invasive surgery... Justice. Patient centered care.

SVC: Subclavian steal syndrome

Hep E: Hepatitis E virus

Primary sclerosing: Primary biliary cirrhosis

reassurance: Bone tumors, osteochondroma

Meduloblastoma can manifest with signs of increased intracranial pressure, including headaches, nausea, vomiting, and papilloedema. F.uchterman, a hemorrhage from a neoplasm could manifest with the sudden onset of a severe headache and altered mental status. However, since most meduloblastomas arise within the cerebellum, the lack of any cerebellar signs (e.g., gait ataxia, truncal instability, intention tremor, double vision) makes meduloblastoma less likely. Fever and a 3-week history of both pain suggest another etiology for this patient's symptoms.



Africa ma female circumcision huncha Culture respect garera autonomy. Aba nagame - non maas kina silaidyo - beneficence kti lai sodhera silayo --- autonomy kti ko parwar/kti ko against ma gayera silaidyo --- autonomy violate bhayo

KEY FACT: 1. Enlarged, tender, axillary, subclavian lymphadenopathy, BCRCL, and... 2. Dysphagia, odynophagia, weight loss, anorexia, TIA or stroke... 3. Change in voice, hoarse... 4. Change in voice, hoarse... 5. Spontaneous pneumothorax, rib fracture, etc.

Subclavian steal syndrome occurs when there is a significant stenosis or occlusion of the subclavian artery proximal to the origin of the vertebral artery. The resulting reversal of blood flow in the vertebral artery of the affected arm "steals" blood from the brain and leading symptoms such as dizziness, fatigue, and syncope, especially during arm activity. The presence of a bruit over the vertebral artery and the absence of a bruit pressure between the arms suggest the diagnosis.

16-year-old male presents to the primary care physician with complaints of dizziness, fatigue in his left arm, and occasional episodes of pruritus, particularly when using the left arm for activities such as combing the hair or styling it back. He reports that these symptoms have been progressively worsening over the past few months. He has no other symptoms, there is no family history of liver disease, and he has never traveled to the right areas, with the left arm showing rigidity over swelling. A bruit is heard over the left subclavian artery. Which of the following is the most likely diagnosis?

Subclavian steal syndrome occurs when there is a significant stenosis or occlusion of the subclavian artery proximal to the origin of the vertebral artery. The resulting reversal of blood flow in the vertebral artery of the affected arm "steals" blood from the brain and leading symptoms such as dizziness, fatigue, and syncope, especially during arm activity. The presence of a bruit over the vertebral artery and the absence of a bruit pressure between the arms suggest the diagnosis.

- Seq of Interstitial cystitis a) investigation cystoscopy then diagnosis(interstitial cystitis)

USG garera confirm garne then Catheterization hoina

Small prostate <40ml and serum psa <1.5ng/ml alpha blocker
Large prostate >40ml and serum psa > 1.5ng/ml finasteride

- Seq. GA mein operation, anuria. Inv? Us then b) ~~alpha blocker~~ Intermittent catheterization

- Covid mein excellent work by physician wala. Bipolar walay symptoms? Take to ER?

(Choice C) The depressive and hypomanic episodes in bipolar II disorder may resemble the mood instability seen in **borderline** personality disorder. However, the labile mood states in **borderline** personality disorder are brief, typically lasting hours to days (rather than weeks to months). There is insufficient evidence to diagnose **borderline** personality disorder in this patient; she does not exhibit a lifelong pattern of identity disturbance, feelings of emptiness, intense anger, splitting, or self-mutilating behavior.

(Choices D and E) Major depressive disorder and persistent depressive disorder (dysthymia) are unipolar disorders; these patients do



- Borderline mein sab se ziada risk? A) cyclothymia b) bipolar

Yedi curette garera falisakeko follow up ma ho bhane GTN jam , pet mai chha POC bhane mole jaam k bhannu huncha
Less bhaye partial mole

- Bhcg 64000, kis cheez ka risk? Hydatidiform mole? Choriocarcinoma?

- 1st time 16weeks gestation, kia approp hai to establish WOG? Fetal height, ya ultrasound?

Laboratory evaluation
hCG — a quantitative serum hCG should be measured. The serum hCG concentration in patients with mole is usually higher than that observed with digitalis concentration or actual pregnancy of the same gestational age.
If the serum hCG level is high (>10000 U/L) a nonpregnant ultrasound should be performed and will likely demonstrate molar disease if present. If the hCG level is high and the ultrasound shows an apparently normal pregnancy, the ultrasound and hCG should be repeated in one week to exclude the possible presence of a nonconception with normal fetus and undetected molar pregnancy. Markedly elevated hCG levels are more commonly seen in patients with complete mole in contrast to partial mole (5% of an average 10

- Trauma, xray mein dono taraf pneumo. Needle compression

- Still murmur, AS ka , MR ka. PDA vayo still murmur

- Bacha ko continuous murmur, kia dogay? Indomethacin.

PDA khula rakhna — Alprostadil

- Sturge weber rash? Kia develop ho sakta? Epilepsy

PDA banda garna — indomethacin

- NF 1 ka rash decribled, kia develop hoga? Perpheral sheath tumor ya Schwannoma?

- Osteo ka sawal. Kia karengey? Vigourous exercise, light exercise, resistance exercise two times a week.

- 65 yrs old, kisi cheez ka player, weight gain ho raha tha, ostarthritis settle nai ho raha tha conservative se, ab kia karogay, 6 weeks more exercise ya topica diclo etc

- RA, hath bhi dikhaya hua, beti ka jora seena hai? Kia dogay? Splint ya steroid? Occupational therapy

- Baba marnay wala, beti kehti continue treatment, beta(who is attorney) kehta no? Kia karogay? Go with beta ya family meeting? Family meeting more then son

- Varicella lesions pictures, most approp next step in management karogay? Coughing aur sneezing se bacho ya oral acyclovir.

Timing and regimen — If antiviral therapy is initiated (eg, immunocompetent patients at increased risk of developing complications from varicella), treatment should be started within 24 hours after the rash develops, if possible. Clinical trial data found greater benefits when antiviral therapy was initiated within 24 hours of symptoms onset [5,21]. In addition, in immunocompetent hosts, viral replication typically stops by 72 hours after the onset of rash.



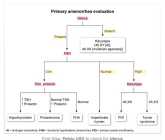
- 46 yr old homosexual, kia karogay? HPV?

yo sab garisakya hunxa ... nothing ma jane
Hiv hepatitis syphilis garni garnu xa yani

- DM type 1 and pregnant? Sab se ziada risk kis ka? Accreta, previa, abruption, preeclam?

within 72 hours or within two weeks after onset
• Indications (21)
• Age (20)
• Moderate or severe rash and/or
• Immunocompetent patients
• High or increasing fever and/or
• Multiple complications
• High risk for development of serious complications
• Regimen (21)(2)
For immunocompetent patients
Choose one of the following:
• Acyclovir
• Valacyclovir
• Famciclovir
For immunocompetent patients and/or
those with immunocompetent hosts (2)
Duration
Acyclovir should be initiated as early as possible after the effectiveness of antiviral treatment decreases in the absence of response.

System	Organ	Common Pathogens	Common Pathogens
Respiratory	Trachea	Staphylococcus aureus, Klebsiella pneumoniae, Pseudomonas aeruginosa, Haemophilus influenzae, Moraxella catarrhalis, Streptococcus pneumoniae, Legionella pneumophila, Mycoplasma pneumoniae, Chlamydia pneumoniae, Coxiella burnetii, Histoplasma capsulatum, Cryptosporidium parvum, Isospora belli, Cyclospora cayentensis, Pneumocystis carinii, Toxoplasma gondii, Aspergillus fumigatus, Aspergillus niger, Aspergillus terreus, Aspergillus nidulans, Aspergillus glaucus, Aspergillus versicolor, Aspergillus nidulans, Aspergillus terreus, Aspergillus nidulans, Aspergillus glaucus, Aspergillus versicolor, Aspergillus nidulans, Aspergillus terreus, Aspergillus nidulans, Aspergillus glaucus, Aspergillus versicolor	Staphylococcus aureus, Klebsiella pneumoniae, Pseudomonas aeruginosa, Haemophilus influenzae, Moraxella catarrhalis, Streptococcus pneumoniae, Legionella pneumophila, Mycoplasma pneumoniae, Chlamydia pneumoniae, Coxiella burnetii, Histoplasma capsulatum, Cryptosporidium parvum, Isospora belli, Cyclospora cayentensis, Pneumocystis carinii, Toxoplasma gondii, Aspergillus fumigatus, Aspergillus niger, Aspergillus terreus, Aspergillus nidulans, Aspergillus glaucus, Aspergillus versicolor
Respiratory	Lung	Staphylococcus aureus, Klebsiella pneumoniae, Pseudomonas aeruginosa, Haemophilus influenzae, Moraxella catarrhalis, Streptococcus pneumoniae, Legionella pneumophila, Mycoplasma pneumoniae, Chlamydia pneumoniae, Coxiella burnetii, Histoplasma capsulatum, Cryptosporidium parvum, Isospora belli, Cyclospora cayentensis, Pneumocystis carinii, Toxoplasma gondii, Aspergillus fumigatus, Aspergillus niger, Aspergillus terreus, Aspergillus nidulans, Aspergillus glaucus, Aspergillus versicolor	Staphylococcus aureus, Klebsiella pneumoniae, Pseudomonas aeruginosa, Haemophilus influenzae, Moraxella catarrhalis, Streptococcus pneumoniae, Legionella pneumophila, Mycoplasma pneumoniae, Chlamydia pneumoniae, Coxiella burnetii, Histoplasma capsulatum, Cryptosporidium parvum, Isospora belli, Cyclospora cayentensis, Pneumocystis carinii, Toxoplasma gondii, Aspergillus fumigatus, Aspergillus niger, Aspergillus terreus, Aspergillus nidulans, Aspergillus glaucus, Aspergillus versicolor
Respiratory	Alveoli	Staphylococcus aureus, Klebsiella pneumoniae, Pseudomonas aeruginosa, Haemophilus influenzae, Moraxella catarrhalis, Streptococcus pneumoniae, Legionella pneumophila, Mycoplasma pneumoniae, Chlamydia pneumoniae, Coxiella burnetii, Histoplasma capsulatum, Cryptosporidium parvum, Isospora belli, Cyclospora cayentensis, Pneumocystis carinii, Toxoplasma gondii, Aspergillus fumigatus, Aspergillus niger, Aspergillus terreus, Aspergillus nidulans, Aspergillus glaucus, Aspergillus versicolor	Staphylococcus aureus, Klebsiella pneumoniae, Pseudomonas aeruginosa, Haemophilus influenzae, Moraxella catarrhalis, Streptococcus pneumoniae, Legionella pneumophila, Mycoplasma pneumoniae, Chlamydia pneumoniae, Coxiella burnetii, Histoplasma capsulatum, Cryptosporidium parvum, Isospora belli, Cyclospora cayentensis, Pneumocystis carinii, Toxoplasma gondii, Aspergillus fumigatus, Aspergillus niger, Aspergillus terreus, Aspergillus nidulans, Aspergillus glaucus, Aspergillus versicolor



- 32 wog pe kia vaccine? Tdap **27 to 36**

Routine Prenatal Care	
First Trimester (first visit by 13 weeks gestation)	<ul style="list-style-type: none"> Lab Tests: CBC, blood type, Rh screen, UA and culture, Rubella, Varicella antibody screen, RPR/VDRL, chlamydia, HIV, HBsAg, PTV, Consider high ALT, TORCH and factors on symptoms. NIPT or APO screening for possible hemoglobinopathy Thrombophilia analysis (for APS or DVT of legs, Test dual if more carriers concerning genes) Concern over family history of genetic conditions (Cystic Fibrosis, Tay Sachs) Genetically test screen. Test dual if more carriers concerning genes.
Second Trimester	<ul style="list-style-type: none"> Anomaly screening (either late first or early second trimester) Serial fetal defect screen (15 + AFP) Fetal anatomy (19-22 weeks) Children screen (16-20 weeks): 1-hour glucose challenge RhesusG for Rh- women (at 28 weeks), HIV, GBS Culture (35-37 weeks)
Third Trimester	<ul style="list-style-type: none"> RhesusG for Rh- women (at 28 weeks), HIV, GBS Culture (35-37 weeks)

Note: Visit qd week (up to 28 wks), q2 week (29-34 wks), q1 week (34 wk - birth)

- Pregnancy se pehle kia check karogay? Varicella??(confirm urself)

- Aurat ko bachpan se myelomeningo, vegan, pregnant incidental? In addition to prenatal vitamins, kia dogay? **Folic acid, vit d, b12?**

Primary amenorrhea : First do US
Secondary: bHc

- 48 yrs old, amenorrhea, kia karogay? **Bhcg, fsh, do nothing(confirm urself)**

- Pt alcohol addicted, ab chhor rahi thi, kal raat ko pee k neend aajaye, kal court meeting pe jaana, kia karogay? **Blood alcohol, usg, do nothing? Ast 80 tha, alt 40. (Ratio 2:1 ban rahi)**

- Aortic dissec, 90/60. CT angio ya echo **complication ma gayo Rupture TEE while preparing for OT**

- Aik banda, intermittent explosive disorder, bachi ko belt se **maara maheena pehle jo 6 saal** ki thi but remorse. Psychotherapy ya **cps? Regret garera kaha sukha pahixa ra ...**

- Seizures after party, garmi lag rahi, BP high, Temp? Cocaine, lsd, **mdma.**(confirm urself)

Hyponatramic seizure
mdma
hyperthermia
bruxism
hyponat

Euta MDMA chai party ma chill hanna khane wala Low sod sodhya xa

- Alzheimer—rivastigmine

- Parkinson ki dawai start karatay, symptoms improve but develop psychosis, he says dawai kamm nai karni, kia karogay? Quetiapine.

- smoker, hematuria, histo given(RCC), kia assoc hoga? **Hb>20, ya urea nitrogen >80?**

- IgG bohat ziada, ca normal. **Bone biopsy he did for myeloma.**

- Bandy ko stones, ulcer bhi jo khany se theek hojata hai? Kia check karogay?

Gastrin. Duodenal ulcer

Men1 3p

- Urine metaneph increase, ca increased? Kia karogay? **Calcitonin.**

Men2

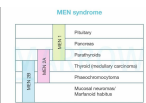
Pituitary tumor
Parathyroid adenoma
Pancreatic endocrine tumor

- Patient ko penicillin allergy, kia contraindicated hai? Cephalexin?

CEPHALOSPORIN nadim

- Bartonella k do questions. Aik mein papule and chronic(5-6 weeks) tender unilateral lymphadenopathy. Doosre mein hath mein scratches and again tender lymph

- CGD— staph aur



1. A 35-year-old female presents with a 2-year history of weight gain, fatigue, and decreased libido. She has a family history of hypertension and diabetes. Her physical exam is unremarkable. Laboratory studies show the following: TSH 0.1 mIU/L, FT4 4.5 pmol/L, Prolactin 1500 mIU/L, and HbA1c 6.5%. Which of the following is the most likely diagnosis?

- Pituitary adenoma
- Hyperprolactinemia
- Thyroid adenoma
- Acromegaly

Secondary amebiasis	
Reservoir organisms	<ul style="list-style-type: none"> • <i>Sporella hominis</i> • <i>Cat</i> spores or oocysts (rare) • <i>Trichostema</i>
Etiology	<ul style="list-style-type: none"> • Ingestion of contaminated food • Ingestion of contaminated water • Ingestion of contaminated produce
Manifestation	<ul style="list-style-type: none"> • Acute enteritis • Chronic enteritis • Colitis • Colitis with abscess • Colitis with fistula • Colitis with abscess and fistula
Diagnosis	<ul style="list-style-type: none"> • Stool study with trichostema • Serology
Treatment	<ul style="list-style-type: none"> • Metronidazole or tinidazole • Supportive therapy

Affected lymph nodes are enlarged, tender, and have overlying erythema, suppurative in the common. Lymphadenopathy may take 1-2 months to resolve. Although many mild cases will self-resolve, treatment with rifampin is typically recommended.

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- SCID ka best treat. Transplant.

- Sickle cell— penicillin prophylaxis

- G6pd def scenerio— all ani malarials mentioned, which ones causes it?(highest risk). Primaquine?(confirm urself) **Female sulfa drugs wala autoimmune hemolytic**

Medications that may trigger hemolysis in G6PD deficiency	
Use with caution (probably safe)	<ul style="list-style-type: none"> Chloramphenicol Chloroquine Quinine Trimethoprim-sulfamethoxazole
Avoid (likely unsafe)	<ul style="list-style-type: none"> Dapsone Fluoroquinolones (eg, ciprofloxacin) Methylene blue Nitrofurantoin Primaquine Riboflavin Sulfonamides

*Some of these medications were previously considered unsafe in G6PD deficiency but are now considered probably safe if given in standard doses. G6PD = glucose-6-phosphate dehydrogenase.

- Cml scenerio, metamyelocytes etc given, asks about pathophys— bcr-abl fusion related.

- Bipolar patient, which drug contraindicated? No ssri in the options, went with **Lorazepam.(confirms urself) Imipramine**

- Personality disorder— starts with something related to ADHD(patient says he has read about adhd in adults etc)— then switches to patient describing himself that he has many newspaperes and does everything perfect, wish everyone was like him etc etc... narcissistic personality disorder, histrionic, ocpd??

Ma nai hu-- Narcissist
Bhane jastai huna paryo --
OCPD

30) Army going for sentry duty...
command + break rules, **stalls vehicle**. Says its not his problem if they can't get their belongings safe, **stalls van**.

Clonus ko funda

- Pt on escitalopram therapy for mdd, takes over the counter cough syrup for 2-3 days. His roommate brings him in with symptoms like DTRs 3+, ankle clonus etc—

- CJD
-serotonin syndrome
- Tay sach dis

dx? **Serotonin Syndrome.**

Diagnosis	Key features	Management
Serotonin syndrome	Agitation, hyperreflexia, rigidity, tachycardia, hyperloquacity, diaphoresis, hyperthermia, clonus, rigidity, hyperreflexia, tachycardia, hyperloquacity, diaphoresis, hyperthermia	Discontinuation of serotonergic agents, supportive care, benzodiazepines, cyproheptadine
Neuroleptic malignant syndrome	Hyperthermia, rigidity, autonomic instability, altered mental status, hyperreflexia, tachycardia, hyperloquacity, diaphoresis, hyperthermia	Discontinuation of neuroleptics, supportive care, dantrolene, benzodiazepines

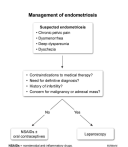
DTR+++ --> Serotonin syndrome
Flu like symptoms --> SSRI discontinuation syndrome

- Pregnant lady, has acne(probably comedonal, not moderate nor severe), Tx? Isotretinoin, oral doxycycline, **benzoyl peroxide(he marked this)**. What if reassurance in the options?(confirm yourself)

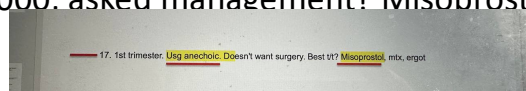


- 6-7 yrs old(age could vary) bacha/bachi, adopted, easily adjust hogya with new family, shopping mall mein strangers k sath ghul mil jata hai, doctor ko bhi "big fat hug" deta hai, Dx asked? **Disinhibited child disorder.**

- Endometriosis scenerio with associated infertility—Tx? No ocp in options, so Laparoscopy (he did)



- Ectopic pregnancy scenerio with bhcg <5000. asked management? Misoprostol, mifepristone, **methotrexate(he marked)**



- Mother ko endometrial cancer, behan ko ovarian, this patient has some menstruation problem, which cancer is she at inc risk for? **Endometrial cancer(he did)**

- Arrows for IDA and Anemia of Chronic disease.

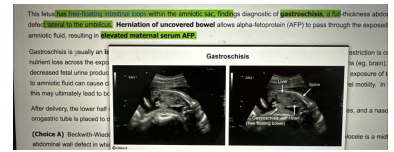
Iron studies in microcytic anemia					
Cause	MCV	Iron	TIBC	Ferritin	Transferrin saturation (Iron/TIBC)
Iron deficiency	↓	↓	↑	↓	↓
Thalassemia	↓↓	↑	↓	↑	↑
Anemia of chronic disease (inflammation)	Normal/↓	↓	↓	Normal/↑	Normal/↓

MCV = mean corpuscular volume; TIBC = total iron binding capacity. ©2019

Cta set May combine page 9

1. Female pregnant came at the 20 weeks of gestation on ultrasound baby Intestinal content were protruding(commig out in the umbilicus) what complication it can develop in future

- A. gastroschesia
- B. omphalocele**
- C. mesentic ishenia



2. . Long history of gerd in endoscopy the squmular to columnar changes in the esophagus mention in CV no pic the beside prescribing th point what will you do

Ans= endoscopic Surveleine

3. 2-3 weeks ago have urti now have hf signs with S3 mention in CV asked for diagnosis

I did myocarditis.

Myocardial Complications of acute myocardial infarction	Time course	Pathological changes	Clinical findings	Investigative findings
Pericarditis	1-2 days	Pericardial inflammation	Pericardial pain, friction rub	ECG: ST-segment depression, PR-segment depression
Papillary muscle dysfunction	1-7 days	Ischemic necrosis of papillary muscles	Acute mitral regurgitation	ECG: ST-segment elevation, Q-waves
Septal perforation	1-7 days	Septal necrosis	Left-to-right shunt	ECG: ST-segment elevation, Q-waves
Free wall rupture	1-7 days	Free wall necrosis	Pericardial effusion with tamponade	ECG: ST-segment elevation, Q-waves
Left ventricular aneurysm	1-6 weeks	Left ventricular aneurysm	Thrombus formation, embolism	ECG: ST-segment elevation, Q-waves

4. Patient 2 to 3 days ago have the Myocardial infarction now have murmur lungs bl crakle caused aksed

I chose papillary muscle rupture.

Papillary rupture- 2- 7 days, hypotension, cardiogenic shock , MR

Myocardial Complications of acute myocardial infarction	Time course	Pathological changes	Clinical findings	Investigative findings
Pericarditis	1-2 days	Pericardial inflammation	Pericardial pain, friction rub	ECG: ST-segment depression, PR-segment depression
Papillary muscle dysfunction	1-7 days	Ischemic necrosis of papillary muscles	Acute mitral regurgitation	ECG: ST-segment elevation, Q-waves
Septal perforation	1-7 days	Septal necrosis	Left-to-right shunt	ECG: ST-segment elevation, Q-waves
Free wall rupture	1-7 days	Free wall necrosis	Pericardial effusion with tamponade	ECG: ST-segment elevation, Q-waves
Left ventricular aneurysm	1-6 weeks	Left ventricular aneurysm	Thrombus formation, embolism	ECG: ST-segment elevation, Q-waves

5. . COPD patients fev1to fvc ratio 50 what will you see in changes in the heart asked in ups and downs arrow question. >sarcomere added in series **>sarcomere added in parallel.**

Parallel ma - increase
Series ma - decrease

If a schizophrenia patient who is already stabilized on medication suddenly wants to get rid of schizophrenia quickly, the likely outcome would be:

- A. His condition further deteriorates.

Schizophrenia is a chronic mental disorder that typically requires ongoing treatment with antipsychotic medications and often involves therapy and support. Stopping medication abruptly or trying to "get rid" of schizophrenia quickly without proper medical supervision can lead to a worsening of symptoms and a relapse of the disorder. It's important for individuals with schizophrenia to continue with their prescribed treatment plan under the guidance of healthcare professionals to maintain stability and manage symptoms effectively.

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6. schizophrenia patient comes to follow already drugs now stabilize doing routine activities going to college taking major courses to complete his wants to get quickly rid of schizophrenia what will happened .

Aru sathi jastai normal feel garxa
Ani khana xodxa medicine

A. His condition further deterots

B. failure of therapy

Hectic schedule , xito thik garna xa vanepaxi , overdose khanxa
hola ni xito thik garna

C. will get better

Compliant chha bhane C jaam

Qn ko vignette pt aauxa stable xa aile. Ma aba khana xaadxu thik vaysakyo jasto kura garxa.

Ans is something like you have to continue the medication to get better estai estai khaalko qn xa

7. Schizophrenia patient on haloperidol develops the signs of **aksthesai**
akxed MOA of drug **4day to 40 days**

A. Sentization of dopamine receptor

Dopamin receptor hypersensitivity vanne auta thiyo ni paila?

B. dopamine blocks

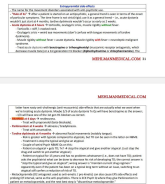
typical antipsychotic that primarily acts by blocking dopamine receptors in the brain. This blockade helps to reduce the positive symptoms of schizophrenia, such as hallucinations and delusions. Over time, however, the prolonged blockade of dopamine receptors can lead to a compensatory increase in the sensitivity of dopamine receptors (sensitization). This phenomenon may contribute to the development of side effects like akathisia, which is characterized by motor restlessness and a compelling urge to move. Therefore, option A is the correct answer regarding the MOA of haloperidol and its relation to the de

Akathisia should be considered if a patient becomes agitated or restless when an antipsychotic is changed or the dosage is increased. Akathisia is the most common extrapyramidal symptom associated with antipsychotic medication. Patients with akathisia describe a subjective sense of inner restlessness that may manifest as pacing or an inability to sit still (eg, walking all day in the neighborhood). In severe cases, patients may become extremely distressed, resulting in increased agitation and overall global worsening. Clinicians must differentiate akathisia from worsening psychotic agitation because akathisia is dose dependent. An increase in antipsychotic dosage would make akathisia worse. Akathisia is more common with high-potency, first-generation antipsychotics, but can also occur with newer second-generation antipsychotics, such as risperidone. The first steps in management of akathisia include the following:

- Cautiously reducing the antipsychotic dosage
- Changing to an antipsychotic with less potential to cause extrapyramidal symptoms
- Adding a beta-blocker (eg, propranolol), an anticholinergic (eg, benztropine), or, less commonly, a benzodiazepine (eg, lorazepam)

Propranolol is a common initial choice; it may work by blocking noradrenergic and serotonergic inputs on dopamine pathways.

The first step in management of akathisia would be a dose-reduction trial (ie, if clinical benefit is not significant, discontinuation of the antipsychotic entirely). The beta-blocker propranolol or a benzodiazepine may be used for symptomatic management of akathisia if antipsychotic reduction or discontinuation was failed. If psychotic symptoms worsen following discontinuation, a trial of a different antipsychotic medication (eg, quetiapine) with less potential to cause extrapyramidal symptoms could be considered.



8. Patient have previous history of 3rd degree skin burned got skin graft now to work as construction have **to work at outdoor** what he is at risk **of**

A. SCC due uv light

B. Ulcer

9. Patient with 2 months history of **acute memory loss** cant remember things short question also there was flat effect asked cause MDD Alzheimer's **crudz jacob disease**

Cruetzfeldt-Jakob Disease
Cruetzfeldt-Jakob Disease is a rare prion protein misfolding disease that causes a rapidly progressive neurodegenerative disorder. It is characterized by rapidly progressive dementia, myoclonus, and motor signs and symptoms. The disease is caused by a misfolded prion protein that induces other prion proteins to misfold, leading to a chain reaction of protein aggregation. The disease is fatal, with a median survival time of approximately 14 months. The disease is named after the German neurologist Hans Christian Cruetzfeldt and the British neurologist Robert W. Jakob.

Causes:

- Sporadic: Most cases of the disease are sporadic, with no known cause.
- Genetic: Some cases are inherited, with a mutation in the prion protein gene (PrP^C).
- Iatrogenic: Some cases are caused by medical procedures, such as the use of contaminated surgical instruments or the consumption of contaminated meat.

Diagnosis:

- Clinical: The disease is diagnosed based on the clinical presentation, including rapidly progressive dementia, myoclonus, and motor signs and symptoms.
- Laboratory: Laboratory tests, such as the detection of PrP^{Sc} in the cerebrospinal fluid or the presence of PrP^{Sc} in the brain tissue, can confirm the diagnosis.

Management:

- Supportive: There is no specific treatment for the disease, and management is primarily supportive, focusing on symptom relief and patient care.

Prognosis:

- Fatal: The disease is fatal, with a median survival time of approximately 14 months.

Key features of the disease include a long incubation period with average onset of age 60-70. Most cases (95%) are sporadic, with the remaining being hereditary or iatrogenic (due to contaminated surgical instruments or contaminated meat). It is a rapidly progressive, fatal neurodegenerative disorder with a median survival time of 14 months. The disease is named after the German neurologist Hans Christian Cruetzfeldt and the British neurologist Robert W. Jakob.

10. Womens with **band like headaches for 8months pain worse** when she goes to job until she comes home and pain has inc in duration previous 2 to 3 times a weeks now has inc frequency dx

Characterization of chronic headache	
Headache subtype	Clinical presentation
Migraine	<ul style="list-style-type: none"> Female predilection Unilateral, throbbing Mechanically, photophobia
Tension type	<ul style="list-style-type: none"> Global or near bilateral pressure Not to increase without previous acute tension headaches
Cluster	<ul style="list-style-type: none"> Male predilection Unilateral, ipsilateral orbitonasal with ipsilateral autonomic symptoms Duration 15 min to 3 hr, recurrent periodically Response with triptan therapy
Medication overuse	<ul style="list-style-type: none"> Develops or worsens with daily medication use Similar pattern to chronic, episodic or tension type headaches

- A. Migraine with out aura
- B. Cluster Verapamil 02,100%
- C. Tension headache

Types of headache	Migraine	Cluster	Tension
Sex predilection	Female > male	Male > female	Female > male
Family history	Often present	No	No
Onset	Variable	During sleep	Under stress
Location	Often unilateral	Behind one eye	Band-like pattern around the head (bilateral)
Character	Pulsatile & throbbing	Excruciating, sharp & steady	Dull, tight & persistent
Duration	4-72 hours	15-90 minutes	30 minutes to 7 days
Associated symptoms	Auras, photophobia, phonophobia & nausea	Sweating, facial flushing, nasal congestion, lacrimation & miosis	Muscle tenderness in the head, neck, or shoulders

11. patient with moter vehicle accident got unconscious at the than got up having headache than Conditions further Detroit with Ipsilateral blow pupil and contralateral hemipersis dx asked

- A. Epidural
- B. Subdural
- C. Subarachnoid

Epidural hematoma	
Pathogenesis	<ul style="list-style-type: none"> Often in temporal area with tearing of middle meningeal artery Onset of consciousness followed by lucid interval Characteristics expansion leads to: <ul style="list-style-type: none"> Intracranial pressure (impairment consciousness, headache, pupilloconstriction) Unilateral hemiparesis Unilateral pupillary dilation & contralateral optic atrophy
Clinical features	<ul style="list-style-type: none"> Headache Unilateral pupillary dilation Unilateral hemiparesis Unilateral optic atrophy
Diagnosis	<ul style="list-style-type: none"> Head CT: dense, biconvex (lens-shaped) hyperdense that does not cross suture lines Signet ring sign They often occur with fracture in the parietal region (the junction of the frontal, parietal, temporal, and occipital bones) and most commonly affect young adults
Treatment	<ul style="list-style-type: none"> Craniotomy and craniectomy (craniectomy) for symptomatic patients

Subdural hematoma	
Pathogenesis	<ul style="list-style-type: none"> Expansion of bridging veins (tear tearing)
Clinical features	<ul style="list-style-type: none"> Often a crescent (crescent-shaped) 1-1.5 cm Unilateral hemiparesis Unilateral pupillary dilation Unilateral optic atrophy
Diagnosis	<ul style="list-style-type: none"> Head CT: dense, crescentic (crescent-shaped) hyperdense that crosses suture lines Signet ring sign They often occur with fracture in the parietal region (the junction of the frontal, parietal, temporal, and occipital bones) and most commonly affect young adults
Treatment	<ul style="list-style-type: none"> Craniotomy and craniectomy (craniectomy) for symptomatic patients

12. Dermatomyocyte question with clear picture of heliotrope rash wat investigation will you do for diagnosis

- A. Skin biopsy
- B. Muscle biopsy

EXERCISE QUESTION

Question: A 45-year-old male patient presents with a heliotrope rash on his eyelids and a Gottron's sign on his fingers. What is the most likely diagnosis?

Options:

1. Dermatomyositis
2. Systemic sclerosis
3. Scleroderma
4. Mixed connective tissue disease
5. Sjogren's syndrome

Answer: 1. Dermatomyositis

Explanation: The patient's presentation is characteristic of dermatomyositis, a systemic autoimmune disease. The heliotrope rash on the eyelids and Gottron's sign on the fingers are classic cutaneous manifestations of this condition.

Question: A 60-year-old male patient presents with a heliotrope rash on his eyelids and a Gottron's sign on his fingers. What is the most likely diagnosis?

Options:

1. Dermatomyositis
2. Systemic sclerosis
3. Scleroderma
4. Mixed connective tissue disease
5. Sjogren's syndrome

Answer: 1. Dermatomyositis

Explanation: The patient's presentation is characteristic of dermatomyositis, a systemic autoimmune disease. The heliotrope rash on the eyelids and Gottron's sign on the fingers are classic cutaneous manifestations of this condition.

13. 4 year child on routine examination abdominal mass on physical exam ct pic given with bid renal mass but respecting mid line was not cross mid line according to me normotensive dx asked

- A. Wilmos tumor
- B. Neuroblastoma **Nephroblastoma**

14. Patient heavy alcoholic asked which marker will be deranged

2 to 3 line question

- A. Alt
- B. Ast
- C. Ggt.

15. 36 year old girl with family history of breast cancer

underwent breast surgery due to breast cancer 2 to 3 months ago comes to doctor pre pregnancy counseling can I get pregnant.

- A. yes you go with pregnancy
- B. wait for 4 year than go for pregnancy
- C. you can not go with pregnancy.



Breast cancer: pregnancy
Diagnosis pachie wait 2 yrs
Chemo pachie wait 8 month
Tamoxifen cessation for 3 mnth
Transtuzumab cessation for 7 months

16. Patient with heart sounds on right side ct given, history of

infection but CFTR mutation is negative what complication can he develop I understand this case as **kartagener syndrome**.

- A. infertility
- B. mesenteric ischemia

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17. Baby 8 weeks year old with history of Nonbilious vomiting and after vomiting feeling hungry asked diagnosis

A. Pyloric stenosis

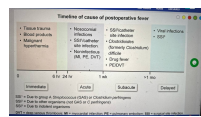
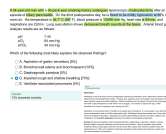
18 patient underwent some surgery 3 to 4 days ago now having

Inc RR pulse rate Tachypnea what will you do.

Chest radiograph is frequently abnormal but has poor sensitivity and specificity for pulmonary embolism, and may be normal. Findings may include atelectasis, infiltrates, pleural effusions, Westermark's sign (peripheral hyperlucency due to oligemia), Hampton's hump (peripheral wedge of lung opacity due to pulmonary infarction on xray), and Fleischner sign (enlarged pulmonary artery). Chest CT scan showing a wedge-shaped infarction is virtually pathognomonic for pulmonary embolism.

Pulmonary embolism A. Ct Spiral ct

B. Xray



Atelectasis ko lagi BP low dinxa

Diagnosis garum Pe - CT jam, atelectasis Vaya x-ray jam

19. Patient work in coal Furness were the burn coal and made something of marble what Organ is he at most like risk

A. Lung

B. Panncrea

C. Bladder Agent orange ,benzene factory dye factory

20 Abstract: Chronic Back pain yoga

GFR Japan

Long scenario,, 32yr female,, at last mentioned her ASCVD score was 2%, what to do?

a. Life style modification

b. Aspirin

c. Ezetimibe

Indication	Recommendation
Established ASCVD	High-intensity statin therapy (LDL-C reduction ≥50%)
Primary prevention	Intermediate-intensity statin therapy (LDL-C reduction ≥30-49%)

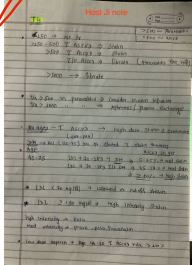
1. All ASCVD patients should be treated to reach a target LDL-C through application of one of the following strategies:

2. High-intensity statin therapy is recommended for patients with ASCVD and LDL-C ≥190 mg/dL. The target LDL-C is <70 mg/dL. High-intensity statin therapy is also recommended for patients with ASCVD and LDL-C ≥100 mg/dL who are at high risk for ASCVD.

3. The target LDL-C for patients with ASCVD and LDL-C <100 mg/dL who are at high risk for ASCVD is <70 mg/dL. For patients with ASCVD and LDL-C <100 mg/dL who are at low risk for ASCVD, the target LDL-C is <130 mg/dL.

Indications for statin therapy in prevention of ASCVD	
Secondary prevention	<ul style="list-style-type: none"> Established ASCVD <ul style="list-style-type: none"> Acute coronary syndrome Stable angina Arterial revascularization (eg, CABG) Stroke, TIA, PAD
Primary prevention	<ul style="list-style-type: none"> LDL ≥190 mg/dL Age ≥40 with diabetes mellitus Estimated 10-year risk of ASCVD ≥7.5%-10%

ASCVD = atherosclerotic cardiovascular disease; CABG = coronary artery bypass grafting; PAD = peripheral artery disease; TIA = transient ischemic attack.



Age group	ASCVD risk	ASCVD risk	ASCVD risk
40-75 years	At least one cardiovascular disease (ASCVD) risk factor	High intensity statin	High intensity statin
40-75 years	At least one cardiovascular disease (ASCVD) risk factor	High intensity statin	High intensity statin
40-75 years	At least one cardiovascular disease (ASCVD) risk factor	High intensity statin	High intensity statin
40-75 years	At least one cardiovascular disease (ASCVD) risk factor	High intensity statin	High intensity statin

2018 American Heart Association/American College of Cardiology Primary Prevention Guidelines for Management of Hypertension

2018 AHA/ACC Secondary Prevention Guidelines for Management of Atherosclerotic Cardiovascular Disease

2018 ACC/AHA Primary Prevention Guidelines for Management of Atherosclerotic Cardiovascular Disease

2018 ACC/AHA Secondary Prevention Guidelines for Management of Atherosclerotic Cardiovascular Disease

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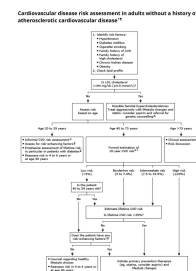
2018 ACC/AHA Primary Prevention Guidelines for Management of Atherosclerotic Cardiovascular Disease

2018 ACC/AHA Secondary Prevention Guidelines for Management of Atherosclerotic Cardiovascular Disease

Another similar one,, around 72 yr,, he or she needs 10% jasto aayo,, similar option

- a. Life style modification
- b. Aspirin
- c. Ezetimibe
- d. Gemfibrozi

ASCVD risk	ASCVD risk
Age 40-75 with diabetes	High intensity statin
Age 40-75 with diabetes	High intensity statin



21. Young female, chronic back pain,, progressive,, pain scale 7-8 out of 10,, she have huge breast F size mentioned in CV,, pain medications not helping that much.. Pq

A. Breast reduction mammoplasty...

B. Not mentioned about supportive bras or other...

22. 47yr female Colon cancer,,, surgery done, biopsy positive for high grade microsatellite instability and MSH-2.. what else advice Pq

A. Prophylactic hysterectomy

Lynch CEOS Skin

B. Screen for pancreatic Ca

C. Screen for prostatic Ca

23. History of urti 1week ago now presents with continuous vertigo, Tinnitus. Due to?

A. Bppv Dix-Hallpike maneuver causes nystagmus

B. Vestibular neuritis

C. Schwanoma

Vestibular neuritis (VN)
VN is a self-limited disorder of the vestibulocochlear nerve (CN VIII) that sometimes follows a viral upper respiratory infection. VN can be associated with significant nausea and vomiting as well as impaired gait, with the patient falling toward the affected side. VN associated with unilateral hearing loss is termed **labyrinthitis**.
The diagnosis of VN is based on clinical findings.
Head thrust test is usually **normal**. In this test, the patient is asked to look at a fixed target while the head is rapidly rotated. Patients with normal vestibular function maintain visual fixation; however, in patients with VN, the eyes move away and then return to the target with a horizontal saccade.
Because VN is usually self-limited, patients can be managed expectantly. Management

- Viral prodrome
- Mycocarditis
- Vestibular neuritis
- Iga nephropathy
- Transient synovitis

Dizziness from inner ear conditions		
Labyrinthitis	VS	Meniere's disease
<ul style="list-style-type: none"> • Constant vertigo • Erratic eye movement • Hearing loss • Tinnitus • Often a recent history of respiratory infection 		<ul style="list-style-type: none"> • Episodic vertigo • Erratic eye movement less likely • Hearing loss • Tinnitus • Not linked to infection
	VS	BPPV
		<ul style="list-style-type: none"> • Episodic vertigo • Erratic eye movement • No hearing loss • No tinnitus • Not linked to infection

24. A physician A with maniac episode? In the viral illness pandemic (in emergency department). What would to do the physician A Send physician A to psychiatric department?.

Don't allow him to treat pt
And
Psychiatric hold wala thyo

25. 40 y.o man with gastric and duodenal ulcers, and a mass in pancreas. What other parameters you have to measure?

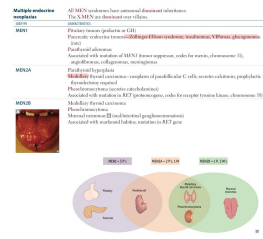
A. Prolactin,

B. PTH, 3pMEN 1

Men 1-prolactin/PTH
Men 2-calcitonin

C. calcium

Classification of multiple endocrine neoplasia	
Type 1	<ul style="list-style-type: none"> Primary hyperparathyroidism (parathyroid adenomas or hyperplasia) Pituitary tumors (prolactin, visual defects) Pancreatic tumors (especially gastrinomas)
Type 2A	<ul style="list-style-type: none"> Medullary thyroid cancer (calcitonin) Pheochromocytoma Primary hyperparathyroidism (parathyroid hyperplasia)
Type 2B	<ul style="list-style-type: none"> Medullary thyroid cancer (calcitonin) Pheochromocytoma Mucosal neuromas/marfanoid habitus



26. Female, 30s History of intubation. Removed awhile back now presents with inspiratory stridor. No other symptoms.

A. Tracheomalacia Expiratory stridore

B. Tracheal stenosis Prolonged intubation

C. Epiglottitis

In patients with a more collapsible intrathoracic airway (eg. tracheomalacia), the decreased pressure widens the intrathoracic tracheal airway. In contrast, expiration increases intrathoracic pressure. In patients with tracheomalacia, the increased pressure narrows the intrathoracic tracheal airway, leading to expiratory stridor. Tracheal stenosis (ie, rigid narrowing) or tracheomalacia (ie, weakness and collapsibility) are complications of prolonged endotracheal intubation (>2 weeks), and can cause dyspnea and noisy breathing.

27. A study conducted to see the efficacy of ear drop. Randomisation done. Before study is begin run in analysis is done, subject where give dyed ear drops and only those with coloures eardrum were chosen. What did run in do?

A. Decrease confounding

B. Decrease generalizability

esma randomize garexa tara feri euta particular arm matra analyse garya xa

C. Increase confounding

D. Increase generalizability

28. Unilateral massive pleural effusion after trauma x ray given

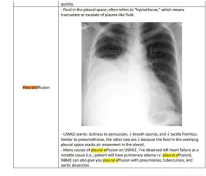
Thoracostomy

a.tube thoracostomy vs b.thoracocentesis

BOX 10-2 Indications for Surgery after Tube Thoracostomy Based on Results of Thoracostomy

Massive hemothorax, >1000- to 1500-mL initial drainage
 Continued bleeding
 >300 to 500 mL in the first hour
 >200 mL/hr for the first 3 or more hours
 Increasing size of the hemothorax on a chest film
 Persistent hemothorax after two functioning tubes are placed
 Clotted hemothorax
 Large air leak preventing effective ventilation
 Persistent air leak after placement of a second tube or inability to fully expand the lung

This is meant to be a guide, and clinical judgment should always be used.



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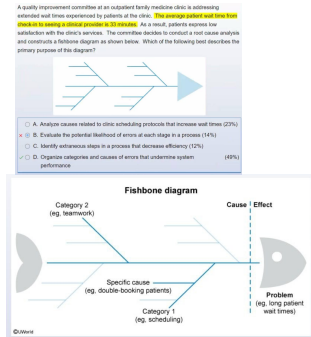
...the procedure is performed under sterile conditions. The patient is positioned in the lateral decubitus position with the affected side up. The tube thoracostomy tube is inserted into the intercostal space, and the chest is aspirated. The tube is secured to the chest wall, and the patient is repositioned to the upright position. The tube is connected to a water seal or a suction system. The patient is monitored for signs of complications, such as bleeding, infection, or respiratory distress.

Trypanosoma cruzi Chags dz

- Hopi of Trapimosigoide.----- Dilated cardiomyopathy

- Hopi Kawasaki disease---- IVIG-aspirin

Pathophysiology & etiology	Diagnosis criteria	Laboratory findings	Treatment	Complications
<ul style="list-style-type: none"> Multi-systemic reaction Onset after age 5 2 weeks to 1-2 months 	<ul style="list-style-type: none"> Fever of 38°C or higher Coronary arterial aneurysms Myocardial infarction Coronary artery stenosis Coronary artery calcification Coronary artery aneurysms Coronary artery dissection 	<ul style="list-style-type: none"> Plasma & WBCs ESR CRP ECG ECG ECG ECG 	<ul style="list-style-type: none"> Aspirin IVIG 	<ul style="list-style-type: none"> Coronary artery aneurysms Myocardial infarction Coronary artery stenosis Coronary artery dissection



- Some error, next step: design fishbone diagram

- Classic cluster headache Rx asked: verapamil sumatriptan

Acute treatment [148171]

- Sumatriptan
- Verapamil
- Other

- Diverticulitis, no improvement - repeat ct

- HA1c 6.2 nbs : repeat in 1 yr (prediabetes should repeat annually)if less than 5.7 then repeat Hba1c in 3 yr

Health maintenance in diabetes mellitus

Diabetic control:

- HbA1c < 7% (individualize target)
- Time in range (TIR) > 70%
- Time below range (TBR) < 4%
- Time at goal (TAG) > 50%

Retinopathy screening:

- Annual dilated retinal examination
- Asymptomatic patients: dilated retinal examination
- Diabetic retinopathy: dilated retinal examination

Neuropathy screening:

- Annual monofilament testing
- Annual 10-g monofilament testing
- Annual 128-Hz tuning fork testing

Cardiovascular:

- Annual lipid panel
- Annual blood pressure
- Annual ECG

Foot inspection:

- Annual foot inspection
- Annual foot inspection
- Annual foot inspection

Diabetes

- >100 to <120 fasting glucose = recheck yearly
- HbA1c < 7% recheck every 3 years, 7.5-8.4 = yearly
- diabetic neuropathy = annually
- DM retinopathy = every 1-3 years (begin at the time of dx for DM2, 5 years after DM1)
- Age > 65 with DM2 > 25 kg/m²

- Ectopic orthostatic hypotension: operative

so laparotomy garnu paryo

- Symptoms of celiac, Nbs: serum antibodies

- Small cell lung cancer, synaptophysin positive

Small blue cell

Small cell lung cancer- synaptophysin, enolase, chromogranin

- Primary enuresis in a 7 yr old: alarm

	Primary	Secondary
Diagnosis	Diagnosis of primary enuresis is based on the presence of at least one wet night per week for at least 3 months.	Diagnosis of secondary enuresis is based on the presence of at least one wet night per week for at least 3 months after a period of at least 6 months of dryness.
Causes	Genetic, developmental, and psychological factors.	Psychological, medical, and hormonal factors.
Treatment	Behavioral therapy, alarm, and medication.	Behavioral therapy, alarm, and medication.

Small cell lung cancer

- Small cell lung cancer is a highly aggressive form of lung cancer that is characterized by its rapid growth and early metastasis. It is often diagnosed at an advanced stage and has a poor prognosis. Treatment typically involves a combination of chemotherapy and radiation therapy.

- FEMALE taking penicillin, sulpha drug what is the cause of hemolysis?

penicilline le AIHA ko kura huna sakcha female ma sulpha nahola

Autoimmune hemolytic anemia

A normocytic anemia that is usually idiopathic and Coombs (+). Two types:

- Warm AIHA - chronic anemia in which primarily IgG causes extravascular hemolysis. Seen in SLE and CLL and with certain drugs (eg, Bactrim, d-methylphen).
- Cold AIHA - acute anemia in which primarily IgM + complement cause RBC agglutination and extravascular hemolysis upon exposure to cold = painful, blue fingers and toes. Seen in CLL, Mycoplasma pneumoniae infections, infectious mononucleosis.

Spherocytes and agglutinated RBCs in peripheral blood smear.

Warm AIHA treatment: steroids, rituximab, splenectomy (if refractory).

Cold AIHA treatment: avoid avoidance, rituximab.

29. Cervical dilation 2 cm, effaced 50% contractions duration 30 seconds occur every 5 mins at 28 week (not sure) dx asked

A. premature contractions

B. premature labor, Cervical insufficiency painless dilation

C. cervical insufficiency

30. Someone with crohns did ileocelectomy some long time ago now presenting with chronic watery diarrhea, Rx asked

cholysteramine vs crohns drugs.

Common causes of Heterotopia	
Pancreatic insufficiency	<ul style="list-style-type: none">• Chronic pancreatitis due to alcohol abuse, cystic fibrosis, or autoimmune/hemolytic pancreatitis• Pancreatic cancer
Diarrhea	<ul style="list-style-type: none">• Irradiation Colitis disease• Bacterial overgrowth• Primary biliary cirrhosis• Primary sclerosing cholangitis• Surgical resection of ileum (at least 50-100 cm)
Impaired intestinal surface epithelium	<ul style="list-style-type: none">• Celiac disease• AGS enteropathy• Giardiasis
Other rare causes	<ul style="list-style-type: none">• Whipple disease• Zollinger-Ellison syndrome• Medication induced

31. A study conducted to see the efficacy of ear drop. Randomisation done. Before study is begin run in analysis is done, subject where give dyed ear drops and only those with coloures eardrum were chosen. What did run in do?

A. Decrease confounding

B. Decrease generalizability

C. Increase confounding

D. Increase generalizability

32. A 30 year male presented with whitish lesion in mouth. History of asthma controlled under ics and arbuterol. What to do for diagnosis?

A. Biopsy

B. KOH mount Candida

C. Gramstain

D. HIV testing

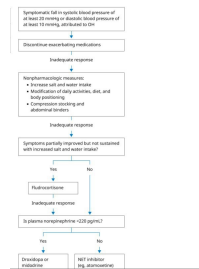
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33. 70year man had a episode of **dizziness after abruptly standing up**. His bp reading while sitting 130/90, standing 100/70. Later he was advised to drink plenty of water and stand slowly. Nbs?

A. Dexamethasone

B. Prednisolone Hydrocortisone

C. Nothing



34. A week old child presented to clinic, **he has smooth philtrum, thin lips**. His mother didn't have routine care during pregnancy. During examination child **has murmur**. What's the most likely cause?

A-VSD Fetal alcohol syndrome

B. PDA

C.TOF

35. A 18yr rugby player is tackled in the field. He was tackled by his neck and shoulder. **He had tingling sensation in right arm for 30mins**, mild head ache for **10mins**. He **didn't lose consciousness**. He has history of being tackled 4 weeks back. Diagnosis?

A. Cervical strain

B. Concussion

C. SDH

36. 42yr female **has completed her family with 2 children** and wants a reliable contraceptive method as she **doesn't want more children**. She has chlamydia. Advice?

A. Hysterectomy

B. Tubal ligation

C. OCP

d. Diaphragm

Post amputation pain

Post-amputation pain	
Acute stump pain	<ul style="list-style-type: none"> Tissue & nerve injury Severe pain lasting 1-3 weeks
Ischemic pain	<ul style="list-style-type: none"> Swelling, skin discoloration Wound breakdown ↓ Transcutaneous oxygen tension
Post-traumatic neuroma	<ul style="list-style-type: none"> Weeks to months after amputation Focal tenderness, altered local sensation ↓ Pain with anesthetic injection
Phantom limb pain	<ul style="list-style-type: none"> Onset usually within 1 week Increased risk in patients with severe acute pain

37. 60yrs female complains of pain in her amputated leg and difficulty to wearing her prosthetic. On examination 3mm wound dehiscence is seen. What will you do to guide the antibiotic treatment?

- A. Blood culture
- B. Bone biopsy and**
- C. culture
- D. Nothing

38. 60yrs Goyb / female → family history of fracture mentioned → female don't have any complication → Vaccinated as per schedule → vitals stable → asked for NBS

A. DEXA Risk x a vani DEXA natra milena

B. colonoscopy

C. No need for any Interention Risk xaina vani opp C MA

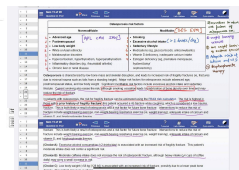
The National Osteoporosis Foundation indications for bone density testing

Women age 65 years and older and men age 70 years and older, regardless of clinical risk factors.
 Younger postmenopausal women, women in the menopausal transition, and men age 50 to 69 years with clinical risk factors for fracture.
 Adults who have a fracture after age 50 years.
 Adults with a condition (eg, rheumatoid arthritis) or taking a medication (eg, glucocorticoids in a daily dose ≥5 mg prednisone or equivalent for ≥3 months) associated with low bone mass or bone loss.

Reproduced with permission from National Osteoporosis Foundation. Clinician's Guide to Prevention and Treatment of Osteoporosis. Washington, DC: National Osteoporosis Foundation; 2013.

ANTIBIOTIC THERAPY

Whenever possible, selection of antibiotic therapy should be based on specimen for culture and sensitivity and specimens for culture are obtained from the site of infection and the site of the infection.
 In general, management of patients with infectious aetiology includes a prudent use of intravenous antibiotic therapy. (see antibiotic therapy) and a prudent use of oral antibiotic therapy (see antibiotic therapy).
 Empirical antibiotic regimens are advised for the following situations:
 1. Empirical antibiotic therapy should be initiated in patients with severe infections who are unable to undergo culture and sensitivity due to their condition.
 2. Empirical antibiotic therapy should be initiated in patients with severe infections who are unable to undergo culture and sensitivity due to their condition.
 3. Empirical antibiotic therapy should be initiated in patients with severe infections who are unable to undergo culture and sensitivity due to their condition.
 4. Empirical antibiotic therapy should be initiated in patients with severe infections who are unable to undergo culture and sensitivity due to their condition.
 5. Empirical antibiotic therapy should be initiated in patients with severe infections who are unable to undergo culture and sensitivity due to their condition.



12) 57 postmenopausal lady comes for regular health care exam. BMI 18, have family history of osteoporosis. Best screening test
 a. DEXA scan
 b. Xray of wrist
 c. Serum Ca/PO4
 13) 8 Sickle cell boy with h/o Otitis media counle week ago. Now he has pain in

39. Inferior wall MI & CV

A. from ECG.

B. pt. had H/O DVT → Presented roith SOB → No chest pain → Very long CV at lact ECG finding hinting towards infecior wall NI.

subtypes.
 Histological subtypes of melanoma [16][11]

Epidemiology

Typical sites

Clinical appearance

Growth

Lentigo maligna [17]

- Peak incidence between 65 and 80 years of age [17]
- Sun-exposed areas (e.g., face, neck)
- Darkly pigmented macule
- Irregular borders and varying size
- Gradual growth, color irregularities, surrounding island-like speckling

• Premalignant lesion

• Slow growth

• Upto 50% of untreated lesions may transform into lentigo

maligna melanoma. [17]

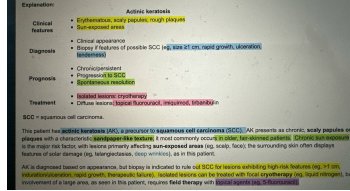
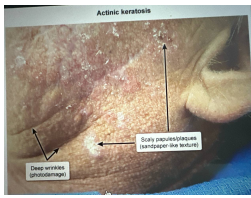
58. PPV FLIPPED ONE 40/ (40+332)

59. PPV straight forward. First q of my 1st block

60. 53 Years female had a fracture of distal radius, no hx of trauma or injury, xray was given after management of case, what will recommend to this patient?
 A. vitamin d
 B. DEXA
60+1/14 → 62.5% → 62.5% → 62.5%



Solar lentigo



40. Young Man → went on vacation during summer → presented with non-itchy, painless hyperpigmented macule → well demarcated margin. → authorji went with actinic keratosis (no pic given)

moisture vairakhne pasina ayuna
thayuma hune body part tira
hune versico



at last EEG finding hinting towards presence of...
① Young Man → Hand on vacation → presented with non-itchy, painless hyperpigmented macule during summer. Well demarcated margin. tinea
② CV of raised ICP → Papilloedema (→ NBS) → Hyperventilation
③ Mannitol
Authorji went with Actinic Keratosis (no pic given).

41. CV of raised ICP → Papilloedema + → NBS

A. Hyperventilation

B. Mannitol.

Interventions to reduce intracranial pressure	
↓ Brain parenchymal volume	• Osmotic therapy (eg, hypertonic saline, mannitol) to extract water
↓ Cerebral blood volume	• Head elevation to ↑ venous outflow • Sedation to ↓ metabolic demand • Hyperventilation to ↓ PaCO ₂ , resulting in vasoconstriction
↓ CSF volume	• CSF removal (eg, external ventricular drain)
↑ Cranial volume	• Decompressive craniectomy

CSF = cerebrospinal fluid.

42. CV of pneumothorax → resolved happens → Presented with fever (101 farh) → x rays findings → half of left lung whiteout ...asked for diagnosis

A. retained pneumoth

B. Pneumonia

C. Pneumothorax.

D. Lung abscess

43. PT ↓ treatment takes Penicilin → after a wk presented with rash. Asked what type of Hypersensitivity?

A. Type I

B. Type II

C. Type III/IV → (both in same option)

44. An old age man presented by himself with complains of forgetfulness. Recently he forgot his granddaughter birthday → Normal old age dementia.

45. Depression 3 question.

1. SIGECAPS (+) → was to diagnose.

2. Depression with psychotic feature(+).

3. Depression scenario → SIGECAPS(+) (suicidal Ideation mentioned). →

→asked for treatment. A. **CBT** B.SSRI Hospital admissions ect hola ki

46. CV of PTSD → ↓SSRI mentioned in STEM → NBS ?

.....**Nightmare**

→ **Prazosin Add**

- Trauma-focused cognitive-behavioral therapy
- Antidepressants (SSRIs, SNRIs)
- Prazosin for nightmares

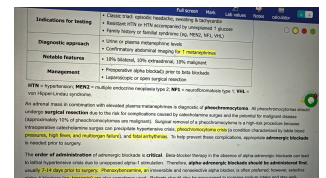
47. Roc curve given → asked about the diagnosis of disease → went with top left (most sensitive & most specific).

48. ↑ed BP | episodic Headache(+) → had family h/o smoking in father & DVT in mother, Asked NBS → VMA, no option for 5-HIAA.

Pheochromocytoma as VHL
sanga ko association ma
sodhiraxa

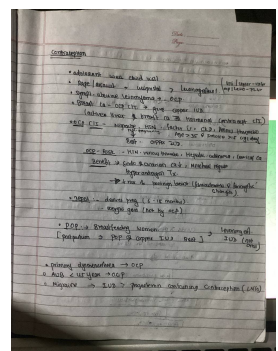
Reds of HOC
HOC malignant
HOC bilateral
HOC extracranial (eg, bladder wall, organ of Zoster) (rare)
HOC calcified
HOC lobulated

Episodic hypertensive syndrome (EHS)
Hypertension (↑ BP)
Pain (headache)
Diaphoresis
Palpitations (tachycardia)
Pallor



49. Young adolescent female raped → history of Migrane(+) → NBS details.?

- LNG.
- **Cu IUD.**



- contraceptive patch.
- contraceptive pill

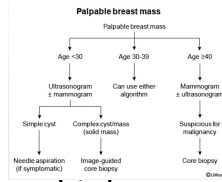
50. HOPI → fluctuant breast mass circumscribed lesion → fibroadenoma to diagnose.

Upper outer
Cyclical changes
mobile tender
huxa

51. Breast Mass + in a women 35yrs → NBS → Mammogram.

Benign breast diseases

Diagnosis	Clinical features
Breast cyst	<ul style="list-style-type: none"> • Solitary, well-circumscribed & mobile mass • ± Tenderness
Fibrocystic changes	<ul style="list-style-type: none"> • Multiple, diffuse nodulocystic masses • Cyclical premenstrual tenderness
Fibroadenoma	<ul style="list-style-type: none"> • Solitary, firm, well-circumscribed & mobile mass • Cyclical premenstrual tenderness
Fat necrosis	<ul style="list-style-type: none"> • Post-trauma/surgery • Firm, irregular mass • ± Ecchymosis, skin/nipple retraction



52. MSM → non receptive → frequent intercourse with partner of STI status unknown. → Asked for NBS.

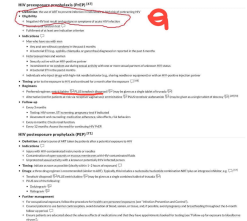
A. Antibiotic for gonorrhea

Pre maa 2, post maa 3

B. Pre exposure HIV prophylaxis.

Entricitabin, tenofovir

emtricitabine tenofovir raltegravir post



53. DMD Seenarto → NBS for diagnosis →

A. CK

B. genetic testing

C. biopsy

DMD

Duchenne Muscular Dystrophy

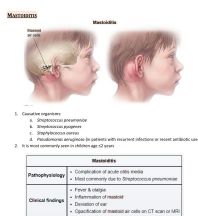
- Loss of dystrophin → **myonecrosis**
- **Creatine kinase** elevation
 - Common in early stages
 - Released from diseased muscle
- Other muscle enzymes also elevated
 - Aldolase
 - Aspartate transaminase (AST)
 - Alanine transaminase (ALT)

54. 2yr child → 20 words → other milestone achieved as per age → language delay.

55. Mastoid tenderness tnt = fever → History of barotrauma → asked for NBS

A. CT scan of Head.

B. CT scan of Head+sinus.



Acute mastoiditis

Pathophysiology	<ul style="list-style-type: none"> • Infection of the mastoid air cells • Complication of acute otitis media • Most commonly due to Streptococcus pneumoniae
Clinical findings	<ul style="list-style-type: none"> • Fever & otalgia • Inflammation of mastoid • Protrusion of the auricle • Opacification of mastoid air cells on CT scan or MRI
Management	<ul style="list-style-type: none"> • Intravenous antibiotics (ampicillin) • Drainage of purulent material (eg. myringotomy, mastoidectomy)
Complications	<ul style="list-style-type: none"> • Extracranial extension (subperiosteal abscess, facial nerve palsy, hearing loss, labyrinthitis) • Intracranial extension (brain abscess, meningitis)

56. CV of ICH → BP 185/100 mm of hg → contraindication of which of following.

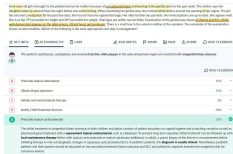
- A. Thrombolysis
- B. Thrombectomy
- C. Alteplase.
- D. Anticoagulation.

57. Lichen Sclerosus Scenario nt in child(8-9yr) → vulva finding → thin | whitish itching | → asked for treatment

A. no trt req

B. chlobetasone

Topical steroid



Vulvar lichen sclerosus	
Epidemiology	• Prepubertal girls & perimenopausal or postmenopausal women
Clinical features	• Thin, white, wrinkled skin over the labia majora/minora; atrophic changes that may extend over the perineum & around the anus • Excoriations, erosions, fissures from severe pruritus • Dysuria, dyspareunia, painful defecation
Workup	• Punch biopsy of adult-onset lesions to exclude malignancy
Treatment	• Superpotent corticosteroid ointment

58. KLCO RA → cv asked which of following value is increased?

A. DLCO

B. FEV₁/fvc

Differential diagnosis based on DLCO			
	Obstructive pattern (FEV ₁ /FVC <80% predicted)	Restrictive pattern (FEV ₁ /FVC >80% predicted & FVC <80% predicted)	Normal spirometry
Low DLCO	• Emphysema	• Interstitial lung diseases • Sarcoidosis • Asbestosis • Heart failure	• Anemia • Pulmonary embolism • Pulmonary hypertension • Pulmonary edema
Normal DLCO	• Chronic bronchitis	• Musculoskeletal deformity • Asthma	• Pulmonary hemorrhage • Polycythemia
Increased DLCO	• Asthma	• Morbid obesity	

DLCO is raised in pulmonary or alveolar hemorrhage (co. pulmonary vasculitis), an abnormal increase in haemoglobin within the alveolar space increases carbon monoxide transfer, resulting in increased DLCO.

Inc DLCO with normal Spic:
• Pregnancy - Left to right shunt

- Inc DLCO
- Asthma
- Morbid obesity
- Pul hemorrhage
- Polycythemia

Relative exclusion criteria	
• Platelets <100,000/mm ³ or glucose <50 mg/dL	• Active PFT
• Anticoagulant use with INR >1.7, PFI >15 sec, or t active PFT	
• Minor or rapidly improving neurodeficits	
• Major surgery/trauma in the past 14 days	
• Myocardial infarction in the past 3 months	
• GI or GU bleeding in the past 21 days	
• Seizure at stroke onset	
• Pregnancy	

Aspirin should be held for 24 hours following thrombolytic therapy due to the increased risk of intracranial hemorrhage. Therapeutic hypothermia is commonly used to prevent hypoxic ischemic brain injury in patients with out-of-hospital cardiac arrest.

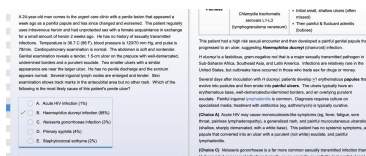
Criteria for thrombolytics in stroke	
Inclusion criteria	• Ischemic stroke with measurable neurodeficit • Symptom onset <4.5 hours before treatment initiation
Strict exclusion criteria	• Hemorrhage or multiple infarct involving >33% of cerebral hemisphere on CT scan • Stroke/brain trauma in the past 3 months • History of intracranial hemorrhage, aneurysm, or vascular malformation • Recent intracranial surgery • Active bleeding or arterial puncture in the past 7 days at noncompressible site • Blood pressure >185/110 mm Hg

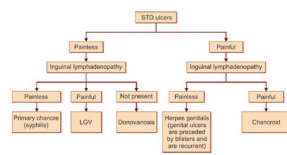
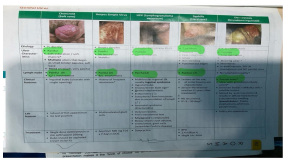
59. LVH scenario → ECG given → Where would you auscultate for this finding?

Apex ma , it 5th intercostal space ,s4 sound

60. femur # → 10cm below the hip → probably along the shaft → No xray given → asked for complication.

- A. AVN.
- B. Malunion
- C. Nonunion





Lymphogranuloma venereum—small, painless ulcers on genitals + swollen, painful inguinal lymph nodes that ulcerate (buboes). Treat with doxycycline.

Infectious genital ulcers	
Primary	<ul style="list-style-type: none"> Herpes simplex virus Chancroid (Haemophilus ducreyi) Lymphogranuloma venereum (Chlamydia trachomatis)
Secondary	<ul style="list-style-type: none"> Syphilis (Treponema pallidum) Chancroid (Haemophilus ducreyi) Lymphogranuloma venereum (Chlamydia trachomatis)

61. Painless ulcer with Painful LAD. →? diagnosis.

Lymphogranuloma Igv, azithro

62. Motor Vehicle accident → loss of consciousness → By the time he reaches hospital regained consciousness → Which initial investigation to be done ?

A. CT Scan

B. MRI brain.

63. Pt. multiple medication one of them being TMP-SMx. → presented with black skin over foot → No H/o fever → asked for its cause.

A. TMP-SMX

B. Clostridium Perfringens

Trimethoprim-sulfamethoxazole (co-trimoxazole): Drug Information
 Frequency not defined:
 Cardiovascular: Polyarteritis nodosa
 Dermatologic: Erythema multiforme, exfoliative dermatitis, skin photosensitivity, skin rash, urticaria
 Gastrointestinal: Abdominal pain, anorexia, diarrhea, glossitis, nausea, pancreatitis, stomatitis, vomiting

Tetanus	
Cause	<ul style="list-style-type: none"> Clostridium tetani spores inoculate skin wound → germinate → produce tetanospasmin → retrograde axonal transport to CNS → blocks inhibitory interneurons Risk: incomplete childhood vaccines or lack of 10-year booster shot
Symptoms	<ul style="list-style-type: none"> Trismus (lockjaw) & difficulty swallowing Intermittent intense muscular spasms Opisthotonus (sustained arched back) Risus sardonicus (facial muscle spasm while smiling)
Prevention	<ul style="list-style-type: none"> Tetanus toxoid vaccination

Black skin vanya ta necrosis nai vayo

yo same set compiled ma pani cha...paila chai tmp-smx gako

Tetanus prophylaxis	Clean or minor wound	Dirty or severe wound
≥3 tetanus toxoid doses	Tetanus toxoid-containing vaccine* only if last dose ≥10 years ago No TIG	Tetanus toxoid-containing vaccine* only if last dose ≥5 years ago No TIG
Unimmunized, uncertain, or <3 tetanus toxoid doses	Tetanus toxoid-containing vaccine* only No TIG	Tetanus toxoid-containing vaccine* PLUS TIG

64. Vit B12 → clinical feature & lab finding suggestive of Vit B12 def → what will you look initially for.

A. Vit B12

B. Methylmalonyl CoA

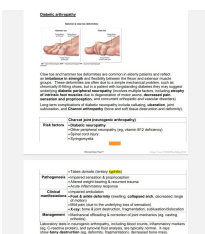
C. Homocystine

65. CV of MCAD (Hypoketotic + Hypoglycemia, mentioned in cv.) → Asked about which lab value hint towards its diagnosis? → (↑ Amonia level.)



Medium-chain acyl-CoA dehydrogenase deficiency → inability to break down fatty acids into acetyl-CoA → accumulates of fatty acid ketones in the blood with hypoketotic hypoglycemia. Causes vomiting, lethargy, seizures, coma, liver dysfunction, hyperammonemia. Can lead to sudden death in infants or children. Treat by avoiding fasting.

66. Sexually active pt → H/o unprotected sexual intercourse → presented with Painful, tingling sensation in lower limbs → x rays showed lytic lesion foot over



Syphilis manifestations	
Primary	<ul style="list-style-type: none"> Painless genital ulcer (chancres) Diffuse rash (palm & soles) Lymphadenopathy (epitrochlear)
Secondary	<ul style="list-style-type: none"> Condyloma latum Oral lesions Hepatitis
Latent	<ul style="list-style-type: none"> Asymptomatic
Tertiary	<ul style="list-style-type: none"> CNS (tabes dorsalis, dementia) Cardiovascular (aortic aneurysm/sufficiency) Cutaneous (gummas)



75. A 57-year-old woman comes to the emergency department because of increasingly severe pain and swelling of the right foot that began 2 weeks ago. She reports that the pain is worse at night. The most recently noted blood urea nitrogen level is 12 mg/dL and her weight is 110 kg (242 lb; BMI 42 kg/m²). Temperature is 101°C (38.3°C), pulse is 98/min, respiratory rate is 18/min, and blood pressure is 150/90 mm Hg. She reports an acute onset of numbness in the right foot that began 1 week ago. She reports that the numbness is worse at night. She reports that the numbness is worse at night. She reports that the numbness is worse at night. She reports that the numbness is worse at night.

Charcot arthropathy

Tabes dorsalis (also called locomotor ataxia) is a disease of the posterior columns of the spinal cord and of the dorsal roots. It has long latent period, primary infection and onset of symptoms of forms of neurosyphilis, with the interval averaging about 20 years, but sometimes as few as three years.

While tabes dorsalis was the most common form of neurosyphilis in the pre-antibiotic era, it is uncommon in the antibiotic era.

The most frequent symptoms of tabes dorsalis are sensory ataxia and lancinating pains. The latter are characterized by sudden, brief, severe stabs of pain that may affect the limbs, back, or face and that may last for minutes or days. Less common symptoms are paresthesia and gastric crises, characterized by recurrent attacks of severe epigastric pain, nausea, and vomiting. Bladder dysfunction with urinary retention and overflow incontinence may occur early in the course of disease.

Pupillary irregularities are among the most common signs in patients with tabes dorsalis, and the Argyll-Robertson pupil accounts for approximately one-half of these. An Argyll-Robertson pupil is small, does not respond to light, contracts normally to accommodation and convergence, dilates imperfectly to mydriatics, and does not dilate in response to painful stimuli.

Other findings seen with tabes dorsalis include absent lower extremity reflexes, impaired vibratory and position sensation, and, less commonly, impaired touch, pain, and optic atrophy.

calcaenium + navicular → author wenr with charcot arthropathy.(other option :tabes dorsalis)

67. Pic of Hyphema given → H/o trauma to Rt-eye during play. → NBS?

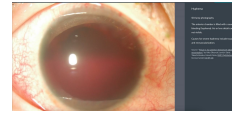
A. Measure IOP

B. Refrac. Error

C. flurosece.

Traumatic hyphema

Traumatic hyphema
• Most common form of traumatic hyphema
• Associated with blunt trauma
• Vision loss
• Pain
• Photophobia
• Elevation of fluid in the anterior chamber
• Possible (peripheral) pupillary block
• Optic nerve avulsion
• Ruptured retinal vessels
• Corneal endothelial eye
• IOP ↑
• Displaced lens
• Vitreal (blood) head
• Seizuring
• Traumatic glaucoma → optic nerve
• Vitreous hemorrhage
• Prolapsed lens
• Retinal vessel tear



A 7-year-old boy is brought to the emergency department due to an eye injury. About an hour ago, the patient was playing catchball and was hit in the face by the ball. He felt pain immediately in his left eye and passed out for 30 minutes. The appearance of the eye on examination is shown in the image below.



This patient's diagnosis increases the risk of which of the following complications?

- A. Cataract formation
- B. Corneal neovascularization
- C. Infectious keratitis
- D. Intraocular hypertension
- E. Subconjunctival hemorrhage

Traumatic hyphema Clinical features and diagnosis
• Traumatic hyphema is a common cause of acute vision loss in children and young adults.
• It is caused by blunt trauma to the eye, which can result in retinal vessel tearing and leakage of blood into the anterior chamber.
• The blood in the anterior chamber can cause a decrease in vision and pain.
• The blood in the anterior chamber can also cause a secondary glaucoma.
• The blood in the anterior chamber can also cause a corneal endothelial injury.
• The blood in the anterior chamber can also cause a vitreous hemorrhage.
• The blood in the anterior chamber can also cause a retinal detachment.
• The blood in the anterior chamber can also cause a lens dislocation.
• The blood in the anterior chamber can also cause a traumatic cataract.
• The blood in the anterior chamber can also cause a traumatic optic neuropathy.
• The blood in the anterior chamber can also cause a traumatic strabismus.
• The blood in the anterior chamber can also cause a traumatic enophthalmos.
• The blood in the anterior chamber can also cause a traumatic exophthalmos.
• The blood in the anterior chamber can also cause a traumatic eyelid injury.
• The blood in the anterior chamber can also cause a traumatic facial injury.
• The blood in the anterior chamber can also cause a traumatic skull fracture.
• The blood in the anterior chamber can also cause a traumatic intracranial hemorrhage.
• The blood in the anterior chamber can also cause a traumatic meningitis.
• The blood in the anterior chamber can also cause a traumatic encephalitis.
• The blood in the anterior chamber can also cause a traumatic stroke.
• The blood in the anterior chamber can also cause a traumatic aneurysm.
• The blood in the anterior chamber can also cause a traumatic arteriovenous malformation.
• The blood in the anterior chamber can also cause a traumatic tumor.
• The blood in the anterior chamber can also cause a traumatic infection.
• The blood in the anterior chamber can also cause a traumatic cancer.

68. RA prolong History → most common ocular symptom?

A. Anti Ureitis

B. Keratitis

C. scleritic

69. Discrepancy between Rt & Left breast. At breast (Tanner3) & left breast (tanner 1) CV asking about long term Complication

A. Malignancy

B. fibroadehoma

C. No complication

70. CV mentioned fluctuant mass with circumscribed lesion on breast.

cv asking regarding its complication.

A. fibroadenoma

B. Malignancy

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Acute compartment syndrome of the extremity	
Pathophysiology	<ul style="list-style-type: none">↑ Pressure within enclosed fascial space↓ Blood flow & tissue perfusion
Common causes	<ul style="list-style-type: none">• Distal injury, long bone fracture, malalignment• Ischemia after prolonged ischemia• Malaligned prosthesis
Clinical features	<ul style="list-style-type: none">• Early:<ul style="list-style-type: none">+ Progressive, severe pain+ Rapidly ↑ skin swelling+ ↑ Pain with passive stretching+ Paresthesia• Late:<ul style="list-style-type: none">+ Sensation+ Weakness (within hours), paralysis (late)+ Loss of pulse (uncommon)
Management	<ul style="list-style-type: none">• Compartment pressure measurement can confirm diagnosis• Emergency fasciotomy

A 65-year-old man involved in a motor vehicle collision undergoes a prolonged surgery to repair a left tibial fracture and popliteal artery injury. He is placed in a knee immobilizer and taken to the intensive care unit for close monitoring following the procedure. The next day, he has increasing pain in the left leg and is treated with intravenous fluids and morphine. Two hours later, the patient complains of severe pain that worsens with passive ankle movement and is associated with a sensation of "pins and needles" in the limb. Examination shows tense, tender swelling of the left calf. There is sensory loss between the great and second left toes. The lower extremity pulses are palpable bilaterally. Which of the following is the most important next step in management of this patient?

A. Elevate the leg and apply ice packs

B. Initiate patient-controlled analgesia

C. Obtain a Doppler ultrasound of the lower extremity

D. Obtain x-rays of the lower extremity

E. Return to the operating room for fasciotomy

Correct
80% answered correctly

Explanation:

Clinical features of compartment syndrome

- **Pain out of proportion to injury**
- **Pain ↑ on passive stretch**
- **Rapidly increasing skin swelling**
- **Paresthesia (early)**

Common

- **↑ Sensation**
- **Motor weakness (within hours)**
- **Paralysis (late)**

Uncommon

- **↓ Distal pulses (uncommon)**

This patient has increasing pain and calf tenderness after arterial injury, a presentation concerning for **compartment syndrome (CS)**. CS is due to elevated pressure within a fascial space, which prevents adequate perfusion of the tissues. It can be caused by trauma or prolonged compression of an extremity or by reperfusion after revascularization of an acutely ischemic limb. Patients typically have **severe pain** that is worsened on passive range of motion. Neurologic manifestations include **paresthesia (early)** and **sensorium and motor**

71. Pt. prolonged Immobilization → well score high → DVT diagnosed → presented with sudden calf pain → distal pulse not palpable → swollen legs, tendon → NBS?.

A. Heparin

B. fasciotomy

72. ILD finding mentioned → CV asked which of following is req for its diagnosis.

A. Fev

B. FvC

C. DLCO

(no option of fev1/fvc)

73. Huntington disease+ in dad → Similar H/o in past Pq Mother worried about her child so did genetic testing → absent in infant → Which principal did you violate

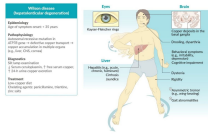
A. autonomy

B. beneficance

C. Non-Maleficance

D. social justice.

74. Clinical Scenario of Wilsons disease → (↑ed ceruloplasmin), → asked for treatment → (Penicillamine.)



Wilson disease	
Pathogenesis	<ul style="list-style-type: none"> Autosomal recessive mutation of <i>ATP7B</i> → hepatic copper accumulation → leak from damaged hepatocytes → deposits in tissues (eg. basal ganglia, cornea)
Clinical findings	<ul style="list-style-type: none"> Hepatic (acute liver failure, chronic hepatitis, cirrhosis) Neurologic (parkinsonism, gait disturbance, dysarthria) Psychiatric (depression, personality changes, psychosis)
Diagnosis	<ul style="list-style-type: none"> ↓ Ceruloplasmin → ↑ urinary copper excretion Kayser-Fleischer rings on slit-lamp examination ↑ Copper content on liver biopsy
Treatment	<ul style="list-style-type: none"> Chelators (eg. D-penicillamine, trientine) Zinc (interferes with copper absorption)

Dec ceruloplasmin
inc urinary copper

75. Cutaneous Larva Migrane → Pic given asked for trt → Albendazole / Ivermectin.

76. Catch-22 scenario given → asked for electrolyte Imbalance (hypocalcemia

mentioned

In question)

- Hyponatremia
- Hypernatremia
- Hypokalemia
- Hyperkalemia
- **Hypomagnesma**
- Hypermagnesemia

77. Red Grey Tongue , HIO ferer & cough → CV hinting towards Diphtheria. No feature of Obstruction → asked regarding its complication → → no option of myocarditis.

Clinical features
Respiratory diphtheria
<ul style="list-style-type: none"> Local features <ul style="list-style-type: none"> Tonsillar and pharyngeal diphtheria <ul style="list-style-type: none"> Greyish-white pseudomembrane over the posterior pharyngeal wall, and/or tonsils Any attempt to scrape off the pseudomembrane exposes the underlying capillaries and results in heavy bleeding Bull neck due to cervical lymphadenopathy and swelling of the soft tissue of the neck → almost obstruction Systemic features (due to dissemination of toxin) <ul style="list-style-type: none"> ↑ Gaiter <ul style="list-style-type: none"> Myocarditis Arrhythmias Reversible polyneuropathy

78. CV of cord compression given. → No History of fever → fetal Heart rate 110 bpm. → Asked for NBS (No option for resuscitation)

- Abx
- CIS
- Tocolytics
- observe

Fetal Distress
Fetal distress is a clinical diagnosis in the fetal heart rate (FHR) tracing. It is defined as a pattern of FHR that is abnormal and suggests fetal hypoxia. It is characterized by late decelerations, variable decelerations, and/or prolonged decelerations.

Normal FHR
Normal FHR is between 110 and 160 bpm. It is characterized by a regular rhythm and a consistent rate.

Early deceleration
Early deceleration is a gradual decrease in FHR that begins before, during, and after contractions. It is caused by head compression.

Variable deceleration
Variable deceleration is a sudden decrease in FHR that is unrelated to contractions. It is caused by cord compression.

Late deceleration
Late deceleration is a gradual decrease in FHR that begins after the start of a contraction and returns to baseline after the contraction ends. It is caused by uterine-placental insufficiency.

NBS
NBS is required if there is evidence of fetal distress. It is used to detect hypoglycemia, hypocalcemia, and other metabolic abnormalities.

Early deceleration - head compression
Variable deceleration - cord compression
Late deceleration - utero placenta insufficiency

Intermittent Variable deceleration - observe
Continuous - mat. positioning, amnioinfusion
if goes into fetal brady/acidosis - Cs

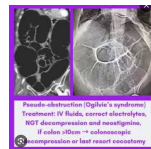
79. sharp chest pain & frequent bout of cough → X-ray shows Mediastinitis → Pt. had past H/o Pneumothorax. → asked for diagnosis.

- Esophageal rupture.
- bronchial rupture

80. Simple CV of Ogilvie syndrome → asked for NBS → look for electrolyte.

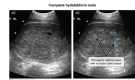
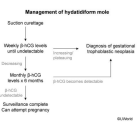
other type of obstruction (before perforation — HYPERACTIVE bowel sounds

Pseudo obstruction — HYPOACTIVE bowel sound



Intussusception	• If the Q gives you Bristol 7 stool (watery diarrhea, no solid pieces) with laxatives but Bristol 1 (severe constipation, pellet-like stools) when on nothing, answer on suspect is 'red flag' with the ans being Bristol 3-6 • Classic follow up surgery for whatever reason • Large bowel pseudo-obstruction • Think of this as less but of the large, not small, bowel. "Hurt" sounds like Hurm, which is part of small bowel. So then (it's used for refer to large bowel pseudo-obstruction). • Abdominal X-ray is what we do first in stable patients when we are looking for suspected obstruction. If patient is unstable, always go right to laparotomy. • Shows up on 2D NBE. You just need to know this is the answer for someone who has a hemoperitoneum + bloody vomiting. • The 2 Hs is because the patient is bleeding internally. • Before remember intussusception (intestine above intestine)
-----------------	--

81. Gestational size increased → H. mole (asked for diagnosis)



Etiologies	• Acute colonic pseudo-obstruction (Ogilvie syndrome) • Major surgery, traumatic injury, severe infection • Electrolyte derangement (K ⁺ , Mg ²⁺ , Ca) • Medications (eg. opiates, anticholinergics) • Neurologic disorders (eg. dementia, stroke, MS, parkinson)
Clinical findings	• Abdominal distension, pain, obstipation, vomiting • Tympanic to percussion, bowel sounds • If perforation, guarding, rigidity, rebound tenderness
Imaging	• X-ray: colonic dilation, normal haustra, nondilated small bowel • CT scan: colonic dilation without anatomic obstruction
Management	• NPO, nasogastric tube decompression • Neostigmine if no improvement within 48 hour or if the cecal diameter on imaging exceeds 12 cm (which is a strong predictor of impending perforation).

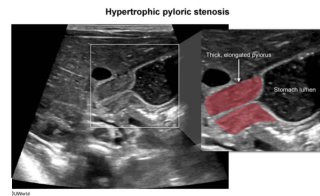
Common causes include electrolyte imbalance (eg. hypokalemia, hypomagnesemia) and factors that lead to autonomic disruption of the colon (eg. major surgery, neurologic disease, anticholinergic medication).

82. Infant 1 mth history → non-billous Vomiting → lab report awaiting → what it is the Suspected diagnosis. → H. Pyloric stenosis

hungry still (or again) after vomiting ie hungry vomiter

Pyloromyotomy
Hungry vomiter

Pathogenesis	• Thickening of pylorus → gastric outlet obstruction • Risk factors: infection, malacia, metabolic exposures
History	• Age 3-6 weeks • Projectile nonbilious emesis • Immediately postprandial • Usually followed by interest in refeeding ("hungry vomiter")
Examination	• Palpable, olive-shaped epigastric mass • Visible peristalsis • Signs of dehydration (eg. sunken fontanelle)
Laboratory	• Hypochloremia, hypokalemia, metabolic alkalosis • Increased hemoglobin • Urinalysis: thickened, stringy pylorus • Intravenous fluids (the hydration & correction of electrolyte abnormalities) • Surgical pyloromyotomy
Treatment	• Laboratory findings may be normal if diagnosed early.



83. Epiglottitis feature + → stridor + → NBS.

A. Abr intubation first if indicated then xone +vanco

B. steroid Airway edema Laryngospasm

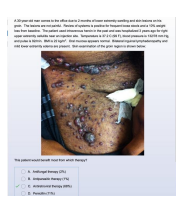
Treatment
• Endotracheal intubation (if needed)
• Intravenous antibiotics (ceftriaxone plus vancomycin)

Diagnosis is confirmed via direct visualization of an edematous epiglottis. Respiratory compromise is less common in adults but can progress quickly; therefore Patients with epiglottitis who develop rapid-onset respiratory failure (eg. tripod position, hypoxia, drooling, tachypnea) require urgent airway management. This includes bag-valve-mask ventilation with 100% oxygen followed by endotracheal intubation with advanced equipment (eg. video laryngoscope). A single failed attempt at video-assisted endotracheal intubation should prompt surgical cricothyrotomy, which bypasses the epiglottic swelling and potential obstruction.

If intubation without a video laryngoscope fails, intubation with a video laryngoscope should be attempted.

Dehydration of the uvula and unilateral lymphadenopathy can be helpful in distinguishing a peritonsillar abscess from epiglottitis.

84. Reddish-purple rash present on foot No H/o immunocompromised → Author went with Kaposi .

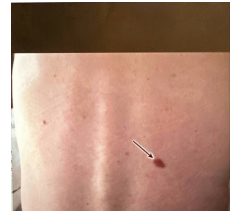
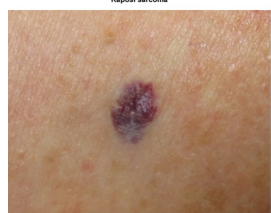


Kaposi sarcoma
Kaposi sarcoma is a type of cancer that causes abnormal blood vessels to form in the skin and other tissues. It is most common in people with weakened immune systems, such as those with HIV/AIDS. The disease is named after Dr. Moritz Kaposi, who first described it in 1872.

Signs and symptoms
• Skin lesions: The most common sign is a reddish-purple rash on the skin. These lesions can be flat or raised and may bleed or crust. They often appear on the face, hands, and feet.
• Swelling: Swelling of the hands and feet is common.
• Pain: Painful, swollen lymph nodes are common.
• Weight loss: Unexplained weight loss is common.
• Fatigue: Extreme fatigue is common.

Diagnosis
• Physical exam: A doctor will examine the skin for lesions.
• Biopsy: A small piece of skin is removed and examined under a microscope.
• Blood tests: Blood tests can help identify HIV/AIDS and other conditions that may lead to Kaposi sarcoma.

Treatment
• Antiretroviral therapy: If the patient has HIV/AIDS, treatment with antiretroviral drugs can help control the disease.
• Local therapy: For skin lesions, treatments include cryotherapy, laser therapy, and chemotherapy.
• Systemic therapy: For more advanced disease, systemic chemotherapy and targeted therapy are used.



Psychosis (eg, hallucinations, delusions, paranoia) in PD is common and may be due to the underlying disease process, medication, or a combination of the two. Two of the most common anti-Parkinson medication classes associated with psychosis are dopamine precursors (eg, levodopa) and dopamine agonists (eg, pramipexole). PD may be complicated by visual hallucinations, which can be treated with dose reduction of antiparkinsonian agents and/or a low-potency, second-generation antipsychotic (eg, quetiapine, pimavanserin).

Treatment of hallucinations in PD often begins with an initial trial adjustment of a medication regimen (eg, switching pramipexole to levodopa), but regimen changes and dose reduction are often limited due to resurgent motor symptoms. When hallucinations persist despite initial efforts, a low-dose antipsychotic may be tried. The most frequent antipsychotics used in PD are low-potency, second-generation antipsychotics (eg, quetiapine, pimavanserin).

89. Parkinson cog wheel rigidity bradykiesia 2 weeks before levodopadose increased had syntomatic releif now patient presents with visual hallucinations

NBS:

A. give dopamine

B. give quetapine,

C. reduce dose of levo and carbidopa?

Ans:decrease dose

Reduce the dose first then quetiapine

Dopamine dherai bho
Hallucinations bho
Aba reduce garne hola

Dose dec garda motor syptoms worsen hola , low dose ma anti psychotic jada thik

Systems relief cha but hallucinations cha as a side effect so Dose decrease

Parkinson disease.

Central features of idiopathic Parkinson disease

Cardinal features

- Tremor
- Rigidity
- Bradykinesia
- Postural instability

Supporting features

- Dysphagia
- Depression
- Sleep disturbance
- Autonomic dysfunction (eg, constipation, orthostatic hypotension, urinary incontinence)
- Visual hallucinations
- Delusions
- Anxiety
- Apathy
- Impaired attention
- Impaired executive function
- Impaired verbal fluency
- Impaired visuospatial ability
- Impaired memory
- Impaired social cognition
- Impaired quality of life

A pathophysiologic model of PD involves an accumulation of alpha-synuclein within the nucleus of the striatal nigral cells (SNc), leading to neuronal dysfunction and cell death. The underlying genetic process involves mutations in the SNc, but the exact mechanism of neuronal dysfunction remains unclear. It is thought that the pathophysiology of PD involves a combination of genetic and environmental factors, including exposure to pesticides and herbicides, and a loss of striatal dopamine neurons. The loss of dopamine neurons leads to the cardinal features of PD, which are tremor, rigidity, bradykinesia, and postural instability. Supporting features include dysphagia, depression, sleep disturbance, autonomic dysfunction (eg, constipation, orthostatic hypotension, urinary incontinence), visual hallucinations, delusions, anxiety, apathy, impaired attention, impaired executive function, impaired verbal fluency, impaired visuospatial ability, and impaired memory.

Pathologic features of PD

PD is characterized by the presence of neurofibrillary tangles and Lewy pathology in the SNc and other brain regions. The presence of these features is a hallmark of PD and is associated with neuronal dysfunction and cell death. The pathophysiology of PD is thought to involve a combination of genetic and environmental factors, including exposure to pesticides and herbicides, and a loss of striatal dopamine neurons. The loss of dopamine neurons leads to the cardinal features of PD, which are tremor, rigidity, bradykinesia, and postural instability. Supporting features include dysphagia, depression, sleep disturbance, autonomic dysfunction (eg, constipation, orthostatic hypotension, urinary incontinence), visual hallucinations, delusions, anxiety, apathy, impaired attention, impaired executive function, impaired verbal fluency, impaired visuospatial ability, and impaired memory.

• Used OCP electrolyte abnormalities of Na and K?

The effects of hormonal contraceptives on electrolytes has shown that it exerts severe electrolyte derangement as there were significant reductions in serum sodium and chloride concentrations and elevation in serum potassium and bicarbonate concentration Angiotensin activity increase garchan..

• OCP complications:3 questions

• Typical dementia question

• Hospital acquired deliri um question gets better in 2 to 3 weeks old pq

• One single hypoplastic polyp on colonoscopy in normal screening asymptomatic:repeat in 7 to 10 years

US multi-society task force recommendations for post-colonoscopy follow-up in average-risk adults with normal colonoscopy or adenoma*

Baseline colonoscopy finding	Recommended interval for surveillance colonoscopy	Strength of recommendation	Quality of evidence
Normal	10 years*	Strong	High
1 to 2 tubular adenomas <10 mm	7 to 10 years*	Strong	Moderate
3 to 4 tubular adenomas <10 mm	3 to 5 years	Weak	Very low
Adenoma >10 mm	3 to 5 years	Strong	Moderate
6 to 10 tubular adenomas <10 mm	3 years	Strong	High
Tubular adenoma >10 mm	3 years*	Strong	Moderate
Tubulovillous or villous adenoma	3 years*	Strong	Moderate
Adenoma with high-grade dysplasia	3 years*	Strong	Moderate
Large (>20 mm) adenoma removed by piecemeal resection	3 years*	Strong	Moderate
1 to 2 adenomas	1 year	Weak	Very low
3 to 4 adenomas	6 months	Weak	Very low

* CRC: colorectal cancer.

• 2 questions of papilledema unilateral pailledema question

?? Jhukako hola sayad

• Player American game 15 degree banda badi hand restricted left shoulder pain not improved with nasids and steriods diagnosis?:?Adhesive capsulitis,?bicep tendionitis No rotator cuff in option

Adhesive capsulitis
Not relieve by lidocaine/ steroids
Stiffness more than pain
Intact reflex
Gradual onset
Decrease passive and active rom

• Right arm pain shoulder blades pain along with numbness in middle finger: ? C5 ? C6 ? C7: Ans: C7

• Old pq osteochondroma treatment:do nothing

Lateral projection



Common causes of shoulder pain

Rotator cuff impingement or tendinopathy	<ul style="list-style-type: none"> • Pain with abduction, external rotation • Subacromial tenderness • Internal rotation of hip with positive impingement tests (eg, Neer, Hawkins)
Rotator cuff tear	<ul style="list-style-type: none"> • Similar to rotator cuff tendinopathy • Pain with lifting, carrying, or overhead reaching • Age >60
Adhesive capsulitis (frozen shoulder)	<ul style="list-style-type: none"> • Decreased passive & active range of motion • Stiffness < pain
Biceps tendinopathy or rupture	<ul style="list-style-type: none"> • Anterior shoulder pain (over bicipital groove) • Pain with lifting, carrying, or overhead reaching • Weakness (flexion, extension)
Glenohumeral osteoarthritis	<ul style="list-style-type: none"> • Uncommon & usually caused by trauma • Gradual onset of anterior or deep shoulder pain • Decreased active & passive abduction & external rotation
Acromioclavicular joint sprain	<ul style="list-style-type: none"> • Pain over AC joint • Passive shoulder abduction provokes pain

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- Round 2nd year surgery resident had alcohol smell you're intern
NBS: report to medical supervisor
- Young female goes to study in college wants to drop out from college as she can't give presentation, but speaks well with normal group friends: social anxiety? , performance anxiety
- Young female cervical cancer diagnosed wants to get pregnant later
NBS: hysterectomy? , chemo?, LEEP?
- Treatment of tardive? valbenazine and clozapine?
- Tell about medical error 3 to 4 questions.
- Swan neck deformity RA mentioned grand daughter marriage hand tremor while working best advise to improve motor function? wrist splint, occupational therapy, steroid?
- Diabetic patient pregnant what is the risk in pregnancy: diabetes before present under insulin controlled: what is the risk to the baby?
- 15 years regular follow up previously vaccinated for HPV and meningitis now what will you give: no Tdap in option HPV dine?
- Bronchiolitis infant discharged after treatment late for vaccination due to bronchiolitis NBS: give vaccine as schedule, give all, postpone, give while discharge? ans: vaccine as schedule?
- Female brings her child due to illness unable to pay won't do treatment can't take help from charity due to religion what will you do?: go through court order
- 82y f staying with her daughter who is poor give her mother rotten foods she also eats same food due to poor financial status mother mental status intact NBS: APS?

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- Round 2nd year surgery resident had alcohol smell you're intern
NBS: report to medical supervisor
- Young female cervical cancer diagnosed wants to get pregnant later
NBS: hysterectomy?, chemo?, LEEP?
- Valvular lesion ulcerative picture under ocp not like lichen planus
sclerosis what might be complications: Fistula?, infertility? Ans: Fistula?
- Sclerosis treatment clobetasone old pq
- Diabetic patient white plaque on tongue like oral thrush what would you
give: cotrimazole?, chlorhexidine mouth gargle? Ans: cotrimazole
lozenges?
- Chlamydia trachomatis young female treatment received pregnant
complication in baby what screening?: ocular herne
- Baby nose bleed no family history bleeding for 20 minutes even after
minor injury factor 8 within range what is diagnosis: VWD1? VWD2?
VWD3? Hemophilia A? hemophilia B?
- Hyperchromatic nuclei with keratin pearl
- 17 y /f bf pregnant want to do the abortion parents know risk and
benefit wants to abort child :go with procedure
- Picture: port wine stain : truncus arteriosus hearing loss ?what would be
the complication.
- ECG : cardiac tamponade : mild chest pain no Beck's triad NBS : Echo

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- DVT and pulmonary embolism : risk factor, treatment, diagnosis wells score .
- Graves disease : feeling hot tsh decrease t3 t4 increase : finding? increase t4 ? increase perioxide antibody,
- ILD:histolgy diyera diagnosis
- Right carotid bruit, numbness tingling rt hand jvp raised: sublavian stress syndrome ?venous insufficiency?
- Thoracic outlet syndrome?
- Hernia inguinal and spigelian old pq
- Somatoform disorder
- Extra pyradimal symtoms bata treatment
- Acute dystonia and tardive dyskinesia treatment
- Treatment of tardive:? valbenazine aliu.
- Dr. gives Anastrozole, and later tells not to take it. Why? No significant difference even with Anastrozole
- Calculate NNT for vasomotor system.
$$\text{NNT} = 1 / \text{YARR (Intervention-control)}$$
$$\text{NNH} = 1 / \text{YAR (exposed-unexposed)}$$
- Blinding is done. whom to be blinded to ↑ better result?

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Patient (Double blind) Data analyzer (Triple blind)✓✓

- Inhaler fluticasone for allergic rhinitis.
- Colonoscopy cutaneous manifestation
- Acute otitis media treatment asked
- Ascending Cholangitis
- Asus screening
- Mucosal neuroma present paternal uncle has thyroid cancer present with symptoms of pheochromocytoma
- Fentanyl patch for pain control in multiple myeloma
- USG for thyroid nodule
- Secondary hyperthyroidism
- Uvula deviated with Duputren contracture
- Lichen sclerosis vulval lesion
- Ruptures ectopic pregnancy
- Mdma intoxication
- Alcohol
- Bulimia nervosa
- Renal stone of oxalate risk factor hypercalcemia with oxaluria

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- Cholecystectomy answer.. plan after discharge from gb stone leading to pancreatitis
- Enteral Vancomycin for clostridium difficile infection
- Pagets disease of breast
- Infertility hysterosalpingography
- Generalised lymphadenopathy..syphilis
- Pancytopenia methotrexate
- Infective endocarditis
- Iv drug user
- Necrotising fasciitis meropenem and Vancomycin
- Venous ulcer compression
- Newborn with heart block? Cause Maternal antibodies of sle
- Roc curve.. most sensitive test for screening a disease outbreak
- History of exposure to asbestosis presented with pleural effusion.. findings
- intrapulmonary mass or pleural plaques
- Toxic megacolon treatment
- Primary prevention
- H/o Migraine contraceptive levonorgesterol IUD
- intrapulmonary mass or pleural plaques

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- Toxic megacolon treatment
- Primary prevention
- Obese pregnant women...weight gain during pregnancy
- Army men gay like symptoms when stress cut his wrist.. mgmt admit the pt Mantoux 18mm..isoniazid therapy
- 1. Ebv symptoms given.Asked when should the child should play sports
 - After fever subsides
 - after splenomegaly subside
 - Afte 4wks
- Penis wart leison given. Asked future course
 - Chronic
 - waxing n wanning
 - Recurrence
- efficient vs effective scenario
- Hospital planned for sepsis prevention / early t/t for which they made protocol which included things like
 - when pt arrives at er take vitals within 5 min
 - lab workup within 30 min.....
 - 2 other points were also there similar to further diagnosis n t/t!
- Tourette syndrome associated with adhd
- Anchoring bias
- ARR calculation
- Prazocin PTSD under sertraline with history of night mares

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- CML cbc report with splenomegaly
- Cutena Larva migrans Albendazole
- Abstract stroke
- Intravenous thrombectomy vs tPA
- Ocp vs thromboembolism
- Dexa scan 68 yr female.
- Sequential AAA screening,
- Ecg
- AF, MI
- MR, VSD, AS
- Incomplete abortion with absent fetap cardiac activity
- Complete abortion rh negative mother
- ILD answer HRCT
- Early marker for lung function assessment in ILD DLCO
- Empyema chest xray given, pt presented with fever cough sob with history of traumatic
- pneumothorax 3 weeks back.
- Allergic rhinitis fluticasone
- MVA 15 question

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- Anal fissure
- Wilson treatment penicillamine eye picture of kf ring
- History of Arcus senile
- Charcot arthropathy
- Subconjunctival hemorrhage
- Osteosarcoma xray biopsy
- ankle sprain splinting
- Bacterial meningitis
- Abscess picture I & D
- Picture genital wart treatment asked
- Chronic granulomatous disease infection risk of stap
- Tampon induced tss
- Inhaler fluticasone for allergic rhinitis

You talk about:-
Diamond ring, Gold ring, Engagement ring, etc

We talk about:-
Signet ring, Inguinal ring, Vascular ring, Schatzki ring, Wimberger's ring etc

We are not same bro !

Pictures:

1. HSV 2 genital pic vesicles, along with CV explaining pain and burning sensation, - diagnosis

Herpes simplex virus (HSV) infection
 Patients with a primary infection often have systemic symptoms (eg, fever) and develop a tender, regional lymphadenopathy. PCR routinely detects nucleic acids in genital, perianal, and orolabial lesions. Immunofluorescence and direct immunofluorescence (DIF) but no bacteria on Gram stain. Serology (eg, IgM, IgG, and IgA) is used to confirm the diagnosis. In addition, some patients may develop acute urinary retention (eg, suprapubic fullness) due to either an immune or mechanical response to viral infection within the lower urinary tract, which causes autonomic nervous system dysfunction and resultant acute urinary retention.

The appearance of genital HSV lesions can vary and mimic other disease processes as the lesions change from vesicles to ulcers. Therefore, a suspected clinical diagnosis of genital HSV requires laboratory confirmation via **viral culture or PCR testing**. Viral culture is most effective in patients with acute HSV lesions but has decreasing sensitivity as lesions heal.

Herpes simplex virus (HSV) infection typically resolves spontaneously within a week of lesion development. However, many patients will experience recurrent recurrences, particularly during the first year after primary infection. Afterward, recurrences become less frequent due to improved cell-mediated immunity. Antiviral (eg, acyclovir, valacyclovir) are used to reduce symptom duration and frequency of recurrences but do not eliminate recurrences.

2. HPV pic given- diagnosis

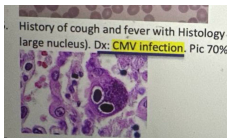
>1cm or >5 warts - Cryo /Ablation
 <1cm or <5 warts - topical .podophylline

Acuminata wart ,hiv test garni que pani cha

Genital warts (condylomata acuminata)	
Etiology	<ul style="list-style-type: none"> • HPV 6 & 11
Clinical features	<ul style="list-style-type: none"> • Multiple pink or red-colored lesions • Lesions ranging from smooth, flattened papules to exophytic cauliflower-like growths
Treatment	<ul style="list-style-type: none"> • Chemical: Podophyllin resin, trichloroacetic acid • Immunology: Imiquimod • Surgical: Cryotherapy, laser therapy, excision
Prevention	<ul style="list-style-type: none"> • Vaccination • Barrier contraception

HPV - human papillomavirus

3. CMV pic- owl eye appearance- CV explaining a renal transplant patient under immunosuppressant what t/t to be given- ans ganciclovir



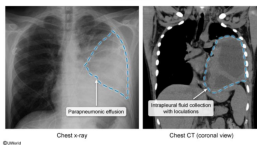
inhibits viral kinase
 Acyclovir inhibits thymidine kinase

4. Condyloma lata pic given- CV explain no pain, lesion present- is sexually active, what happens if not treated- options: recurrence, latent syphilis..

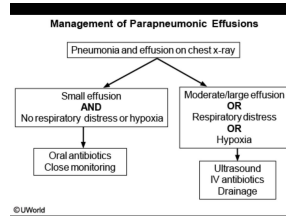


Hpv and hsv- recurrence hunxa
 Syphilis - latent ma janxa

5. Pic of empyema along with CV ... ?



Demographics/etiology	Uncomplicated	Complicated
Etiology	Staphylococcus aureus	Staphylococcus aureus
Risk factors	Small to moderate size	Large
Pathogenesis	Small to moderate size	Large
Diagnosis	Small to moderate size	Large
Management	Small to moderate size	Large



6. Venous ulcer pic given of NBME, what is the treatment? Compression therapy

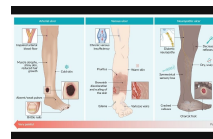
Handwritten notes on venous ulcers, including Brachial Index, arterial ulcer, diabetic ulcer, and venous ulcer locations.

Brachial Index ko value ... For pad and normal?

Arterial ulcer - Bony region

Diabetic ulcer - ventral aspect of foot

Venous ulcer - over shin above the malleolar region

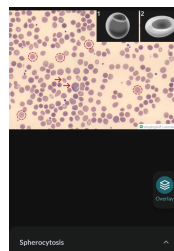


Arterial = Lateral (roughly backward pronunciation)
 Venous : Medial side

7. Peripheral blood smear given, Pic of spherocytes seen, CV explained hepatomegaly, right upper abd pain, fatigue... what will you do to prevent? Splenectomy, blood transfusion..

Hereditary spherocytosis:	
Epidemiology	<ul style="list-style-type: none"> • Usually autosomal dominant • Northern European descent
Clinical presentation	<ul style="list-style-type: none"> • Classic triad: <ul style="list-style-type: none"> • Hemolytic anemia • Jaundice • Splenomegaly
Laboratory findings	<ul style="list-style-type: none"> • ↑ MCHC (due to membrane loss and RBC dehydration) • Negative Coombs test • Spherocytes on peripheral smear • ↑ Osmotic fragility on acidified glycerol lysis test • Abnormal eosin-5-maleimide binding test
Treatment	<ul style="list-style-type: none"> • Folic acid supplementation • Blood transfusion • Splenectomy

AHA ma CLL, SLE, drugs, Drug induced Ma ni spherocyte



Spherocyte (Helmet cell) - Hemolysis - Splenectomy

Hemolysis = pigment gall stone= RUQ pain. T/t= splenectomy.

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Dacrocyte (Tear Drop) - Myelofibrosis - Trapped cell

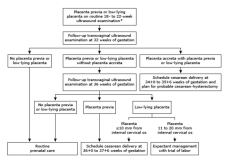
Primary myelofibrosis

Primary myelofibrosis is a condition that is very rare in childhood, with a peak incidence between the 6th-7th decades. Labs can demonstrate pancytopenia and bone marrow hypocellularity, but the condition is chronic in nature and is often accompanied by splenomegaly. A peripheral smear would demonstrate nucleated erythrocyte precursors and immature myeloid cells. This young patient has both a different clinical course and a peripheral smear.

8. Pic of tear drop given diagnosis?.. Myelofibrosis

9. Pic of placenta previa given CV expaining painless vaginal bleeding in a....week pregnant patient, definitive management will be? **C-section**, surfactant delivery, steroid to the mother..

Follow-up of placenta previa or low-lying placenta on routine 18- to 22-week ultrasound examination



The above algorithm applies to asymptomatic women. Emergency cesarean delivery is indicated for women with placental previa and active labor, severe and persistent vaginal bleeding, or a vaginal breech at 36 weeks of gestation.

Placenta previa	
Risk factors	<ul style="list-style-type: none"> Prior placenta previa Prior cesarean delivery Multiple gestation
Clinical features	<ul style="list-style-type: none"> Painless vaginal bleeding >20 weeks gestation
Diagnosis	<ul style="list-style-type: none"> Transabdominal followed by transvaginal sonogram
Management	<ul style="list-style-type: none"> No intercourse No digital cervical examination Inpatient admission for bleeding episodes

Labour ma vayeni cs

10. Pic of Achalasia cardia given, young man, with dysphasia initially **for soling later for liquids** but could drink water, regurgitation of food and even consolidation of right lower quadrant oc chest expanded, **no wt loss**. What will happen? Inc peristaltic movement, **problem in nervous system dec relaxation of sphincter..**

Mechanism vs cause vayo
Symptoms kina dekhlyako cha ques le k vanxa herera gani

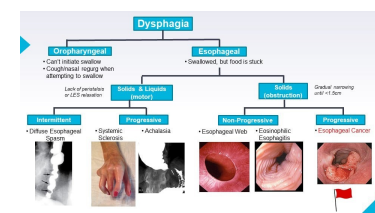
"Obstruction" of passage of the LES due to loss of NO releasing neurons in the Auerbach's plexus (myenteric plexus) in the muscularis propria
Can be observed on NBI or "Bird beak" on barium swallow with consolidation at the gastroesophageal junction.
Patients usually complain of food sticking and regurgitation after eating. This is common in liquids, which will not reach the stomach to get digested, or progressive from solids to liquids. They complain of dysphagia.
Other causes of achalasia can be caused by Chagas disease, loss of NO due to "Chagas disease", which is caused by Trypanosoma cruzi. This may be diagnosed by biopsy of the muscle, which shows a lack of "bird beak" appearance.



Achalasia Bird beak narrowing

Achalasia	
Clinical presentation	<ul style="list-style-type: none"> Chronic dysphagia to solids & liquids, regurgitation Heartburn, weight loss
Diagnosis	<ul style="list-style-type: none"> Manometry: ↑ LES resting pressure, incomplete LES relaxation, ↓ peristalsis of distal esophagus Barium esophagram: Smooth "bird-beak" narrowing at gastroesophageal junction
Management	<ul style="list-style-type: none"> Upper endoscopy to exclude malignancy Laparoscopic myotomy or pneumatic balloon dilation Botulinum toxin injection, nifedipine & CCB

CCB - calcium channel blocker; LES - lower esophageal sphincter.



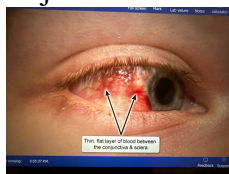
Hamle khako food lai overcome gamma ta increase prestalsists nai hune vayo... (badhne nai vayo) booz there is obstructi... relaxation decrease vayo

11. Contact dermatitis pic given, T/T? Steroid

Contact dermatitis	
Pathophysiology	<ul style="list-style-type: none"> Aggravated by hypersensitivity Friction
Triggers	<ul style="list-style-type: none"> Plants: urticarial Plants: irritant contact
Appearance	<ul style="list-style-type: none"> Pruritic Erythematous Blister Cystic Small vesicles Small blisters Bullae Large blisters



12. Subconjunctival hemorrhage pic, NBS- Tonometry, **Reassurance**

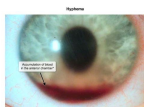


This patient with an atraumatic and **painless red eye** has a thin layer of blood on the eye surface consistent with **subconjunctival hemorrhage (SH)**. SH is a **benign condition** that occurs when a small conjunctival blood vessel ruptures, leading to **blood collecting** in the space **between the sclera** (the white part of the eye) and the **conjunctiva** (the mucous membrane covering the sclera). SH can be due to minor trauma (e.g., **rubbing the eyes vigorously**), but in many cases there is no apparent cause.

Patients with SH are typically **asymptomatic** and have a normal eye examination (eg, normal pupillary reflex, extraocular movements, and visual acuity) except for the presence of a flat and focal area of blood on the ocular surface. SH is a **clinical diagnosis** that does not require any **further work-up**. Patients can be **reassured** that **no medications** or interventions are necessary; the hemorrhage typically spreads, fades, and becomes yellow in color, and then disappears completely **within 1-3 weeks**.

13. Hypema pic given NBS- Tonometry, observe, antibiotics..

Traumatic hypema	
Mechanism	<ul style="list-style-type: none"> Blunt ocular trauma (eg, baseball, softball, deployment) Penetrating trauma (eg, gunshot)
Clinical presentation	<ul style="list-style-type: none"> Blurred vision Pain Photophobia Decreased intraocular pressure Bleeding (rougeau pupil) Optic disc hyperemia Blurred vision Decreased intraocular pressure Optic disc hyperemia Bleeding (rougeau pupil)
Initial management	<ul style="list-style-type: none"> Observation and monitoring Topical corticosteroids Topical NSAIDs (eg, ketorolac) Analgesics Antibiotics (eg, cefazolin) Pressure-lowering agents
Complications	<ul style="list-style-type: none"> Secondary angle-closure glaucoma Retinal detachment Optic atrophy Chronic uveitis Blurred vision Decreased intraocular pressure Optic disc hyperemia Bleeding (rougeau pupil)



A 19-year-old male is brought to the emergency department due to an eye injury. He reports being struck in the eye with a baseball. He has pain, blurred vision, and a large amount of blood on the surface of his eye and on the sclera. The patient is alert and oriented. The appearance of the eye is consistent with a hypema.

This patient's diagnosis increases the risk of which of the following complications?

- A. Cataract formation
- B. Corneal endothelial abrasion
- C. Intraocular hemorrhage
- D. Retinal detachment
- E. Subconjunctival hemorrhage

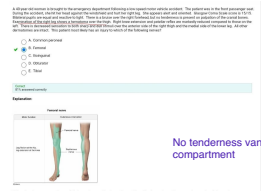
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14. Pic of hematoma on leg, what is he likely to develop? Compartment syndrome, Varicose vein, neurological deficits **Abscess**

Anteriori part of leg ma hematoma vanera airathyo ani varicose vein vanera gairathem But sathi haru abscess ma gairaxan re ?????

Q1. Trauma to skin, hematoma picture given. What will be complication?
a) Abscess
b) Varicose veins
c) DVT.

Hemorrhage/hematomas located beneath the fascia can cause compartment syndrome in the absence of trauma UTD



No tenderness vayo so no compartment

Compartment pani jana chai milxa..d/t increase pressure within the fascial compartment which may arise from hematoma

Hematoma vayo- bacterial growth- pus - cavity pressure rise - compartment - point tenderness

Sequentials:

1. Pic given of hemothorax after RTA, on auscultation, dec air entry to affected lung one, heard sound could be heard, no murmur, no JVP distention, no tracheal deviation BP dec, tachycardia..- diagnosis asked

Later after thoracocentesis 1500ml non clotted blot seen and vitals unstable NBS: chest tube, thoracotomy..

1500ml in 24hour
2 hour ma 200 ml every 1 hour
Thoracotomy

Penetrating trauma accompanied by shock (eg, severe hypotension) is attributed to hemorrhage until proven otherwise.

Although **tube thoracostomy** is often sufficient to manage hemothorax, some patients (up to 15%) require **emergent thoracotomy** for extreme bleeding, including those with:

- Initial bloody output >1,500 mL (>20 mL/kg)
- Persistent hemorrhage: >200 mL/hr for >2 hours, or continuous need for blood transfusion to maintain hemodynamic stability

Massive hemothorax^[4]

- Etiology: most commonly caused by injury to large intrathoracic vessels
- Clinical features
 - Hemorrhagic shock (e.g., need for multiple blood transfusions)
 - Chest tube output ≥ 1500 mL immediately upon placement
 - Chest tube output ≥ 200 mL/hour for 2–4 hours
 - Tracheal deviation may be present.^[4]
- Management: urgent thoracotomy

Penetrating trauma accompanied by shock (eg, severe hypotension) is attributed to hemorrhage until proven otherwise.

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- Initial bloody output >1,500 mL (>20 mL/kg)
- Persistent hemorrhage: >200 mL/hr for >2 hours, or continuous need for blood transfusion to maintain hemodynamic stability

2. RTA blood in meatus, pelvis fracture with unstable pelvis explained XRAY finding, retrograde urethrography done, no spillage seen, blood in catheter, NBS- CT abd and pelvis, Retrograde cystoscopy... No spillage observation
Retrograde cystoscopy done ..NBS?...

Retrograde urethrogram --- Retro cystogram --- Retro pyelogram or CT IVU --- CT Abdominal/Pelvis (KUB) ?

CT pelvis with IV contrast is generally insufficient for evaluation of bladder injury Spill bhacha a bhney blood tah acha kata bata ayo herna paryo

Management of bladder injury

Urine leaked from **extraperitoneal bladder rupture** stays contained within adjacent tissues (eg, retropericitoneal). Therefore, **FAST** is negative for intraperitoneal free fluid and pain typically remains localized to the pelvis (eg, suprapubic tenderness). In contrast, **intraperitoneal bladder rupture** spills urine into the peritoneal space, causing a positive **FAST** and, if undiagnosed, abdominal distention (eg, urinary ascites) and/or chemical peritonitis (Choice B). **Retrograde cystoscopy**, in which the **bladder** is passively filled (eg, via Foley catheter) with water-soluble contrast and imaged (eg, CT scan, fluoroscopy), can confirm the diagnosis.

(Choice C) Urethral injury may present with hematuria. However, it is overwhelmingly caused by iatrogenic trauma (eg, pelvic surgery) or penetrating trauma rather than blunt trauma. In addition, flank/abdominal pain (depending on the location of the injury) is more common than suprapubic fullness and tenderness.

(Choice D) Renal laceration may present with hematuria, but suprapubic fullness and tenderness are unlikely. Patients with renal laceration typically have flank pain and/or hemorrhage into the retroperitoneal space (eg, flank ecchymosis) instead.

Bladder injury

Always present with blood in the urethral meatus.

Urethral injury

- Always present with blood in the urethral meatus.
- Blood on external meatus = pathologic finding.
- Blood on external meatus = pathologic finding.

Bladder injury

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Bladder injury

- Always present with blood in the urethral meatus.
- Blood on external meatus = pathologic finding.
- Blood on external meatus = pathologic finding.

Blood in meatus
1. Retrograde urethrogram first

Blood at urethral meatus than gurgly
Urthral rupture with extravastation
- (if spillage).
suprapubic catheter and do surgical intervention
2. If spillage, not present/pass urthral catheter than hematuria do cystogram
Extravastation...is no...do observation
Extravastation of dye occur
Extraperitoneal...catheter drainage
Intraperitoneal...surgical intervention.

- a. Then do cystogram
- if spillage present
i. Extraperitoneal: catheter drainage
ii. Intraperitoneal: surgical

3. CV along with lab findings given- platelet dec, hb dec, other parameters normal- diagnosis ?- ITP TTP ma ho anemia ra tp Ttp vanera gako yo ma ma
What will be the treatment?... antibiotics given, steroid, immunosuppressants..

Immune thrombocytopenia	
Etiology	<ul style="list-style-type: none"> • Platelet autoantibodies • Preceding viral infection
Clinical findings	<ul style="list-style-type: none"> • Petechiae, ecchymosis • Mucosal bleeding (eg, epistaxis, hematuria)
Laboratory findings	<ul style="list-style-type: none"> • Isolated thrombocytopenia $<100,000/\text{mm}^3$ • Few platelets (size normal to large) on peripheral smear
Treatment	<ul style="list-style-type: none"> • Children <ul style="list-style-type: none"> ◦ Observe if cutaneous symptoms only ◦ Glucocorticoids, IVIG, or anti-D if bleeding • Adults <ul style="list-style-type: none"> ◦ Observation if cutaneous symptoms AND platelets $\geq 30,000/\text{mm}^3$ ◦ Glucocorticoids, IVIG, or anti-D if bleeding or platelets $<30,000/\text{mm}^3$

IVIg = intravenous immunoglobulin.

Anti-D immune globulin (if Rh-positive and Coombs-negative)

Thrombotic thrombocytopenic purpura	
Pathophysiology	<ul style="list-style-type: none"> • ADAMTS13 level \rightarrow uncleaved vWF multimers \rightarrow platelet trapping & activation • Acquired (autoantibody) or hereditary
Clinical features	<ul style="list-style-type: none"> • Hemolytic anemia (\uparrow LDH, \downarrow haptoglobin) with schistocytes • Thrombocytopenia (\downarrow bleeding time, normal PT/PTT) • Sometimes with: <ul style="list-style-type: none"> • Renal failure • Neurologic manifestations • Fever
Management	<ul style="list-style-type: none"> • Plasma exchange • Glucocorticoids • Rituximab

Thrombotic thrombocytopenic purpura

- ADAMTS13 level \rightarrow uncleaved vWF multimers \rightarrow platelet trapping & activation
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Clinical features

- Hemolytic anemia (\uparrow LDH, \downarrow haptoglobin) with schistocytes
- Thrombocytopenia (\downarrow bleeding time, normal PT/PTT)
- Sometimes with:
 - Renal failure
 - Neurologic manifestations
 - Fever

Management

- Plasma exchange
- Glucocorticoids
- Rituximab

LDH = lactate dehydrogenase; vWF = von Willebrand factor

The patient likely has **thrombotic thrombocytopenic purpura**, a life-threatening thrombotic microangiopathy that classically causes a pentad of manifestations:

- Thrombocytopenia
- Microangiopathic hemolytic anemia (MAHA)
- Renal insufficiency
- Neurologic changes (eg, headache, confusion, coma, stroke)
- Fever

However, most patients do not have all 5 manifestations at the time of diagnosis. Cases are most common in previously healthy individuals in middle age who are female, black, pregnant, or obese.

ITP is a diagnosis of exclusion; patients typically have a low platelet count with no other abnormalities

Empiric therapy^[4]

- Prompt initiation of **plasma exchange therapy (PEX)**
- High-dose **glucocorticoids**
 - Prednisone
 - OR methylprednisolone
- Patients with a high pretest probability of TTP (based on clinical judgment)
 - Consider early caplacizumab.
 - Consider **rituximab**.

Others:

1. A child with his parent shifts to a grandmom house built in 1980, he eats a lot of junk foods, he is fatigue and many things explained, lab findings given what could be the cause- options- lead toxicity, eating habits

Estimated population distribution of lead: A risk factor for elevated lead levels is living in a home built before 1978. However, if you have a home built after 1978, there are other factors that could be causing a child to have elevated lead levels.

Children with elevated lead levels: Children with elevated lead levels may have anemia, abdominal pain, and developmental delay. Children with elevated lead levels may also have behavioral problems.

Children (10 years old) with elevated lead levels: Children with elevated lead levels may have behavioral problems, developmental delay, and learning difficulties.

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Lead poisoning	
Risk factors	<ul style="list-style-type: none"> Home built before 1978 Flaking peeling or chipping paint Mouthing behaviors (eg, infants, developmental delay) Soiling with lead poisoning Low socioeconomic status Immigrant or international adoptee
Clinical features	<ul style="list-style-type: none"> May be asymptomatic Abdominal pain/constipation Cognitive impairment/behavioral problems Encephalopathy Obtain venous sample (if screening performed by capillary sample) Environmental surveillance (identify & remove lead sources)
Management	<ul style="list-style-type: none"> Chelation therapy if lead level >45 µg/dL Chelation therapy if lead level >45 µg/dL

How lead toxicity might be distractor
Aru milne options huncha jasto lagyo mai
Estai qtn katai thyo but lead T was both the answer

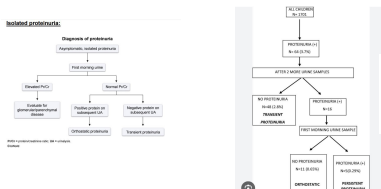
1978 samma ko lead natra aru herumla

2. ECG of Atrial fibrillation given along with pulse irregular, and patient is symptomatic, what could be the source of the ECG finding asked, options:

- Right atrial appendage
- Left atrial appendage
- Right ventricle
- Left ventricle

Flirting is always right . Flutter right
Fibrillation left

3. 1+ proteinuria in a child- ans orthostatic proteinuria



- Transient proteinuria is typically caused by fever, exercise, seizure, stress, or volume depletion and results from variations in glomerular blood flow. This condition usually resolves once the provoking factor is removed.
- Orthostatic proteinuria is most common in adolescent boys and presents with proteinuria when the patient is upright (ie, daytime) and absent urinary protein after a prolonged recumbent period (ie, morning).

Orthostatic proteinuria is diagnosed if the urinary protein excretion rate is normal for the nighttime collection (for children <4 mg/m² per hour and for adults <50 mg over an eight-hour period) and the daytime collection exceeds the normal protein excretion rate.

Orthostatic proteinuria (postural proteinuria): increased protein excretion only in the upright position

4. 2yrs child with giardiasis what is the treatment?

Giardia t/t
<1 yr = metronidazole
1-3 yrs = nitazoxanide
>3 yrs = tinidazole
Pregnancy = paramomycin

Drug	Dose	
	Adults	Children
Single of choice		
Tinidazole*	2 g orally, single dose*	Age 3-5 years: 50 mg orally, single dose (maximum dose: 3 g) Age 6-11 years: 100 mg orally 3 times per day for 3 days Age 12 years: 200 mg orally 3 times per day for 3 days
Nitazoxanide†	500 mg orally 2 times per day for 3 days	Age 3-5 years: 100 mg orally 2 times per day for 3 days Age 6-11 years: 200 mg orally 2 times per day for 3 days Age 12 years: 400 mg orally 2 times per day for 3 days
Alternative agents		
Metronidazole*	500 mg orally twice daily OR 250 mg orally 3 times per day, duration 5-10 days*	10 mg orally divided 3 times per day for 5 to 7 days (maximum 400 mg per dose)
Albendazole	400 mg orally once daily for 3 days	10 to 15 mg orally once daily for 3 days (maximum 400 mg per dose)
Ibuprofen	200 mg orally 3 times per day for 5 days	200 mg orally 3 times per day for 5 days
Paromomycin†	15 mg orally 3 times per day for 5 to 10 days	10 mg orally 3 times per day for 10 to 15 days
Quinacrine†	100 mg orally 3 times per day for 3 to 5 days	2 mg/kg orally 3 times per day for 10 to 14 days (maximum 100 mg per dose)
Quinine	100 mg orally 3 times per day for 5 days	2 mg/kg orally 3 times per day for 5 days (maximum 100 mg per dose)

Metro-nita-zole

5. ASCVD 7, father MI at age 50yrs, LDH mildly raised, what lead you to prescribe statin?

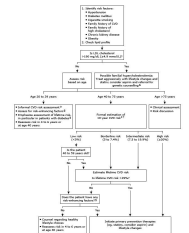
LDL more than 190 cha vannya stain gako hola

LDL more than 160 bhako bhaye dinu paryo

LDL > 190
ASCVD ≥ 7.5
age ≥ 40 with DM

Population	Recommendation	Grade
Adults aged ≥20 to 75 years who have 1 or more cardiovascular risk factors and an estimated 10-year ASCVD risk of ≥7.5% or ≥20% if a cardiovascular event of 10% or greater.	The USPSTF recommends that clinicians prescribe a statin for the primary prevention of CVD for adults aged ≥20 to 75 years who have 1 or more CVD risk factors or a cardiovascular event of 10% or greater.	B
Adults aged ≥20 to 75 years who have 1 or more cardiovascular risk factors and an estimated 10-year ASCVD risk of 5% to 7.4% or 10% to 19.9% if a cardiovascular event of 10% or greater.	The USPSTF recommends that clinicians selectively offer a statin for the primary prevention of CVD for adults aged ≥20 to 75 years who have 1 or more CVD risk factors or a cardiovascular event of 10% or greater, and an estimated 10-year ASCVD risk of 5% to 7.4% or 10% to 19.9% if a cardiovascular event of 10% or greater.	C
Adults 70 years or older.	The USPSTF concludes that the current evidence is insufficient to assess the balance of benefits and harms of initiating a statin for the primary prevention of CVD events and mortality in adults 70 years or older.	I

Cardiovascular disease risk assessment in adults without a history of atherosclerotic cardiovascular disease*



6. Atrial fibrillation, CHADVASC score 4(HTN, DM, 74 yrs,+ AF), what is NBS in management options- Aspirin, Rivaroxaban, Station, Enoxaparin

In patients with mild hypertriglyceridemia (ie, 150-500 mg/dL) who have known or are at high risk for atherosclerotic cardiovascular disease (ASCVD), initiation of a statin (eg, rosuvastatin, atorvastatin) is the recommended first-line pharmacologic therapy. Although moderate alcohol intake (no more than 2 drinks a day for men, 1 drink a day for women) may be associated with reduced cardiovascular mortality, heavy intake can significantly increase triglyceride levels.

Fibrates (eg, gemfibrozil, fenofibrate) are the most effective pharmacologic therapy for reducing triglyceride levels and are recommended in patients with severe or sometimes moderate hypertriglyceridemia (eg, 500-1000 mg/dL). Due to increased rates of adverse effects (eg, myopathy) and lack of proven cardiovascular benefit, the addition of a fibrate to a statin is rarely indicated.

Fibrates and niacin reduce triglyceride levels but have not been shown to improve cardiovascular outcomes.

Estimated 10-year risk of ASCVD
The generally accepted classification scheme for risk of cardiovascular events. Guidelines for the general population are based on the Framingham Heart Study. Risk factors include age, sex, race, total cholesterol, HDL cholesterol, systolic blood pressure, and smoking status. Risk is stratified into 3 categories: low, intermediate, and high. Risk is stratified into 3 categories: low, intermediate, and high.

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Adults 70 years or older.	The USPSTF concludes that the current evidence is insufficient to assess the balance of benefits and harms of initiating a statin for the primary prevention of CVD events and mortality in adults 70 years or older.	I

- Jelly fish sting in Hawaii, presence of tentacles, tentacles removed now what is NBS in management?
- Temp- 36.5, BP > 100 mmHg, under phenylephrine, all test done, no narcotics detected in blood, no spontaneous respiration on disconnecting the ventilator, calor test and other tests done negative. **NBS.. Declare death, Do other lab tests...**

Declaring brain death

- Declaring brain death requires all of the following:
- Establish irreversible coma and the possible cause (e.g., acute severe damage to the CNS consistent with brain death as established by clinical or radiologic evidence).
 - Brain death may not be established if one of the following is present in the patient:
 - Abnormal core temperature
 - Abnormal systolic blood pressure
 - Drug effects of CNS-depressing or neuromuscular blockade drugs
 - Abnormal laboratory values (e.g., severe electrolyte imbalance, acid-base disturbance)

- Apnea testing**
 - An essential part of the evaluation of brain death, proving the absence of respiratory-respiratory control system reflexes
 - Performed after all criteria for brain death have been met
 - The following must be ensured for safe and accurate apnea testing:
 - Normothermia
 - Normotension
 - Eucapnia
 - Eucoria
 - No evidence of hypoxia or CO₂ retention
 - After preoxygenation with 100% oxygen, the patient is disconnected from the ventilator and observed for evidence of respiratory drive (such as gasps or chest movement).
 - After 8-10 minutes, an arterial blood gas reading is obtained.
 - pCO₂ > 50 mm Hg and/or decreased pH < 7.30 when mechanical ventilation assistance is removed signifies an absence of respiratory drive, and the apnea test is considered positive.
 - If the test is inconclusive, the procedure can be repeated for a longer period of time if the patient is hemodynamically stable.

9. Polysomnography

10. Lower UTI explained, urine analysis also given, sulpha allergy, what t/t will you prescribe? **TMP/ SMX, Nitrofurantoin, Fosfomycin, Ciprofloxacin..**

Management	Options	Management
Single course (amoxicillin)	Effective compared to double or triple courses in patients with uncomplicated cystitis	First line: TMP-SMX, Nitrofurantoin, Fosfomycin
Complicated	(1) Treat with TMP-SMX (2) Treat with Ciprofloxacin	See next page
Prevention	Oral nitrofurantoin	
Prophylaxis	Document UTI (2 infections in six months or 3 UTIs in one year) Stable: Prophylactic nitrofurantoin (500mg once daily) Recurrent: Prophylactic TMP-SMX (160/800mg once daily)	Behavioral modification (post-coital voiding, stop sexual activity) Phasix, TMP-SMX or other drug can be used. Use can be daily post-coital or intermittent at treatment.

UTI in pregnancy
 Nitro pregnancy ma pari milcha, just avoid close to delivery date
 TMP smx avoid in 1st trim and after 32 weeks
 -amboss

UTI in pregnancy
 Nitro pregnancy ma pari milcha, just avoid close to delivery date
 TMP smx avoid in 1st trim and after 32 weeks
 -amboss

Nitrofurantoin jam unless these features

UTI in pregnancy

Cystitis & asymptomatic bacteriuria during pregnancy	Management
Asymptomatic bacteriuria	<ul style="list-style-type: none"> Positive urine culture (≥100,000 colony forming units/mL) in asymptomatic patient Screening usually done at 12-16 weeks gestation Treatment reduces progression to UTI & complications (eg. pyelonephritis, low birth, low birth weight)
Acute cystitis	<ul style="list-style-type: none"> Symptomatic patient (eg. dysuria, urgency) with positive urine culture Considered a complicated UTI

11. Basophil... % diagnosis. CML, CLL..

Leukemia	cytopenia	histology	cell treated	morphology	biological progression	CRF results	diagnosis
acute myeloid leukemia (AML)		myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle	cytarabine + ATRA	myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle	myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle	myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle	myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle
acute lymphoblastic leukemia (ALL)		lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle	cytarabine + vincristine + prednisone	lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle	lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle	lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle	lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle
chronic myeloid leukemia (CML)		myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle	imatinib	myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle	myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle	myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle	myeloid leukemia with 20% of blasts after chemotherapy of 1st cycle
chronic lymphocytic leukemia (CLL)		lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle	flutamide + prednisone	lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle	lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle	lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle	lymphoid leukemia with 20% of blasts after chemotherapy of 1st cycle

12. Membrane defect increase in unconjugated bilirubin diagnosis- **G6PD**

Memb defect haina ni Oxidative injury ho G6PD Def ma

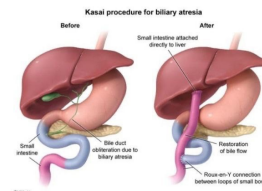
HS, G6pd

13. Neonate with incr vomiting, inc bilirubin, acholic stool, dark urine explained. What other findings will you see? Inc unconjugated bilirubin..

Biliary atresia

- One of the highest yield Peds diagnoses. Answer on USMLE for T direct bilirubin in a kid under the age of 6 weeks.
- Caused by lack of development of the intrahepatic bile ductules and biliary tree.
- Ultrasound will be done first, but USMLE wants **liver biopsy** to confirm the diagnosis.
- Kasai** procedure is done to treat, followed by liver transplant if unsuccessful.

Inc Conjugated bili



Autoimmune liver conditions

- Inflammation of bile ducts within the liver, leading to their destruction.
- Answer in a woman 20-50 who has generalized pruritis, ↑ serum cholesterol, ↑ ALP, ↑ direct bilirubin.
- USMLE loves to mention **Hx of autoimmune disease in the patient or family**

14. Congestive heart failure, Arrow BNP, Renin, Aldosterone

All inc

15. Features of pulmonary embolism in a post op case 4 hrs after OT, ECG given ST elevation in lead II, III and avF **diagnosis?.. MI**

16. CRAB features explained diagnosis? **Multiple myeloma**

17. Child with urinary incontinence failed alarm therapy, NBS in management? **Imipramine,**

**Desipramine ciprofloxacin, prednisolone
Desmopressin**

**@BED
B=behavior modification
E=enuresis alarm
D=Desmopressin**

**Secondary incontinence ma chai —
check urine (dm) / psychosocial
factors**

**<7 yr old enuresis alarm
>7 desmopressin**

Primary nocturnal enuresis	
Definition	<ul style="list-style-type: none"> Nighttime urinary incontinence age ≥5 No prior prolonged period of overnight dryness
Pathogenesis	<ul style="list-style-type: none"> Delayed maturation of bladder control 1 Nocturnal urine output (eg, 1 evening fluids, ↓ ADH) ↓ Bladder capacity
Risk factors	<ul style="list-style-type: none"> Family history Boys age 5-8
Evaluation	<ul style="list-style-type: none"> Urinalysis (to exclude other causes) Voiding diary
Management	<ul style="list-style-type: none"> Treatment of comorbid conditions (eg, constipation) Behavioral modifications (eg, restrict evening fluids) Enuresis alarm Desmopressin therapy

18. Female with features(lump in right upper outer quadrant of breast, firm, .. cm nodule) **fibroadenoma-**
what is she likely to develop? **Ductal carcinoma, intraductal papilloma, fibroadenoma mobile hunchha**

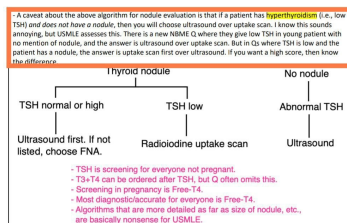
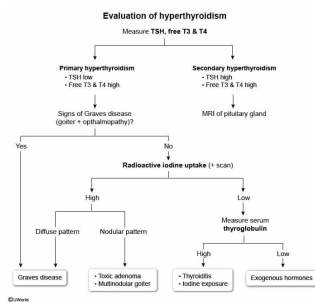
19. Female with red skin, O/E fluctuant mass on breast NBS? I and D, **Dicloxacillin,**
Ciprofloxacin

**Mastitis vayera Radioicloxacin desakyo but pachi Abscess formation
vaisakyo so -> I & D**

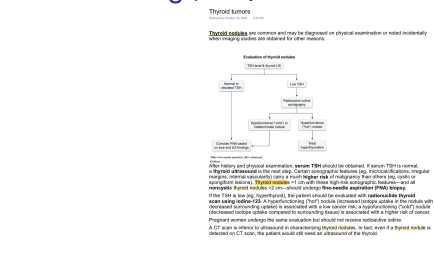
20. Thyroid nodule with dec TSH NBS? **RAIU, USG, FNAC, Propylthiouracil**

**hyper ma no nodule - USG..... hyper ma
nodule - RAIU**

Usg ,tsh paxi tsh low cha vani raiu tira jani



Toxic adenoma is the most likely diagnosis in this patient with increased serum thyroxine, low serum thyroid-stimulating hormone, and a palpable, single thyroid nodule on examination. Additionally, the patient has symptoms consistent with thyrotoxicosis, including irritability, weight loss, and diarrhea. A toxic adenoma presents as a solitary thyroid nodule accompanied by symptoms of hyperthyroidism such as heat intolerance, sweating, hair loss, warm, flushed skin, amenorrhea, diarrhea, palpitations, and weight loss. Vital signs may show hyperthermia, tachycardia, tachypnea, and hypertension. The adenoma autonomously produces triiodothyronine (T₃) and thyroxine (T₄), unregulated by the normal negative feedback mechanism. Thyroid-stimulating hormone (TSH) will be low, as it remains governed by the levels of serum T₃ and T₄. Definitive diagnosis requires evaluation with ultrasonography to identify the thyroid nodule and a radioactive iodine uptake scan, which will demonstrate a solitary hyperfunctioning ("hot") nodule. Treatment generally requires partial thyroidectomy.



21. Teenage male right and left breast unequal, tanner stage I and 3 respectively no pain on examination, no nodule or discharge. NBS. USG, Karyotype, Serum prolactin, **Reassurance**

22. 65Y/F vaginal atrophy, dyspareunia, menopause at 52 yrs, hot flashes, hysterectomy done at .. yrs d/t..fibroid? NBS? MHT, Progesterone pills, **OCPS**, IUDs Estrogen ???

Menopausal hormone therapy

In patients with no contraindications to estrogen, menopausal hormone therapy (MHT) is first-line treatment.

- Patients with an intact uterus require estrogen-plus-progesterone (MHT) (eg, estrogen-progestin pills), which decreases the risk of endometrial cancer associated with unopposed estrogen (Choice B).
- In contrast, patients without a uterus (eg, prior hysterectomy) can receive estrogen-only (eg, transdermal estrogen patch), which is preferred in these patients because estrogen-plus-progesterone (MHT) has a small increased risk of breast cancer with long-term (>3-5 years) use.

23. DMD case explained, NBS.. Aldolase Ck inc

24. Gower sign explained, muscle weakness, his other siblings normal, NBS in confirming the diagnosis? Muscle biopsy, genetic testing

25. Alcohol Use disorder t/t Naltrexone Acamprostate for maintenance

Llver c/i: naltrexone
renal c/i: acamprostate

alcohol+opiod: acamprostate
alchoh: naltrexone
Renal failure ma acamptosate contra

@ RN (Registered Nurse) will take you to LA (Los Angeles)

(R)enal impairment ma
(N)altrexone dine

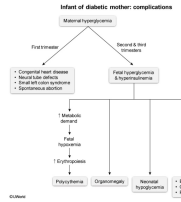
(L)iver impairment ma
(A)camprostae dine

26. AAA size 3.9 what is NBS? F/U in 2-3 yrs, surgery, F/U in 5 yrs

Maximum diameter of the abdominal aorta	Recommended follow-up interval
2.5–2.9 cm	• Repeat ultrasound after 10 years.
3–3.9 cm	• Ultrasound every 3 years
4–4.9 cm	• Ultrasound every 12 months
5.0–5.4 cm	• Ultrasound every 6 months

FEEDBACK

27. Cutaneous larva migrans T/t Albendazole (no Ivermectin in option)



28. Lady with Type I DM, pregnant, what will be the complications (Not clear in whom- fetus or mother), options; CHD, Preterm delivery, **pre-eclampsia**, Abruption placentae

29. DM mother, pregnant now what will you check? **HbA1c**, Lipid profile...

Routine Prenatal Care	
First Trimester (start care by 10 weeks gestation)	Lab Tests: CBC, blood type, Rh screen, UA and culture, Rubella, Varicella antibody screen, BPPV/VDRL, chlamydia, HIV, HBsAg, PTG, Consider HbA1c, TSH, Group factors or prophylaxis. - If MCV low AND suspicious for possible hemoglobinopathy: - Hemoglobin analysis (via HPLC or IEP) if iron. Test lead if iron carrier concerning genes. - If concerns over family history of genetic condition (Cystic Fibrosis, Tay-Sachs): - Genetically test now. Test lead if iron carrier concerning genes.
Second Trimester	- Anomaly screening (either late first or early second trimester) - Repeat alpha-fetoprotein (AFP) - Fetal anatomy US (18-22 weeks) - Diabetes screen (24-28 weeks) - 1-hour glucose challenge
Third Trimester	Rho-GAM for Rh- women (at 28 weeks), H/I, GB Culture (35-37 weeks)

Note: Visits of week (up to 28 wk), 2 week (28-34 wk), 1 week (34 wk - birth)

30. Child with splenectomy, what will you give? **Penicillin**

31. Newborn with NVD, around 10-15 lesions on mouth, uncle has also... what is he likely to develop. **MEN II B**

32. 3rd year resident is performing OT under the guidance of attending resident, 1st year resident is helping in holding the scalpel under their guidance of 3rd year resident, 1st year resident injures ureter, OT extended to correct the injury. What is the error? Decision error, Surgeon error, Surgery error, **Guidance error**

33. Cross- sectional study (what study design will you use)

34. Tolerance, dependence,..? In case of opioid

→ R had Cancer Last Stage taking short & Long Acting Opioids as needed. He asks for more, what he has developed
 (A) Tolerance (B) Dependence (C) Addiction

Table 2. Definition of terms commonly used in the literature relating to opioid dependence.

Term	Definition
Dependence	A diagnosed drug-use disorder characterised by a pattern of maladaptive behaviours, such as loss of control over use, craving and preoccupation with non-therapeutic use, and continued use despite harm. [17]
Addiction	A commonly used term to describe a condition characterised by craving and continued use despite harm. [16] Sometimes not preferred due to associated stigma. [17]
Dependence syndrome	ICD-10 diagnostic term. Describes a cluster of physiological, behavioural and cognitive phenomena in which the use of a substance takes on a much higher priority for a given individual than other behaviours that once had greater value. [18]
Substance Use Disorder	DSM-V diagnosis to describe a mild to severe state of chronically relapsing, compulsive drug taking. [18]
Tolerance	The physical adaptation of the body such that more drug is required to reach desired effect. [16]
Physical dependence	Characterised by the presence of a withdrawal syndrome on abrupt cessation or rapid dose reduction of a drug. [16]
Misuse	Use of a drug in a way other than prescribed intentionally or unintentionally. [18]
Abuse	Misuse with medical, social or legal consequences. [18]

35. Zika virus – abstinence through out pregnancy

Male 3 month
 Female 2 month

Zika virus

Women who have had unprotected sex and do not wish to become pregnant due to concerns about Zika virus infection should have ready access to emergency contraceptive services and counseling. Pregnant women should practice safer sex (including correct and consistent use of condoms or abstinence from sexual activity for at least the entire duration of pregnancy).

For couples with no active transmission of Zika virus, WHO recommends practicing safer sex or abstinence for a period of three months for men and two months for women who are returning from areas of active Zika virus transmission to prevent infection of their sex partners. Sexual partners of pregnant women living in or returning from areas where local transmission of Zika virus occurs should practice safer sex or abstinence from sexual activity throughout pregnancy.

Prevention of sexual transmission of Zika virus

WHO response

WHO network continues to conduct

36. Patient under salbutamol what will happen- Hypermagnesemia, Hypokalemia, Hypercalcemia

- B agonist hypokalemia
- B antagonist hyperkalemia

37. Splenectomy done now 4 days, what vaccine will you give? Meningococcal (no pneumococcal, Hib vaccine) other diff vaccines given

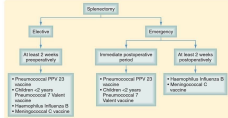


FIGURE 30-7 Splenectomy immunization algorithm flow chart. (From Hajj DJ, Iannico SJ, Mitty P, Neasegler PH. Immunization guidelines in asplenic patients. *Am J Surg* 2002;184:202-205.)

ppsv23 chai 8 weeks after PCV15

Abrasion pani hunxa body ma vanya hunxa so tt

annual flu vaccine ni dina parcha hai splenectomy ma

38. Facial nerve palsy

Prednisone dini first ma

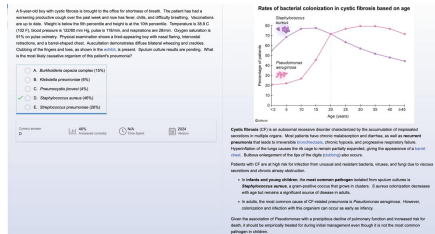
A 32-year-old man who is in the US Marine Corps is being discharged from the hospital after sustaining injuries 2 weeks ago in Southwest Asia when a rocket exploded a few yards away from him. He sustained fragmentation wounds to the right upper extremity, abdomen, and lower extremities bilaterally. At the time of injury, he received a dose of tetanus toxoid and underwent debridement and irrigation of his soft tissue wounds. Exploratory laparotomy showed a hepatic laceration and splenic laceration requiring splenectomy. He has no other history of major medical illness and takes no medications. He does not smoke. He drinks four 12-oz beers every weekend. Today, he reports no concerns. Vital signs are within normal limits. Physical examination shows a well-healing midline abdominal incision and well-healing wounds over the right upper and lower extremities. Which of the following is the most appropriate vaccination for this patient at this time?

- A. Hepatitis B
- B. Meningococcal conjugate
- C. Tetanus immune globulin
- D. Tetanus-diphtheria-acellular pertussis (Tdap)
- E. Trivalent influenza

39. Arrow of COPD- Lung complaine, lung elasticity, FEV1?

Idd

Vitamin E	Includes tocopherol, tocotrienols
FUNCTION	Antioxidant (protects RBCs and neuronal membranes from free radical damage)
DEFICIENCY	Neurologic presentation may appear similar to vitamin B ₁₂ deficiency, but without megaloblastic anemia, hypersegmented neutrophils, or ↑ serum methylmalonic acid levels. Hemolytic anemia, acanthocytosis, muscle weakness, demyelination of posterior columns (↓ proprioception and vibration sensation) and spinocerebellar tract (ataxia). Closely mimics Friedreich's ataxia.
EXCESS	Risk of enterocolitis in infants (infants) with excess of vitamin E. High-dose supplementation may alter metabolism of vitamin K – enhanced anticoagulant effects of warfarin.



Cystic fibrosis	
Pathogenesis	• Mutation (ΔF508) of CFTR gene • Autosomal recessive
Clinical features	• Recurrent sinopulmonary infections (in almost all) • Intestinal obstruction (eg, meconium ileus) • Pancreatic insufficiency & diabetes • Male infertility
Diagnosis	• Elevated sweat chloride levels • CFTR mutation on genetic testing • Abnormal nasal potential difference
Management	• Nutritional support • Airway clearance • Antibiotic coverage (Staphylococcus aureus, Pseudomonas aeruginosa)

CF should be considered in a patient with failure to thrive and recurrent respiratory symptoms (eg, frequent coughing episodes, cough with bilateral rhonchi), regardless of screening results.

Classic CFTR gene mutations are identified on newborn screening and typically cause respiratory symptoms and failure to thrive in infancy and early childhood. However, milder variants due to less common mutations may not appear on universal screening and generally have atypical presentations in adulthood. Infertility may be the only feature of atypical CF, and a male patient with congenital absence of the bilateral vas deferens should have sweat chloride and CFTR gene sequencing to confirm the diagnosis.

1. Young guy with cystic fibrosis. Is unable to perform rapid alternating movements of his hands. Labs show:

Hb: 10

MCV: 90

RDW: was slightly above the upper limit they gave cant remember exact value

Asked which vitamin was deficient = Iron, **Vit E**, B12, and a couple others.

2. Man presents with lower limb weakness and numbness. Gives history of URTI 4 weeks ago. O/E has classic transverse myelitis picture of low reflexes, weakness, loss of sensation to umbilicus level. Asked next step in diagnosis.

Options included both MRI and lumbar puncture. I went with **MRI**

3. Woman gets a hysterectomy for some reason I can't remember, but it wasn't any malignant/cancer type reason. Afterwards histology shows endometrial hyperplasia with atypia. Asked next step in her management

a. Follow with TVUS in 1 year

? b. No intervention needed

c. Cant remember the rest rip sorry

4. 2 heart murmur questions

a. First one was a downs syndrome kid, about 7 years old I think. Audio was holosystolic, I think loudest at mitral and tricuspid area, so I thought VSD. Question asked next step in his evaluation.

i. **Echo**

ii. No tests needed

iii. Some random blood tests

b. Next was a young man coming for dental procedure. Question asked about which antibiotic to give for prophylaxis. Murmur sounded like MR to me so I picked **no prophylaxis needed**.

5. Mom with complain of her kid throwing tantrums in the store when she doesn't buy him what she wants. She tries to ignore but ends up giving in because the kids an annoying little shit. You advise her to continue ignoring. Question asked what is the likely outcome of the kids tantrums if she ignores them?

Up to one-third have nasal polyps; management includes intranasal glucocorticoids (for symptomatic relief) and, in some cases, surgical resection. In addition to the respiratory features, pancreatic insufficiency leads to malabsorption, resulting in poor growth and deficiency of the fat-soluble vitamins A, D, E, and K (eg, epistaxis).

Staphylococcus aureus is the most common cause of bacterial pneumonia in young children with cystic fibrosis, especially in that with coexisting influenza infection. For patients with severe pneumonia, frequent hospitalizations, or recurrent skin infections, intravenous vancomycin should be included for empiric therapy against methicillin-resistant S aureus. Cefepime, a fourth-generation cephalosporin, provides coverage for both methicillin-resistant S aureus (MRSA) and Pseudomonas aeruginosa. Azithromycin is not used in acute CF exacerbations, but chronic maintenance therapy with azithromycin in patients with CF has been shown to slow lung function decline.

	Transverse myelitis	Guillain-Barré syndrome
Motor	• Early flaccid, late spastic paralysis • If quadriplegia, weakness in LE > UE	• Ascending paralysis • Weakness in LE > UE in early disease
Sensory	• Clearly identifiable sensory level	• Mild sensory loss • No spinal cord level
Autonomic	• Bowel & bladder dysfunction	• Cardiovascular instability
Cranial nerves	• None	• Oculomotor, glossopharyngeal, or facial paralysis
Electromyography/nerve conduction velocity	• Mostly normal	• Peripheral motor &/or sensory NCV reduced
MRI	• Focal enhanced area of T2 signal	• Normal • Enhancement of anterior nerve roots or cauda equina
Cerebrospinal fluid	• Pleocytosis • ↑ increased IgG index	• Absence of pleocytosis • Elevated protein

LE = lower extremity; NCV = nerve conduction velocity; UE = upper extremity.

Idiopathic transverse myelitis	
Pathophysiology	• Immune-mediated destruction of spinal cord • Often post-infectious (molecular mimicry)
Clinical features	• Bilateral motor weakness, classically early flaccid (LMN) progressing to late spastic (UMN) • Bilateral sensory dysfunction (all somatosensory systems) • Distinct sensory level • Autonomic dysfunction (bowel &/or bladder)
Diagnosis	• MRI of spine: no compressive lesion, T2 hyperintensity • Lumbar puncture: ↑ WBCs, ↑ IgG index
Treatment	• High-dose intravenous glucocorticoids • Plasmapheresis

LMN = lower motor neuron; UMN = upper motor neuron; WBCs = white blood cells.

Treatment of endometrial hyperplasia based on histology

	Premenopausal women	Postmenopausal women
Endometrial hyperplasia without atypia	<ul style="list-style-type: none"> • Progestin therapy: cyclic or continuous administration from the 12th to the 25th day of the menstrual cycle • In women with PCOS: administration of hormonal contraceptive 	<ul style="list-style-type: none"> • Progestin therapy
Endometrial hyperplasia with atypia	<ul style="list-style-type: none"> • All patients receiving progestin therapy should undergo: Ultrasonographic follow-up after 3-6 months • Hysterectomy with biopsy in the case of suspicious findings on ultrasonogram • Just regular surveillance without progestin therapy is indicated if progestin therapy is: <ul style="list-style-type: none"> • Contraindicated (eg, hormone-responsive cancer, progestin receptor positive breast cancer) • Not tolerated or declined by the patient 	<ul style="list-style-type: none"> • Total abdominal hysterectomy with bilateral salpingo-oophorectomy

Approach to antibiotic prophylaxis against infective endocarditis for patients undergoing dental procedures

Table 1: High-risk conditions and required devices

Patients have one or multiple high-risk features: mitral valve, bicuspid aortic valve, prosthetic mitral or aortic valve, infective endocarditis, heart failure, congenital valvular disease, or aortic aneurysm/dissection.

Table 2: Oral antibiotic regimen for prevention of endocarditis prior to dental procedures

Antibiotic	Adult dose	Child dose (if applicable)
Amoxicillin	2 g	50 mg/kg
Cephalexin	2 g	50 mg/kg
Clindamycin	600 mg	30 mg/kg
Doxycycline	200 mg	4-6 mg/kg, 2-2.5 mg/kg qd for age < 8 yr

Table 3: Oral antibiotic regimen for prevention of endocarditis prior to dental procedures

Patients with low-risk conditions and no high-risk features do not require prophylaxis.

Table 4: Ineffective endocarditis

- The following dental procedures and events do not require prophylaxis: anesthetic injections through noninfected tissue, taking dental radiographs, placement or adjustment of removable prosthodontic or orthodontic appliances, placement of orthodontic brackets, cleaning of primary teeth, and banding from trauma to lip or oral mucosa.
- Antibiotics should be administered as a single oral dose 30 to 60 minutes prior to the procedure. If inadvertently not given prior to the procedure, antibiotics may be administered up to 2 hours after the procedure. For guidance on parenteral regimens (eg, for patients unable to take oral/parenteral agents), refer to separate antibiotic content.
- A referral to UpToDate content on other measures to reduce the risk of IE.
- A Class III or class IV recommendation indicates that the benefits, risks, and costs of the intervention are uncertain. Caution should not be used for patients with history of immediate allergic IgE, angioedema, urticaria, hypotension or any severe reaction to a penicillin antibiotic.

Mitral valve disease, usually mitral valve prolapse with coexisting mitral regurgitation, is the most common valvular abnormality detected in patients with infective endocarditis. Pulmonary stenosis is not associated with IE. Antibiotic prophylaxis to prevent endocarditis is indicated prior to dental procedures in patients with high-risk cardiac conditions (eg, prosthetic heart valve). Overall, the risk of IE following dental procedures in patients with acquired valvular defects due to rheumatic fever is low, and antibiotic prophylaxis is not indicated. However, prophylaxis is indicated in patients with a history of IE.

How to respond during a temper tantrum

The following are helpful hints regarding the most appropriate ways to respond during your child's temper tantrum:

- Stay calm.
- Ignore the child until he or she is calmer. Keep doing whatever you were doing before the tantrum happens.
- Do not hit or spank your child.
- Do not give in to the tantrum. When parents give in, children learn to use inappropriate behavior to get their way.
- Do not bribe your child to stop the tantrum. The child then learns to act inappropriately to get a reward.
- Remove potentially dangerous objects from your child or your child's path.

- a. Gradually decrease
- b. Initially get worse
- c. Stay the same
- d. Continue to get worse

6. Old dementia man. Nonverbal. Lives on his own.

Neighbors help with chores and daughter comes around to help with finances. Come to clinic where man looks kinda okay I think but I remember they mentioned his lips were chapped or something. Daughter asks you to write in paper that considering her dad is non verbal, that she can sell his house and take all the money and what not. What should you do?

- a. Call the police
- b. Report to APS
- c. Agree with the daughters request
- d. Get a court appointed guardian

7. Another old 72 year dementia man, lives with his wife.

Wife complains he has trouble falling asleep and when he does, he wakes up in the night and paces the entire house, and doesn't know what time of day it is. He's able to nap easily in the day. What do you recommend?

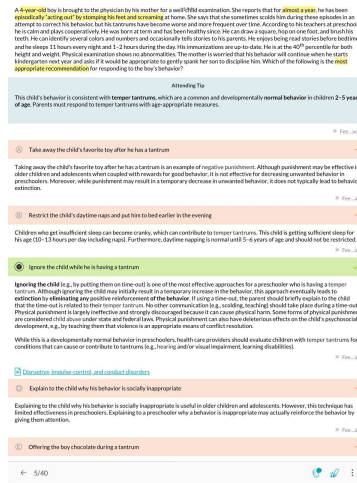
- a. A strict schedule of activities and exercise in the day and sleep at night to fix his sleep habits
- b. Advise the wife to make safe arrangements in the house considering he can get hurt walking around at night
- c. Cant remember the rest of the options soz

8. Question about an older lady whose a retired nurse and she looks after her dementia husband. Does all the chores and care for him, giving him baths, changing his catheters etc. She complains of increase fatigue and difficulty with sleep. Also says she doesn't enjoy hanging out any more with her friends/ doesn't enjoy her usual activities. O/E everything is fine, her affect is kinda flat. Labs all normal.

- a. Caregiver stress
- b. MDD
- c. Some other psych diagnoses that didn't fit at all

9. Ethics case of 2 patients with exact same name, same date of birth, same gender. Nurse A appointed to patient A, Nurse B to the other. Nurse B takes wrong IV medication bag from the pharmacy after verifying name and DOB. Patient says bruh this don't look like the usual medication drip I get. Nurse B comes running saying yo you took my patients medication wth. What can be changed to prevent

Δ: sleep walking



this next time.

- a. Cant remember the options, but the best one was about using the medical record number or **something like that on the patients arm band to verify** *(Also barcode can be used)*

10. Stats and QI questions that I can recall were

- a. Committee at a medical school take a bunch of kids with super poor clinical performance and designed a special training program for them where they would do clinical examinations under supervision, get scored, get feedback and what not. Analysis shows a significant improvement in their performance after the training program so they decide to adopt the program for all the medical students. But when they run the analysis on this new group of kids, results weren't significant. Why is that?
 - i. Lack of power
 - ii. Lack of generalizability**
 - iii. Cant remember da rest, but they didn't make sense at all
- b. QI question on some hospital director asking for there to be a record on the time it takes for every step when a patient with sepsis presents to the ER. He wanted all the time metrics up till when the patient would get the first serum lactate level measured. Whats he trying to optimize?
 - i. **Efficiency**
 - ii. Efficacy
 - iii. Equity/ Justice something like that

	Internal validity	External validity
Characteristics	Describes causality (ie, if change in independent variable causes change in dependent variable) ↑ As study becomes more tightly controlled ↓ As study becomes more like the real world	Describes generalizability (ie, if observed relationship applies to situations or people outside study) ↓ As study becomes more tightly controlled ↑ As study becomes more like the real world
Threats to validity	Bias due to: • Confounding • History • Maturation • Measurement • Regression toward the mean • Repeated testing • Selection	Bias due to: • Artificial research environment • Measurement effects • Nonrepresentative sample

Keywords: Efficacy, Effectiveness, Efficiency, Health management

ABSTRACT

Efficacy, Effectiveness, and Efficiency are widely used term in health care management. Efficacy means getting things done (is it working?), effectiveness means doing the proper things (is it actually working well?), and efficiency means doing things right (is it working within the most economical way?). It's helpful to consider them during this particular order. First, confirm the answer can actually achieve the specified result, albeit that efficacy requires very specific conditions. Then, test your solution during a real-world environment. Finally, if the answer is effective, find out ways to form it more economical more efficient. This article describes the meaning and usage of these three terminologies in context of health care setup.

Timeline > efficiency

Dimensions of health care quality	
Dimension	Description
Safety	<ul style="list-style-type: none"> Minimizes preventable errors Avoids harms from care
Effectiveness	<ul style="list-style-type: none"> Adheres to scientific guidelines evidence Avoids undertreatment & overtreatment
Patient-centeredness	<ul style="list-style-type: none"> Identifies patient values, goals & preferences Tailors care delivery to expressed patient values
Timeliness	<ul style="list-style-type: none"> Avoids delays in care, reduces wait times
Efficiency	<ul style="list-style-type: none"> Avoids wasting or overusing resources
Equity	<ul style="list-style-type: none"> Provides quality care to all individuals regardless of demographic attributes (eg, ethnicity, age, gender)

- c. Patient gets a new insurance type thing that takes a super high deductible. Your patient is dirt poor and cant afford to pay the deductible for the medications she needs. What can you do to help?
 - i. Prescribe only the generic brands
 - ii. **Arrange for a social worker to help her figure out her options**

11. Picture of what I think was Basal cell cancer. Asked next step in management.

- a. **Excision of entire lesion along with clear skin margins**
- b. Shave biopsy
 - i. I got super confused because they didn't say excisional biopsy lol I thought they were implying just remove the **whole thing and I picked shave biopsy**. Probably incorrect in **hindsight**

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→ explore & choose accordingly

12. 3-4 year old child with respiratory distress, low saturations. There's expiratory wheezing, costal retractions and what not. You start giving him some O2. CXR shows peribronchial cuffing. (basically bronchiolitis but a tad severe). Asked next step in management.

a. Listed a bunch of antibiotics and one option to just observe so I went with that

13. First trimester pregnant woman comes with complain of bleeding since she had sex with her husband. I think she was under the age of 30. She had her last pap smear 6 months ago which was normal. Examination showed a friable cervix with a giant ectropion which was bleeding. Asked next step in management.

- Get colposcopy and biopsy
- Cone biopsy
- Leave it alone
- Test for STIs

Ectropion does not need to be treated except in the rare occurrence of excessive mucous discharge or spotting that is very bothersome to the patient. In such cases, malignancy should be excluded before undertaking any treatment.

Since cases that require treatment are rare, there is no evidence to guide choice of treatment. In our experience, a two-week trial of an acidifying agent, such as boric acid suppositories 600 mg vaginally at bedtime for two weeks, may be effective. Another option is an ablative procedure using cryosurgery or electrocautery, but this is invasive and will result in copious vaginal discharge until healing is completed, which may take weeks. Ablative treatment can also result in cervical stenosis, which can adversely affect future fertility and, if pregnancy is achieved, labor and delivery.

14. Woman just delivered baby. Comes with complain of red swelling on her right breast. They gave a picture too. Mentioned that she had fevers and that it was fluctuant so I thought abscess tha.

- Start antibiotics
- I&D
- Some other dumb options

15. Woman comes to ER with complain of persistent nosebleeding. Past history unremarkable. ER people pack it but its still bleeding constantly. O/E they cant identify the bleeding point. There's blood in the posterior pharynx too and theres blood dripping out the front. Vitally stable, Hb was 10.8 ish I think. NSIM?

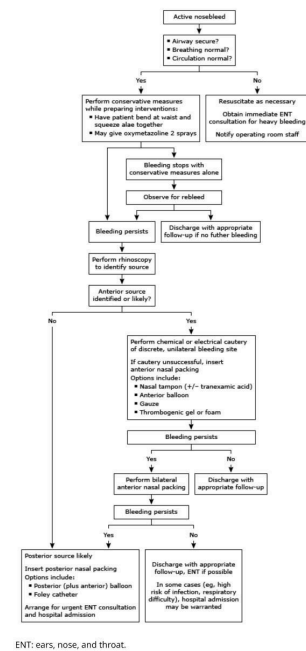
- Endoscopic cautery
- Silver nitrate stick
- Angioembolise
- Pack it some more (posterior nasal packing)

16. HOPI of old lady with complain of lots of vomiting for the past day. Mentioned she's been having symptoms of bloating and abdominal pain since 5 days. 1 week of loss of appetite. Didn't mention anything about constipation sadly. She had a history of gastric bypass surgery like 15 years ago. She was vitally stable, O/E was classic bowel obstruction type with decreased sounds, distended, tympanic sounds etc. Question asked what would you most likely see on abdominal xray.

- Dilated small bowel loops with air fluid levels
- A loop of dilated colon with no air rectum

Bronchiolitis	
Epidemiology	<ul style="list-style-type: none"> Age <2 years RSV most common cause
Clinical presentation	<ul style="list-style-type: none"> Antecedent nasal congestion/discharge & cough Wheezing/crackles & respiratory distress (eg, tachypnea, retractions, nasal flaring)
Treatment	<ul style="list-style-type: none"> Supportive care
Complications	<ul style="list-style-type: none"> Apnea (especially age <2 months) Respiratory failure
Prevention	<p>Palivizumab for selected infants:</p> <ul style="list-style-type: none"> <29 weeks gestation Chronic lung disease of prematurity Hemodynamically significant congenital heart disease
RSV = respiratory syncytial virus.	

Flow diagram for management of epistaxis in adults



Management of ongoing bleeding (7/8/09)

- For anterior epistaxis with the bleeding site identified, consider:
 - Topical vasoconstrictors (e.g., oxymetazoline, phenylephrine)
 - Nasal cautery
 - If bleeding persists or the bleeding site cannot be identified, place nasal packing.
 - For refractory or recurrent bleeding, consider arterial embolization or endoscopic ligation of the bleeding vessel. I.e.:
 - Anterior ethmoidal artery for anterior epistaxis
 - Sphenopalatine artery for posterior epistaxis
- In rare cases, retained nasal packing can cause toxic shock syndrome. [11]

Small bowel obstruction	
Clinical presentation	<ul style="list-style-type: none"> Colicky abdominal pain, vomiting Inability to pass flatus or stool Hyperactive bowel sounds Distended & tympanic abdomen
Diagnosis	<ul style="list-style-type: none"> Dilated loops of bowel with air-fluid levels Partial: Air in colon Complete: Transition point (abrupt cutoff), no air in colon
Complications	<ul style="list-style-type: none"> Ischemia/necrosis (strangulation) Bowel perforation
Management	<ul style="list-style-type: none"> Bowel rest, nasogastric tube suction, intravenous fluids Surgical exploration for signs of complications

Acute colonic pseudoobstruction (Ogilvie syndrome)	
Etiologies	<ul style="list-style-type: none"> Major surgery, traumatic injury, severe infection Electrolyte derangement (↓K, ↓Mg, ↓Ca) Medications (eg, opiates, anticholinergics) Neurologic disorders (eg, dementia, stroke, MS, parkinson)
Clinical findings	<ul style="list-style-type: none"> Abdominal distension, pain, obstipation, vomiting Tympanic to percussion, ↓ bowel sounds If perforation: guarding, rigidity, rebound tenderness
Imaging	<ul style="list-style-type: none"> X-ray: colonic dilation, normal haustra, nondilated small bowel CT scan: colonic dilation without anatomic obstruction
Management	<ul style="list-style-type: none"> NPO, nasogastric/rectal tube decompression Neostigmine if no improvement within 48 hour or if the cecal diameter on imaging exceeds 12 cm (which is a strong predictor of impending perforation).

c. Normal nonspecific gas pattern

d. Free air under diaphragm

i. Now I was sooo confused because, lots of vomiting is SBO, but such a slow onset of symptoms is LBO/volvulus. Cant even remember what I ended up picking last minute rip.

17. Patient comes to ER with same bowel obstruction complaints. There was an abdominal xray picture of what looked like SBO to me. O/E no peritonitis or anything like that, wahi BO signs. NSIM?

a. CT with *rectal* contrast

b. **NG decompression**

18. Some dude had a kidney transplant and was on immunosuppressants. Which of the following is he at greatest risk of infection from due to reactivation of a virus?

a. EBV

b. HHV-8

c. **JC virus**

CMV > BK > EBV

19. Little baby with complain diarrhea. Mention that she had no complain of blood stool or abdominal pain. Vital signs were stable except for a fever of 100.4. O/E she had mild diffuse tenderness of the abdomen. Stool cultures grew Salmonella species. Next step?

a. Start antibiotics

b. **Just observe**

20. Some dude with cancer. Had discussions with his oncologist about chemotherapy. He comes to you (his GP) and he tells you he doesn't wanna try chemo and instead plans to jump a plane to some place to get some pseudo herbal garbage to cure his cancer. What do you do?

a. Assess his decision making capacity

b. Respect his autonomy and let him go eat plants to fix his cancer

c. Call his oncologist and discuss the chemo options

21. This docs wife just died 3 days ago. He comes to the office with his kids with him. His colleagues notice him making a bunch of mistakes. What should they do?

a. Report him to state medical board

b. **Talk to him about his poor performance and that he should get some grief counseling**

i. Sadly there was no option of physician health program

22. 20 ish some girl comes with complain of epigastric pain

Explanation:	
Pathogenesis	<p>Posttransplantation lymphoproliferative disorder</p> <ul style="list-style-type: none"> • (immunosuppressed) following solid-organ or stem cell transplantation → suppressed cytotoxic T-cell immunosurveillance → unchecked viral replication → immunoblastic lymphocyte or plasma cells • Epstein-Barr virus causes >90% of cases, but other human herpesviruses (eg, HHV-8) can also trigger the disease
Manifestations	<ul style="list-style-type: none"> • Fever • Lymphadenopathy & hepatosplenomegaly • Leukopenia • Masses in nonlymphatic tissue
Diagnosis	<ul style="list-style-type: none"> • High viral titers • Biopsy evidence of lymphoid or plasma cell proliferation
Treatment	<ul style="list-style-type: none"> • Reduce (immunosuppression) • B-cell immunotherapy (eg, rituximab) • Epstein-Barr virus titer monitoring

This patient developed a febrile illness with lymphadenopathy and leukopenia while taking immunosuppressive medications following solid-organ transplantation. Although many pathogens can cause febrile illness in the setting of immunosuppression, the presence of an elevated Epstein-Barr virus (EBV) titer suggests **posttransplantation lymphoproliferative disorder (PTLD)**.

PTLD is a plasmacytic or lymphoid proliferation that occurs in the weeks or months following **solid-organ** or hematopoietic stem-cell transplantation. It is triggered by the use of **immunosuppressive medications** that impair cytotoxic T-cell immunosurveillance, which leads to the unchecked replication of oncogenic viral pathogens in transplanted donor tissue or host cells. Most cases (>95%) are caused by the **reactivation of EBV**, a human herpesvirus that establishes **life-long latent (EBV-L) B** and T lymphocytes following initial **infection**.

EBV is an oncogenic virus because it produces proteins (eg, latent membrane proteins) that lead to **B-cell proliferation** and generates proinflammatory nuclear transcription factors (EBNAs) that lead to **B-cell immortalization**. Subsequent cell growth through the mitogen-activated protein kinase pathway results in **lymphadenopathy**, hepatosplenomegaly, bone marrow infiltration (eg, leukopenia, anemia), and/or thrombocytopenia. Diagnosis is often suspected due to a markedly **elevated EBV** titer, but biopsy may be required for

Treatment [1][4][4]	
• Supportive therapy for gastroenteritis: e.g., bland diet, oral rehydration therapy	
• Antibiotic therapy	<ul style="list-style-type: none"> • Not routinely indicated (usually not indicated in immunocompetent individuals) • Indications: severe cases of nontyphoidal Salmonella (consider also for high-risk patients) [3][11] • Preferred regimen usually given for 7–10 days [3][12] • Fluoroquinolones: e.g., ciprofloxacin (off label) severe [3] • CR cephalosporins: e.g., ceftriaxone (off label) severe [3]
• Alternatives	<ul style="list-style-type: none"> • Trimethoprim-sulfamethoxazole (off label) severe [3] • CR azithromycin (off label) severe [3]
• Antibiotic treatment for salmonellosis prolongs fecal excretion of the pathogen. Therefore, it is only indicated for severe nontyphoidal Salmonella infections (eg, in patients with systemic manifestations or a > 3 episodes of diarrhea per day, and those who require hospitalization).	

Salmonella		
	Nontyphoidal	Typhoidal
Etiopathology	<ul style="list-style-type: none"> • Major cause of gastroenteritis worldwide (including United States) • Associated with undercooked poultry • Venereal • Swine & feral • Invasive disease rare 	<ul style="list-style-type: none"> • Most common in resource-limited regions with poor sanitation (eg, unchlorinated treated) • Associated with contaminated food or water
Clinical	<ul style="list-style-type: none"> • Fever & bacteremia • Abdominal pain & rose spots • Life-threatening sequelae (reactive arthritis) 	<ul style="list-style-type: none"> • Blood cultures
Diagnosis	<ul style="list-style-type: none"> • Stool culture 	<ul style="list-style-type: none"> • Potentially fatal • Antibiotics (eg, ceftriaxone) • Drug resistance common
Outcome & treatment	<ul style="list-style-type: none"> • Usually self-limited • Antibiotics rarely needed 	

?a

since 3 days. She has a lot of headaches and she takes a bunch of ibuprofen for it. Shes tried antacids and it hasn't worked. O/E theres tenderness in epigastric region. Next step?

already mentioned in CV apparently. So, NBS => H. pylori related Inx ??

a. EGD

b. Switch ibuprofen to acetaminophen

If no alarming features, go to PPI

c. Start triple therapy *(if A is MALToma)*

23. The linked wala question set where first question was about some 30 year old guy who was having SOB and chest tightness everytime he would go on the eliptical at his gym. This started ever since he joined the gym for a new fitness program. He has a history of coughing with exposure to cold air, and he sometimes wakes up night coughing. Next best step?

i. Get a stress ECG

ii. Get PFTs

iii. Get a CXR

iv. Stress echo I think

b. Answer was PFT because the next question showed PFT results. FEV1 and FVC were about 80 or 85%, TLC FRC were like 98 or 99%. Next step in management?

i. Start him on albuterol before exercise

ii. And a bunch of options related to cardio issues

24. Man with syphilis chancre. Has a RPR titre of 1:16. If you start treatment, what will the lab results look like when you repeat them in 4 months?

a. RPR 1:32 FTABS +ve

b. RPR 1:1024 FTABS +ve

c. RPR -ve FTABS +ve

d. Some options with FTABS negative

Amu -> ? RPR 1:4 & FTA-ABS +ve

A fourfold decline in the nontreponemal titer, equivalent to a change of two dilutions (eg, from 1:32 to 1:8 or from 1:16 to 1:4), is considered an adequate serologic response.

	Syphilis: diagnostic serology
Nontreponemal (RPR, VDRL)	<ul style="list-style-type: none"> • Antibody to cardiolipin-cholesterol-lectin antigen • Quantitative (titers) • Possible negative result in early infection • Decrease in titers confirms treatment • Non specific
Treponemal (FTA-ABS, TP-EIA)	<ul style="list-style-type: none"> • Antibody to treponemal antigens • Qualitative (reactive/nonreactive) • Greater sensitivity in early infection • Positive even after treatment

	RPR	RPR titer	TPHA
Active infection	+	>1:8	+
Latent syphilis	+	Often <1:4	+
False positive	-	Usually <1:4	-
Successful treatment	+ or -	2 titers decrease (e.g., from 1:16 to 1:4)	-

25. Dude currently has pneumonia with neisseria. Previously has had meningococcal meningitis. Asked what to test for the underlying cause. I picked Haemolytic Complement Activity

∆. Terminal complement deficiency (C5 - C9)

26. This 20 something woman who kept having sinopulmonary infections like pneumonia, sinusitis etc in the past couple of years. Has an ANA titre of 1:160. What should you test next?

a. Anti Ds DNA

b. Anti smith

c. Immunoglobulin levels

titre > 1:80 -> auto-immune explore full CV

∆. ? CVID

Pathophysiology	Common variable immunodeficiency
	<ul style="list-style-type: none"> • Abnormal differentiation of B cells into plasma cells — decreased immunoglobulin production
Clinical manifestations	<ul style="list-style-type: none"> • Symptom onset classically age 20-40, as early as puberty • Recurrent respiratory infections (eg, pneumonia, sinusitis, otitis) • Recurrent GI infections (eg, Salmonella, Campylobacter, Clostridi) • Chronic disease: <ul style="list-style-type: none"> • Autoimmune (eg, RA, thyroid disease) • Pulmonary (eg, bronchiectasis, fibrosis) • GI (eg, chronic diarrhea, IBD-like conditions)
Diagnosis	<ul style="list-style-type: none"> • No IgG, IgA, IgM • No response to vaccination
Management	<ul style="list-style-type: none"> • Immunoglobulin replacement therapy

Rheumatologic diseases & commonly associated autoantibodies	Sensitivity (%)	Specificity (%)
Rheumatoid arthritis	RFC 25-50	Anti-CCP 90
Systemic lupus erythematosus	ANA 95	Anti-dsDNA (anti-DNA) 90
Drug-induced lupus	ANA 95	Anti-histidyl-lysine 95
Diffuse systemic sclerosis	ANA 95	Anti-Scl-70 95
Limited systemic sclerosis	ANA 95	Anti-centromere 95
Polymyositis/dermatomyositis	ANA 70	Anti-Jo-1 95
ANA - antinuclear antibodies; anti-CCP - anti-cyclic citrullinopeptides; anti-dsDNA - anti-double-stranded DNA; anti-Jo-1 - anti-Jo-1 antibody; anti-Scl-70 - anti-Scl-70 antibody		

27. Blunt trauma case on a young man. They gave an xray KUB with contrast and asked what was injured. It looked

like the contrast was leaking out of the side of the bladder, so I chose **bladder rupture**. Options had everything from urethral injury, ureteral injury, kidney injury etc.

28. 17 something girl comes in for check up before joining some sports team. Her heart rate I remember was around 52 bpm. Her brother had an implantable defibrillator places for long QT syndrome. Question asked what test would you advise for the girl?

Roman's ND = No SNNHL

Hereditary channelopathies
Brugada syndrome: Autosomal dominant, most commonly due to loss of function mutation of Na⁺ channels. ↑ prevalence in Asian males. ECG pattern of pseudo-right bundle branch block and ST-segment elevations in leads V₁-V₃. Prevent SCD with ICD.
Congenital long QT syndrome: Most commonly due to loss of function mutation of K⁺ channels (affects repolarization). Includes Romano-Ward syndrome – autosomal dominant, pure cardiac phenotype (no deafness); Jervell and Lange-Nielsen syndrome – autosomal recessive, sensorineural deafness.

Causes of acquired long QT syndrome	
Medications	<ul style="list-style-type: none"> Quinidine (due to electrolyte imbalances) Antiemetics (eg, ondansetron) Antipsychotics (eg, haloperidol, quetiapine, risperidone) Tricyclic antidepressants Selective serotonin reuptake inhibitors (eg, citalopram) Antiarrhythmics (eg, amiodarone, sotalol, flecainide) Antibiotics (eg, erythromycin, clarithromycin) Anticholinergics (eg, atropine, scopolamine, benztropine)
Metabolic disorders	<ul style="list-style-type: none"> Electrolyte imbalances (↓ potassium, ↓ magnesium, ↓ calcium) Starvation Hypothyroidism
Bradyarrhythmias	<ul style="list-style-type: none"> Sinus node dysfunction Av-block (2nd or 3rd degree)
Others	<ul style="list-style-type: none"> Hypothermia Myocardial ischemia/infarction Intracranial disease HFH infection

- ECG**
- Echo
- No tests shes good to go
- Some other dumb options

29. Middle aged woman presenting to ER with chest pain, tightness, sweating etc. They do an ECG and it shows ST elevations in leads II, III, avF. They load her up on the aspirin and what not and do a repeat ecg a little while later, which they gave as a picture. From what I could tell, she no longer had elevations in II, III, avf but there were elevations in V2, V3. Question asked her diagnosis.

Vasospastic angina	
Pathogenesis	<ul style="list-style-type: none"> Hypercontractility of coronary smooth muscle
Clinical presentation	<ul style="list-style-type: none"> Young patients (age <40) Smoking (initial other CAD risk factors) Recurrent chest discomfort <ul style="list-style-type: none"> Occurs at rest or during sleep Spontaneous resolution <15 minutes
Diagnosis	<ul style="list-style-type: none"> Antibulatory ECG: ST elevation Coronary angiography: No CAD
Treatment	<ul style="list-style-type: none"> Calcium channel blocker (preventive) Sublingual nitroglycerin (abortive)

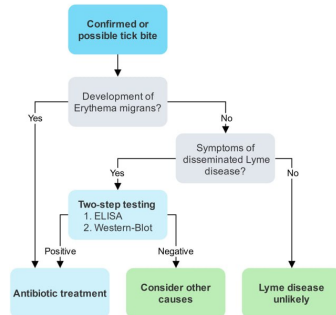
- Right sided MI
- VARIANT ANGINA** (*but explore full CX*)
- And some other options I cant remember rip

INFARCT LOCATION	LEADS WITH ST SEGMENT ELEVATIONS OR Q WAVES
Anteriorseptal (LAD)	V ₁ -V ₂
Anterolateral (LAD)	V ₁ -V ₄
Anterolateral (LAD or LCA)	V ₁ -V ₆
Lateral (LCA)	I, aVL
Inferior (RCA)	II, III, avF
Posterior (PDA)	V ₁ -V ₃ , ST depression in V ₁ -V ₃ with tall R waves

30. Dude presents with complain of mass in his elbow area and they gave a picture. Looked like a lipoma to me so I picked that option. *Δ: ? olecranon bursitis (CV herera jaane)*

31. Kid presents with history of fevers, arthralgias, large macular rash since 6 weeks, and a 2 week history of pain in his knee. Gives history of going camping 6 weeks ago.

- Options had borrelia, scarlet fever, rickettsia
- I went with **borrelia** because the large macular rash sounded like erythema migrans to me



32. Pregnant wife who wants to travel with her husband to Africa. Question says that she is advised not to travel due to zika outbreak there. Her husband is still going to go. What else would you advise her in addition to this?

- Avoid intercourse with her husband for 4 weeks when he gets back *men 3 months, female 2 months*
- Use condoms throughout her pregnancy (went with this)**

Prevention	
• Vaccine against Zika virus does not exist yet	
• Vector control and safer sexual practices	
• Individuals traveling to endemic regions should be told to use insect repellents, mosquito nets, and long-sleeved clothing	
• During pregnancy <ul style="list-style-type: none"> Avoid visiting endemic regions Avoid from unprotected intercourse until the end of the pregnancy if a partner has recently traveled to an endemic region 	

33. 90 ish year old lady comes in for a general health check up. Questions mentions that she had her last mammo in her

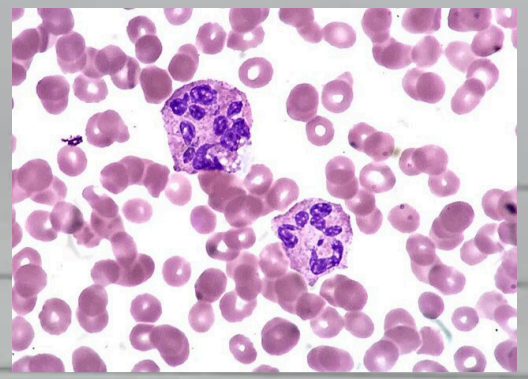
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70s, her last colonoscopy about 7ish years ago. What screening test would you advise at this visit?

- a. Mammo
- b. Colonoscopy
- c. DEXA**
- d. No testing required

23/05/2024

→ folic acid + Vit. B12



1) Hypersegmented neutrophils pic
Anemia

Ans: Folic acid def:

2) Question on **some vitamin toxicity**, was taking double dose.

3) Chest CT picture, some lesions B/L,
Sweat chloride test was -ve.
(explore full CV)

4) Sx of **down syndrome** - asked for Dx.

5) Sx of **DiGeorge Syndrome** - asked about gene deletion.

Ans: 22q11 deletion

kawasaki disease

6) Sx of ~~CHF~~, asked about Rx:

- a) IVIG + low dose aspirin
- ✓ b) **IVIG + High dose aspirin**
- c) IVIG + small dose aspirin
- 2 other irrelevant options.

7) Sexually active young adult, pneumonia Sx,
nasal bleed.

a) **disseminated gonococcal inf.**
other non-infectious options.

Disseminated gonococcal infection	
Manifestations	<ul style="list-style-type: none"> Purulent monoarthritis OR Triad of tenosynovitis, dermatitis, migratory polyarthralgia
Diagnosis	<ul style="list-style-type: none"> Detection of <i>Neisseria gonorrhoeae</i> in urine, cervical, or urethral sample Culture of blood, synovial fluid (less sensitive)
Treatment	<ul style="list-style-type: none"> 3rd-generation cephalosporin intravenously

This patient likely has **disseminated gonococcal infection (DGI)**, which typically presents with purulent arthritis or the following **triad**:

- Polyarthralgia:** Asymmetric pain in multiple distal and proximal joints. Examination usually reveals pain with movement and palpation; multiarticular joint swelling, erythema, and warmth are uncommon.
- Pustular rash:** Most patients have 2-10 pustular or vesiculopustular lesions on the distal extremities; trunk lesions can also occur. The palms and soles may or may not be affected.
- Tenosynovitis:** Patients report pain over the flexor tendons of multiple distal joints (eg, wrists, ankles, fingers, toes) and/or pain with passive range of motion of the joint.

Date: _____

21) Pt. is chest pain, CXR was given, ECG of inferior wall MI was given. Now a new heart sound is present on auscultation (mentioned in cv).

→ MR (d/t papillary muscle rupture)

- a - Mitral regurg
- b - Aortic regurg
- c - ASD
- d - lateral wall rupture

Surgical management and transcatheter mitral repair

- Chronic primary MR
- Indications: severe primary MR with any of the following
 - Asymptomatic patients with LV systolic dysfunction (LVEF < 60% and/or LVESD > 40 mm)
 - Symptomatic patients irrespective of LV systolic function

	Surgical indications for severe chronic MR
Primary MR	<ul style="list-style-type: none"> • Surgery if LVEF 30%-60% (regardless of symptoms) • Consider surgery if successful valve repair is highly likely: <ul style="list-style-type: none"> • Symptomatic & LVEF < 30% • Asymptomatic & LVEF > 60%
Secondary MR (MI, dilated CMP)	<ul style="list-style-type: none"> • Medical management, valve surgery rarely indicated

22) Some mass in hilar region, metastasis to LN mentioned. Histology pic was given. Asking for diagnosis?

- a) Adenocarcinoma
- b) Small Cell CA
- c) Malignant lymphoma.

↳ explore full cv & histo pic.

(in ant. mediastinum)

Abstracts:

1) Correlation of GFR is overall mortality, stroke, HTN

2) Comparison between tPA and standard therapy.

23) Immunocompromised lady living in a building, a TB pt: living in same building on different floor. what if you do:

- a) Isolate TB pt.

(in negative pressure room) & N95 mask

	Infection control isolation precautions
Airborne	<ul style="list-style-type: none"> • Bacterial (tuberculosis) • Viral (varicella, SARS, measles)
Contact	<ul style="list-style-type: none"> • Multidrug-resistant organism colonization (MRSA, VRE) • Enteric (<i>Clostridioides difficile</i>, <i>Escherichia coli</i> O157:H7) • Parasitic (scabies) • Viral (RSV)
Droplet	<ul style="list-style-type: none"> • Bacterial (<i>Neisseria meningitidis</i>, <i>Haemophilus influenzae</i> type B, <i>Mycoplasma pneumoniae</i>) • Viral (influenza, adenovirus)

Droplet: When aerosolized particles are >5 microns (considered relatively large), they cannot stay suspended in the air for long periods and are associated with transmission within only 3-6 feet of the source. **Droplet precautions** require the use of surgical masks within this range.

Airborne: When aerosolized particles are <5 microns, they stay suspended in the air for prolonged periods. Such cases require airborne precautions (ie, negative-pressure rooms, respiratory masks with a minimum 95% filtering capacity [eg, N95 masks]) to prevent inhalation

Type	Pathogen*	Key requirements
Airborne	<ul style="list-style-type: none"> • Bacterial: tuberculosis • Viral: primary VZV (chickenpox), disseminated VZV reactivation (shingles/zoster), dermatomal VZV reactivation (shingles/zoster) in immunocompromised patients, COVID-19, measles 	<ul style="list-style-type: none"> • Negative pressure room • N95 respirator
Contact	<ul style="list-style-type: none"> • Multidrug-resistant organism (eg, MRSA, VRE, ESBL producing) • Bacterial: <i>Clostridioides difficile</i>, <i>Escherichia coli</i> O157:H7 • Viral: RSV, primary VZV (chickenpox), disseminated VZV reactivation (shingles/zoster), dermatomal VZV reactivation (shingles/zoster) in immunocompromised patients 	<ul style="list-style-type: none"> • Gloves & gowns • Single-use equipment (eg, stethoscope)
Droplet	<ul style="list-style-type: none"> • Bacterial: <i>Neisseria meningitidis</i>, <i>Haemophilus influenzae</i> type B, <i>Mycoplasma pneumoniae</i> • Viral: influenza virus, adenovirus 	<ul style="list-style-type: none"> • Mask within 1-2 m (3-6 ft) of patient
Standard	<ul style="list-style-type: none"> • Followed for all patients even when no infection is apparent 	<ul style="list-style-type: none"> • Hand hygiene • Gowns/gloves/masks when appropriate (eg, during procedures)

*Precaution is started immediately when infection with pathogen is suspected.

COVID-19 = coronavirus disease 2019; ESBL = extended-spectrum beta-lactamase; MRSA = methicillin-resistant *Staphylococcus aureus*; RSV = respiratory syncytial virus; VRE = vancomycin-resistant *Enterococcus*; VZV = varicella-zoster virus.

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Explanation:

Differential diagnosis of DSM-5 psychotic disorders	
Brief psychotic disorder	21 days & <1 month, sudden onset, full return to function
Schizophreniform disorder	21 months & <6 months, same symptoms as schizophrenia, functional decline not required
Schizophrenia	26 months (includes 21 months of active symptoms, can include prodromal & residual periods), requires functional decline
Schizoaffective disorder	Mood episode with concurrent active-phase symptoms of schizophrenia + 32 weeks of delusions or hallucinations in the absence of prominent mood symptoms
Delusional disorder	21 delusions & 21 months, no other psychotic symptoms, normal functioning apart from direct impact of delusions

	Schizoaffective disorder
DSM-5 criteria	<ul style="list-style-type: none"> Major depressive or manic episode concurrent with symptoms of schizophrenia Lifetime history of delusions or hallucinations for 22 weeks in the absence of major depressive or manic episode Mood episodes are prominent & recur throughout illness Not due to substances or another medical condition
Differential diagnosis	<ul style="list-style-type: none"> Major depressive or bipolar disorder with psychotic features: Psychotic symptoms occur exclusively during mood episodes Schizophrenia: Mood symptoms may be present for relatively brief periods

CV on diagnosis of Schizoaffective disorder.

31) CV of ADHD Rx: **Methylphenidate**.

32) Hx of some dysplasia at transitional zone of cervix, wasn't treated. Now comes after 3-4 years what if you do.

- a) **Colposcopy**
- b) LEEP
- c) Conization.

Giant cell arteritis: clinical manifestations	
Systemic symptoms	<ul style="list-style-type: none"> Fever, fatigue, malaise, weight loss
Localized symptoms	<ul style="list-style-type: none"> Headaches: located in temporal areas Jaw claudication: most specific symptom of GCA PMR Arm claudication: associated bruits in subclavian or axillary areas Aortic wall thickening or aneurysms CNS: TIAs/stroke, vertigo, hearing loss
Visual symptoms	<ul style="list-style-type: none"> Amaurosis fugax: transient vision field defect progressing to monocular blindness AION: most common ocular manifestation
Laboratory results	<ul style="list-style-type: none"> Normochromic anemia Elevated ESR & CRP Temporal artery biopsy
Treatment	<ul style="list-style-type: none"> PMR only: low-dose oral glucocorticoids (eg, prednisone 10-20 mg daily) GCA: intermediate- to high-dose oral glucocorticoids (eg, prednisone 40-60 mg daily) GCA with vision loss: pulse high-dose IV glucocorticoids (eg, methylprednisolone 1,000 mg daily) for 3 days followed by intermediate- to high-dose oral glucocorticoids

AION = anterior ischemic optic neuropathy; CRP = C-reactive protein; ESR = erythrocyte sedimentation rate; GCA = giant cell arteritis; IV = intravenous; PMR = polymyalgia rheumatica; TIA = transient ischemic attack.

33) CV on diagnosis of temporal arteritis

34) Heart sound on Aortic stenosis.

	Valve replacement in aortic stenosis
Severe AS criteria	<ul style="list-style-type: none"> Aortic jet velocity ≥ 4.0 m/sec, or Mean transvalvular pressure gradient ≥ 40 mm Hg Valve area usually ≤ 1.0 cm² but not required
Indications for valve replacement	Severe AS & ≥ 1 of the following: <ul style="list-style-type: none"> Onset of symptoms (eg, angina, syncope) Left ventricular ejection fraction $< 50\%$ Undergoing other cardiac surgery (eg, CABG)

35) Case of Aortic stenosis, signs of heart failure. Asked about. Sarcomeres II be added in **parallel** or longitudinally or added in both ways.

→ concentric hypertrophy

36) Some scenario on a pt. having prothrombotic state, recurrent inflammation on arm/hand/eye. Also somewhere mentioned about bleeding. what is reason for this ??

- a) **Colorectal cancer**
- b) COPD
- c) Crohn's disease.

37) Some questions on Lynch Syndrome. (HNPCC).

-APC

38) Adolescent is difficulty breathing through nose. normal breathing from mouth.

- a) Bil nasal polyps
 - b) DNS
 - c) Edema turbinates
- (all options are plausible)*

39) Hx of rhinorrhea, unilateral L cheek pain (suggesting sinusitis). How to investigate.

- a) CT Paranasal sinuses
- b) X-ray skull

40) Signs of L heart failure (hepatomegaly, pulmonary edema, dyspnea, murmur of MS). what will you do next ??

- a) Loop diuretic
- b) Thiazide diuretic
- c) ACE inhibitors
- d) β -blocker

41) Recurrent Syncope, dizziness. CV of BPPV ?

Dx \Rightarrow Dix-Hallpike

- a) Epley maneuver \rightarrow Rx

42) Acute Cystitis picture in young female ?? Rx.

- a) Nitrofurantoin

Abstract

Anastrozole effect in Breast Cancer

(a) 57 Ys female had no symptoms but her mother had breast cancer at the age 57 Ys and sister had breast cancer at 42 Ys. Her menopause at 47 Ys. Which is the strong factor for risk of breast cancer in this patient?

(a) family Hx (I marked this)

(b) Age of menopause

(c) Age

(2) Asked about RR?

(3) Validity of study?

b) Abstract

Estrogen only hormone causing breast cancer they gave multiple types of estrogen hormones like vaginal estrogen, transdermal estrogen, oral estrogen. The study compared b/w them and calculated RR. It was so difficult don't remember exactly.

(a) Which of the following limits study?

(a) Exclusion criteria (b) outcome

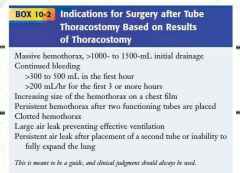
③ Sequential

① MVA case had chest trauma showed X-ray showing left sided opacity + tracheal deviation to opposite side asked Diagnosis?

① hemothorax (correct)

② After diagnosing hemothorax chest tube placed but after some time still patient's condition didn't improve what to do next?

① thoracotomy (marked this).



④ Sequential

① Case of Acute Rhinosinusitis had to diagnose this.

② asked its treatment don't remember options.

⑤ Pt: underwent stem cell transplantation develop rash + diarrhea + Jaundice → GVHD (marked)

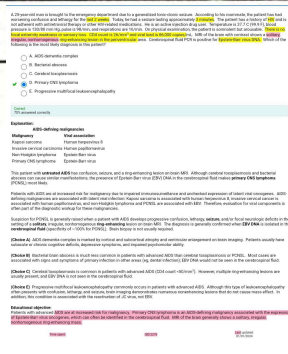
⑥ Pt: underwent kidney transplant has developed SOB pneumonia Histo Pic given showing owl eye inclusions asked treatment?

① oral acyclovir (b/c of CMV) ganciclovir

⑦ Patient was diagnosed w HIV asked this patient is most likely to develop cancer due to what?

① EBV (marked this) ② Kaposi sarcoma ③ CMV

SOLO



2000

⑤ 17 Yrs old patient → Vaccinations upto date what to do next → Meningococcal booster.

⑦ 11 Yrs old girl every thing normal what to do next?
- give HPV vaccine
<15yrs = 2dose 6monthn apart
>15yrs = 3dose

⑩ Scenario of Crohn's disease given Pic given it looked like Perianal fissure. So I marked Perianal fissure.

⑪ another Qs about Crohn disease pt undergone ileocollectomy but after some days had abdominal pain + diarrhea now next step in management?
a) give Ceftriaxone b) give prednisone c) mesalamine

⑫ Patient had pneumonia + recurrent skin abscess diagnosis Ko E. Rese confirm Roma hair?
- I did Dihydroamine test (CGD thought)

⑬ Clostridium difficile prevention → hand washing

⑭ Clostridium difficile treatment → a) IV Vancomycin
b) oral fidaxomicin (marked this) vanco plus metro for fulminant

⑮ Patient had Rti upper Quadrant pain + diarrhea watery first later bloody CT given showing liver abscess
→ E. histolytica (marked this)

t/t : metronidazole

Adequate hand hygiene, which is a standard precaution, should always be performed correctly. Health care personnel (HCPs) for infection control should use either alcohol with a peroxide, quaternary ammonium or chlorhexidine as the preferred disinfectant. In most instances, the use of alcohol-based hand rubs (ABHR) is preferred over soap and water. ABHR should be used for hand hygiene when the hands are not visibly soiled. Soap and water should be used when caring for patients with acute infectious diarrhea or infections due to spore-forming organisms such as C. difficile because ABHRs are ineffective against spore-forming organisms and a given use of ABHRs does not prevent cross-contamination with spores. Use of soap and water before putting on gloves and gloves as well as after removing them.

When caring for patients with certain infectious diseases such as C. difficile infection, in addition to standard precautions, the HCPs should follow contact precautions (e.g., isolating patients, wearing gloves and a gown upon room entry).

18) Scenario of Pseudotumor Cerebri, funduscopy shows papilledema what next? → LP

17) Patient had lg lung cancer had several lymphadenopathies and also some findings in liver asked the poor prognostic factors? → Secondary lesions (marked this thinking of mets)

16) Trisomy 21 markers → β hcg, inhibin b, estriol, α -fetoprotein

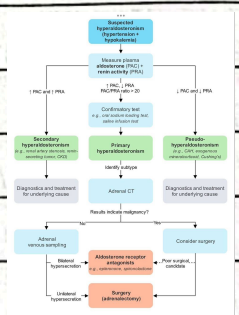


also known as painful bladder syndrome. suprapubic pain worsen by bladder filling, relieved by voiding

15) Diagnosis of interstitial cystitis asked.

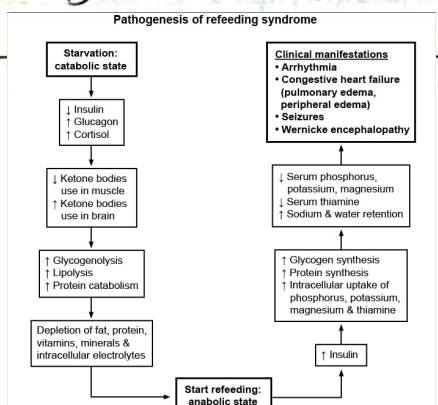
14) Young aged male had HTN not treated medications asked Cause of Refractory HTN. abdominal bruit absent.

- a) fibromuscular dysplasia
- b) Renal atherosclerosis
- c) Essential HTN
- d) Hyperaldosteronism ??



21) Case of Rhabdomyolysis → girl doing excessive exercise had pink urine no other symptoms

22) Refeeding syndrome case asked first laboratory change u get → ↓ phosphate levels (marked)



Date _____

(23) Anorexia nervosa Case give asked lab changes?
→ ↓ K⁺

(24) Patient of MVA got 6 Pints of blood transfusions + 6 Pints of FFP, after 30 minutes he developed SOB + BIL lung infiltrates
asked Cause? Always TACO if underlying cardiac disease ya vane irrespective of timing

(a) TACO >6hrs (b) TALI <6hrs (c) pulmonary embolism

(25) Hematuria + flank pain going to scrotum
next step of diagnosis?

(a) CT scan of kidney (thought nephrolithiasis)

(26) 3-4 Questions on ~~over~~ urinary incontinence

(27) child had 2nd episode of UTI age was <1 yrs what to do next?

(a) Renal ultrasound (b) Voiding cystourethrogram
USG tes pachhi VCUG

(28) 12 WOG - pregnant lady came had Hx of chronic hypertension. What is she at risk of?
→ Preeclampsia (marked this)

(29) 38 WOG pregnant lady had vaginal bleeding but now stopped U/S shown what to do next?

(a) plan for C/S (I thought this as placenta previa)

SOLO



Post tussive vomiting in infants
Post tussive syncope in adults

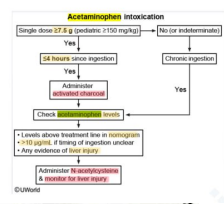
Indications (pre/post exposure prophylaxis)	Treatment
<ul style="list-style-type: none"> Close contact (eg, household members, direct contact with individuals with symptomatic disease) High-risk patients, even with limited exposure (eg, pregnant, child, immunocompromised) 	<ul style="list-style-type: none"> Age < 1 month: azithromycin Age ≥ 1 month: azithromycin, clarithromycin, or erythromycin

30) Scenario of Pertussis case give Paroxysmal Cough & Vomiting asked what to do next?
 (a) give azithromycin to room mate (I marked that)

31) other Scenario on pertussis asked it's treatment?
 (a) Macrolides
 Macrolides
 Vac natalo lai vaccine pani Dina parchha

32) patient present & Vomiting abdominal pain Constipation ↑ bowel sounds asked diagnosis
 (a) Small bowel obstruction

33) Young girl found unconscious by her mother mother found acetaminophen bottle next to her ~~was~~ Activated Charcoal given next step?
 (a) Gastric lavage (b) Intubation (c) fluids
 Lavage contraindicated in unconscious



34) Confusing question on Hypercalcemia and ↑PTH asked Cause but the options were so difficult I couldn't interpret it as it was the case of primary hyperparathyroidism but options were difficult.

35) other Qs on hypercalcemia but it was moderate and asymptomatic asked treatment.
 (a) I/V normal saline

Management of hypercalcemia	
Severe (calcium >14 mg/dL) or symptomatic	Short-term (immediate) treatment <ul style="list-style-type: none"> • Normal saline hydration plus calcitonin • Avoid loop diuretics unless volume overload (heart failure) exists Long-term treatment <ul style="list-style-type: none"> • Bisphosphonate (zoledronic acid)
Moderate (calcium 12-14 mg/dL)	<ul style="list-style-type: none"> • Usually no immediate treatment required unless symptomatic • Treatment is similar to that for severe hypercalcemia
Asymptomatic or mild (calcium <12 mg/dL)	<ul style="list-style-type: none"> • No immediate treatment required • Avoid thiazide diuretics, lithium, volume depletion & prolonged bed rest

SOL 0

* Abstracts:

1. Breast cancer Prevention by Anastrozole.

Inclusion criteria: Post-menopausal women of certain age (50-60) maybe. A very narrow range.

The study showed that there was significant difference. Anastrozole actually prevented breast cancer at the end. A lot of extra stuff like side effects and their CIs were given.

Question 1: Post menopausal woman & family hx of breast cancer. No age of onset of menopause mentioned.

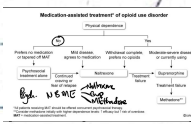
Doctor prescribes Anastrozole. Why

- Age of onset of menopause.
- Family history.

Don't remember other options.

Don't remember other questions.

1. Opioid toxicity.



1. Opioid use disorder. Current user. What to give.

a) Buprenorphine b) Naltrexone

c) Acamprosate

2. Patient had Opioid use disorder. Now has reduced craving. What must he have taken?

a) Buprenorphine b) Naltrexone c) Acamprosate

3. Seq. 1 → Case of GVHD → T cell response

2 → What to do?

a) Start steroids b) ↑ Immunosuppressants c) Stop immunosuppressants

4. 16 year old guy. Parents came concerned saying "You know what goes on teenagers these days". What to counsel them about?

a) STIs b) Bullying c) Accidents d) Drug abuse

most common cause of death in this age group is accidents > suicide > homicide

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5. Pt. had exposure of something (that causes ILD) ~~maybe~~ What is the measure of earliest lung involvement.

a) FEV₁ b) FVC c) DLCO d) RV.

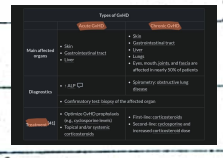
6. ECGs: * 2 ECGs of Inf. MI in 2 dip. blocks.

Inf wall + ST elevation in lead 3 > lead 2
RCA occlusion

* Apib

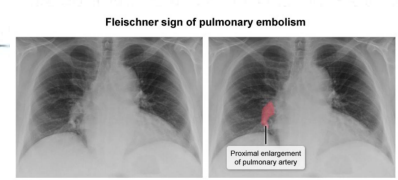
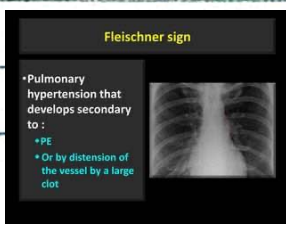
* P.E (No S1Q3T3) case helped.

Alcohol withdrawal = chlorazepate
Opioid withdrawal = buprenorphine
Opioid abuse = naltrexone, mirtazapine, naltrexone, Bupropion, naltrexone
Alcohol abuse = naltrexone, clonidine, naltrexone
Acamprosate = alcohol or liver disease



Category	Validated screening instrument
Mental health	Validated depression questionnaires
Social health	Validated instruments about social activity, substance use, and sexual activity
Substance use	Validated instruments about tobacco, alcohol, and other substances
Safety	Inquiry about history of injury, falls, and violence

PULMONARY FUNCTION TESTS
Measurement of FEV1 and FVC is essential in the evaluation of obstructive pulmonary disease. A restrictive defect is defined by a reduced total lung capacity (TLC) and a proportionally reduced residual volume (RV). In the normal adult, TLC is approximately 2.5 L and RV is approximately 1.5 L. In the normal adult, FEV1 is approximately 1.2 L and FVC is approximately 2.0 L.

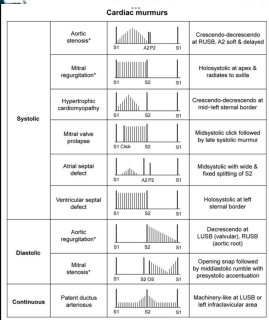


Mitral valve prolapse
Most common murmur. Described as mid-systolic click. "Mitral valve prolapse" is a busy term that refers to connective tissue degeneration causing MVP in Marfan and Ehlers-Danlos. Almost always asymptomatic. On ZCK forms, they want you to know about "mitral valve prolapse syndrome," which is symptomatic MVP that presents as repeated episodes of "heaving chest pain" on the left side in an otherwise healthy patient 20-30s. They might say there is a Ho of MI in the family, but this is MVP, not MI. Answer on surgery form is "no treatment necessary".
USMLE loves using MVP as a distractor in panic disorder questions, particularly on the ZCK Psych CMS forms. They will give long paragraphs about panic attack disorder + also mention there's a mid-systolic click, they'll ask for cause of patient's presentation → answer = panic disorder, not MVP → student is confused because they say mid-systolic click, but the MVP on the cause of the patient's presentation, the panic disorder is MVP's are usually incidental, benign, and asymptomatic. MVP does not progress to mitral regurg almost always. So don't think that MVP and MR are the same.
Eccentric mitral regurgitation (eccentric mitral regurgitation) can also be described

7. Heart sounds : Couldn't really understand from the sound. Case didn't help either.

either.

1. Old pt. has only dyspnea → AS
2. Young lady → Aortic sclerosis
3. Complain only palpitations → MR. (No MI history). Maybe wrong.



8. Sub-Arachnoid hemorrhage CT.

It clearly showed SAH. Asked NBS.

- a) MRI b) LP c) MR angio d) Repeat CT

No management option. I didn't get the point. CT clearly shows SAH, why investigate further? Anyways.

Best initial test: immediate head CT without contrast [19][21]
 Confirmation of SAH: Obtain angiography to confirm source of bleeding and plan treatment.
 Nondiagnostic head CT but persisting suspicion: Perform second-line diagnostic tests

Clinical features	Complications
<ul style="list-style-type: none"> Most commonly due to ruptured arterial aneurysm ("berry" aneurysm) Severe headache at onset of neurologic symptoms Meningeal irritation (eg, neck stiffness) Focal deficits uncommon Rebleeding (first 24 hr) Hyponatremia (after 3 days) Hydrocephalus/increased intracranial pressure Seizures Hypotension (usually from syndrome of inappropriate antidiuretic hormone secretion) 	<ul style="list-style-type: none"> Rebleeding Hydrocephalus Seizures Hypotension

9. Asplenic patient had meningitis. What could have prevented this?

- a) Prophylactic antibiotics.
 b) Pneumococcal vaccine.

No option of meningococcal. Don't remember if time of splenectomy was given.

10. 17 year old boy. Last visit to a doctor was when he was 10 yrs old. At that time fully immunized. Now what to give him?

- a) Meningococcal Booster (No mention of 1st dose)
 b) HPV.

Human papillomavirus vaccination

Routine vaccination

- All persons up through age 26 years: 2- or 3-dose series depending on age at initial vaccination or condition
- Age 9-14 years at initial vaccination and received 1 dose or 2 doses less than 5 months apart: 1 additional dose
- Age 9-14 years at initial vaccination and received 2 doses at least 5 months apart: HPV vaccination series complete, no additional dose needed
- Age 15 years or older at initial vaccination: 3-dose series at 0, 1-2 months, 6 months (minimum intervals: dose 1 to dose 2: 4 weeks / dose 2 to dose 3: 12 weeks / dose 1 to dose 3: 5 months; repeat dose if administered too soon)

According to this 2 types KO huncha...
 MO localista - more common - after trauma - radiotherapy t/t
 MO progressiva - extremely rare - also after trauma - no t/t or nsaid...
 Amboss

11. Trauma to leg. Then some pain & swelling. Pain gone but swelling still there. X ray given. Looked like Myositis ossificans. What to do?

- a) MRI
- b) Biopsy
- c) ALP
- d) Rest + NSAID.

12. GBS case without hx of URTI. Rx asked?

- a) IVIG.
- b) Steroids

13. A student saw a resident messing up. Whom to report?

- a) Talk to resident.
- b) Talk to other residents.
- c) Talk to clerkship director.
- d) Report to senior resident.

Whose mistake and whom to report:

- 1) Med student - Dean or course director
- 2) Resident: Programme Director/Dept head
- 3) Student in training: Local Supervisor 1st
- 4) Attending: Department/Division Head
- 5) Physician in Private practice: State Board

- e) Report to Attending physician of resident.

Source: 100 Cases

14. A resident saw a physician messing up. What to do.

- a) Go talk to him.
- b) Report to state board.
- c) Talk to residents.
- d) Report to prog. director.
- e) Report to ~~state board~~ some lower board.

15. Sponge left in abd. of pt. Physician halfway closed the pt. And counting showed 1 sponge missing. What to do?

• Retained surgical objects (eg, sponges, instruments) frequently occur even when surgical counts are thought to be reconciled. Interruptions increase the risk of counting error; holding a brief time-out (ie, a pause in activities to announce that the count is about to begin) can improve situational awareness (ie, group vigilance for safety concerns), reduce interruptions, and improve count quality.

- a) Close the patient & do imaging.
- b) Stop suturing & count again.
- c) Open sutures & explore.
- d) Close patient & give antibiotics.
- e) Do imaging now.

16. 2 study reports written. Study A has some cases of same disease & wrote a report about it. Study B took same cases & compared their risk factors to a control group. What study types.

Yesma cases are taken from the published cases, and controls are taken then risk factor frequency is compared.

✓ a) Case Series, Cohort.

b) Case control, cohort

c) Case report, Case-control

Combinations like that.

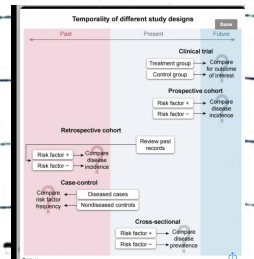
No option of case series, case control. Study A had large number of subjects.

17. A lot of questions on antibiotic of choice. eg: for cellulitis, PPRM, Endometritis, Pneumonia, GI infections. etc.

18. A lot of questions on screening eg: Lung cancer, colon cancer, HPV, AAA. etc.

19. A lot of questions on vaccination.

These were all impossible. Vaccines, Screening & Antibiotics. You just can't pick out of 2.



Seemed absolutely impossible to me.

20. MVA patient came to ED. Died later on. Had no family members there or any other information.

What to do?

- a) Tell his family he is dead.
- b) Look for his cards to see if he is a donor.

21. Trauma to shin. Hematoma picture given. What will be complication?

- a) Abscess
- b) Varicose veins
- c) DVT.

22. Strabismus case. Management asked.

- a) Put patch on normal eye.
- b) Put atropine in affected eye.
- c) Refraction something.
- d) Laser therapy

Strabismus (ocular misalignment)	
Clinical findings	<ul style="list-style-type: none"> • Constant eye deviation at any age • Intermittent eye deviation at age <4 months • Eye deviation on cover test • Asymmetric corneal light reflexes • Asymmetric intensity of red reflexes • Torticollis or head tilt
Treatment	<ul style="list-style-type: none"> • Strengthen deviated eye (eg. patch unaffected eye, cycloplegic drops to blur unaffected eye) • Correct refractive errors (ie, prescription glasses) • Surgery
Complications	<ul style="list-style-type: none"> • Amblyopia (reduced visual acuity) • Diplopia

23. There were questions where BMI was not given. Weight & height were given and something asked in line of management. Like weightloss, starting statin or something.

24. ASCVD risk 7%, LDL <190, Family hx +ve for MI. Asked why to start statin in this pt?

- a) Family hx
- b) ASCVD
- c) LDL

Risk Category	Criteria	Treatment Goal			
		Non-HDL-C (mg/dL)	LDL-C (mg/dL)	Non-HDL-C (mg/dL)	LDL-C (mg/dL)
Low	0-1 major ASCVD risk factors Consider other risk indicators, if known	<130	<100	≥190	≥160
Moderate	2 major ASCVD risk factors Consider quantitative risk scoring Consider other risk indicators*	<130	<100	≥160	≥130
High	≥3 major ASCVD risk factors Diabetes 0-1 other major ASCVD risk factor No evidence of end organ damage CVD stage III or 4 LDL-C >190 mg/dL Quantitative risk score reaching the high risk threshold	<130	<100	≥130	≥100
Very high	ASCVD Diabetes ≥2 major risk factors or Evidence of end-organ damage*	<100	<70	≥100	≥70

Indications for statin therapy in prevention of ASCVD	
Secondary prevention	<ul style="list-style-type: none"> • Established ASCVD • Acute coronary syndrome • Stable angina • Arterial revascularization (eg, CABG) • Stroke, TIA, PAD
Primary prevention	<ul style="list-style-type: none"> • LDL ≥190 mg/dL • Age ≥40 with diabetes mellitus • Estimated 10-year risk of ASCVD ≥7.5%-10%

ASCVD = atherosclerotic cardiovascular disease; CABG = coronary artery bypass grafting; PAD = peripheral artery disease; TIA = transient ischemic attack.

2. Different OCPs and associated risk of VTE.

- * Inclusion: Any OCP leading to even a single episode of VTE.
- * Exclusion: Anyone who had a hx of abortion.

Study was comparing rate of VTE in 4 types of Progestin only OCPs.

OCP	RR	VTE rate	CIs
Levonorgestrol	e	x	a
Denogestrol	f	y	b
etc	g	z	c
etc	h	z	d

Also compared Progestin + Estrogen OCP.

OCP	RR	VTE rate	CIs
Levo + estro	e	x	a
Deno + estro	f	y	b
etc.	g	z	c
etc.	h	z	d

Question 1: A woman came for contraception. Doc. gave her Levonorgestrol instead of Denogestrol. Why?

- a) Study supports levo bez it has lower rate of VTE.
- b) Study doesn't support it bez CI is not significant.

Similar options. Thing is, both levo & Deno had non-significant CIs. But RR with Levo was < 1 . and for Deno was > 1 . So I picked option **A**.

Question 2: Something about Relative Risk decrease. I think they meant RRR. Options has values. One option was "Can't be calculated". Relative risks were given for all OCBs so I guess it can be calculated.

Question 3: Study's drawback?

- a) Didn't take confounding factors under consideration.

Don't remember other option.