

Breadcrumbs

NOTES

SOURCE: MED435 TEAMS. This document covers random notes the doctor's EMPHASIZED on per lecture. IT'S NOT A SUMMARY
 This document also does not cover ALL emphasized notes nor does it cover ALL lectures. Proceed with diligence.

Lecture	Notes
RA	<ul style="list-style-type: none"> - Extra-articular manifestations of prolonged active RA → amyloidosis and usually presents with nephrotic syndrome! - RA by itself won't cause nephrotic syndrome but if complicated by amyloidosis it usually presents with nephrotic syndrome in the form of proteinuria - RA by itself won't cause glomerulonephritis, while SLE can cause glomerulonephritis
SPA	<ul style="list-style-type: none"> - The major gene associated with AS and other forms of SpA is HLA-B27 - Ankylosing spondylitis → non-articular features → anterior uveitis is NOT related to disease activity. Anterior uveitis first line Rx = topical steroid - Reactive arthritis → Sexually acquired infection: chlamydia trachomatis. OR GI infection → shigella, salmonella, yersinia, campylobacter, clostridium difficile.
Osteoarthritis	<ul style="list-style-type: none"> - X-ray showing joint space narrowing, osteophytes, and subchondral cyst → OA
HSV	<ul style="list-style-type: none"> - Recurrent herpes labialis is the most frequent clinical manifestation of HSV 1 reactivation
PTH	<ul style="list-style-type: none"> - HyperPTH → shortened QT interval --- HypoPTH → prolonged QT interval - The most favorable site of calcification in hypoPTH → basal ganglia
Obesity	<ul style="list-style-type: none"> - Management of obesity → Indication of starting pharmacotherapy (BMI > 30) OR (BMI 27-30) with comorbidities¹. - Indication of surgical intervention? (BMI > 40) OR (BMI > 35) with comorbidities.
Pituitary disorders	<ul style="list-style-type: none"> - Pit-1, Prop-1, Pitx2 → transcription growth factors responsible for pituitary cell development - Genetics related to pituitary adenoma are: MEN-1, gs-alpha mutation, PTTG gene, FGF receptor-4 - The 1st line Rx for all pituitary adenomas is surgery EXCEPT for prolactinoma, 1st line → medications
DM1	<ul style="list-style-type: none"> - Patient's with DM1 often first present with acute DKA and as such 1st management is

¹ These include diabetes, sleep apnea, osteoarthritis, cardiomyopathy.

	<p>FLUID.</p> <ul style="list-style-type: none"> - DM1 occurs in younger patients, it is immunological, it is acute, and it involves DKA.
Epilepsy	<ul style="list-style-type: none"> - Phenytoin and carbamazepine aggravate absence and juvenile myoclonic epilepsy. - The drug of choice in absence seizure: ethosuximide or valproic acid. - Sodium valproate is the best for unclassified or specific syndromes. (✘ in pregnancy)
Myopathies	<ul style="list-style-type: none"> - Among inherited myopathies, dystrophin is the most commonly mutated gene. The second most common is DM1 (dystrophia myotonia 1). - If you're presented with a family tree in which only males are affected. You're dealing with an x-linked recessive disease (duchenne and becker dystrophy).
Stroke (2L's)	<ul style="list-style-type: none"> - Lateral medullary syndrome (wallenberg/PICA syndrome): sensory deficit in opposite side + ipsilateral ataxia. - The most common cause for intraparenchymal hemorrhage (IPH) is HTN (and the most common location for a HTN IPH is PUTAMEN [basal ganglia]) - The second most common cause of IPH is amyloid angiopathy (and in the following order are the locations: frontal>parietal>occipital>temporal)
CNS infections	<ul style="list-style-type: none"> - If you have a pregnant women and her husband has meningitis give CEFTRIAXONE as prophylactic therapy.
Anemia	<ul style="list-style-type: none"> - DDX of anemia: <ul style="list-style-type: none"> - <u>MCV < 80 fL (TAILS)</u>: 1. Thalassemia 2. Anemia of inflammation 3. Iron deficiency 4. Lead poisoning 5. Sideroblastic anemia. MOST COMMON OF WHICH is iron/thalas. - <u>MCV normal with low retic. Count</u>: 1. Bone marrow failure (aplastic anemia) 2. BM suppression (toxins/sepsis/organ failure/chronic diseases/chronic inflammation) 3. BM infiltration (lymphoma, leukemia, metastatic solid tumor, granulomatous diseases such as TB). - <u>MCV normal with high retic. Count</u>: 1. Acute bleeding 2. Hemolysis 3. Treated nutritional deficiency (iron, B12). - <u>MCV > 100 fL</u>: 1. Megaloblastic anemia (B12 deficiency, folate deficiency, drugs such as methotrexate) 2. Non-megaloblastic (liver disease, alcohol, myelodysplasia, thyroid disease, myeloma, congenital BM failure). - B12 deficiency/folate deficiency/methotrexate use cause IMPAIRED DNA metabolism.