





Disease of Adrenal glands	
Condition	Investigation of choice
ACTH independent Cushing syndrome Adrenal adenoma	 CORTISOL ↑ ACTH ↓ : CT Abdomen
Primary aldosteronism CONN HTN ± Hypokalemic M. alkalosis Saline infusion Test / salt loading Test	 ALDOSTERONE ↑ : ENac : CD Salt + H ₂ O + 
Pheochromocytoma Screening → loc:	24-hour urinary Fractionated Metanephrine levels PLASMA FREE Metanephrine levels
Adrenal insufficiency Addison →	ACTH stimulation test = Autoimmune TB : AIDS 

↑ 24 hr urinary VMA : NEUROBLASTOMA n-myc
 * ← pheochromocytoma

Metformin, sepsis

HYPERGLYCEMIC
HYPEROSMOLAR

Unconscious unresponsive diabetic patient

	Lactic acidosis	D.K.A	WMA	? CVA
pH ↓	7.3	7.3	7.40	7.4
pCO ₂ ↓ (35-45 mmHg)	30	30	38	38
HCO ₃ ↓ 22-26	15	15	24	24
Urine ketones	⊖	4+	2+	—
RBS	250 mg/dL	300 mg/dL	800 mg	150
Any special investigation ordered ↳	plasma β OH butyrate = (n) S. lactate ↑	plasma β OH butyrate level: ↑	plasma osmolality ↓ 310 mosm	NOCT HEAD

Methods of Insulin administration

	<p>CONTINUOUS GLUCOSE MONITOR</p> <p>* CONTINUOUS INFUSION</p>	<p>C.S.I.I</p> <p>SC INSULIN</p>
	<p>INSULIN se =></p>	<p>PENS : 32G needle 90° angle To skin</p>
	<p>INHALED insulin (Rapid acting)</p>	
	<p>MULTI DOSE * 1ml insulin</p> <p>Storage Temp: +2 To 8°C</p> <p>Shelf life: ~ 4 weeks</p>	<p>VIAL $\frac{100U/ml}{10ml}$</p> <p>Syringe</p>

once a Long-acting weekly insulin: ICODEC

Start insulin if

1. Asymptomatic t2DM with HbA1c > 10% ✓
2. Symptomatic t2DM with weight loss and HbA1c > 9%

3. D.K.A, HYPEROSMOLAR COMA, K^+ ↑
REGULAR sc/iv

Somogyi phenomenon (extra amount of bed time insulin)

* Nocturnal HYPOGLYCAEMIA
* GLUCAGON, CORTISOL, Catecholamines, GH

* Pre breakfast HYPERGLYCAEMIA

Dawn phenomenon (GH peak between 4 am to 7 am)

T2DM Nocturnal EUGLYCAEMIA / mld HYPERGLYCAEMIA

Pre-breakfast HYPERGLYCAEMIA

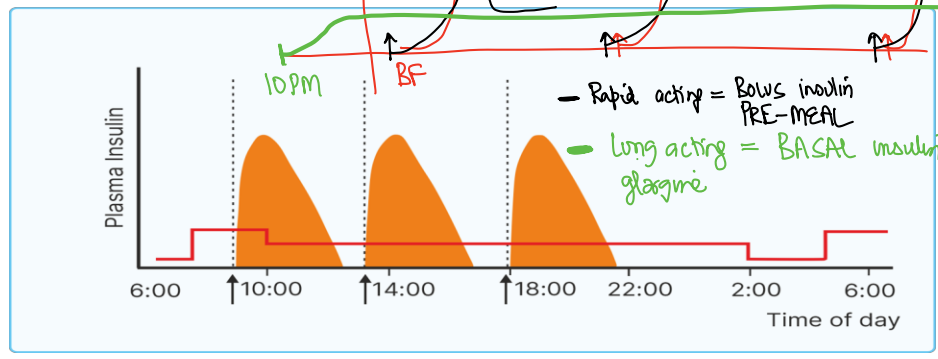
Empagliflozin

* T2 DM: HbA1c: 8-10% : (No) comorbities : Metformin + SGLT2i
 " " " : ASCVD : Metformin + SGLT2i + GLP1RA
 " " " : HF : Metformin + SGLT2i
 " " " : CKD : Metformin + SGLT2i
 eGFR: < 20 : DPP-4i

* metformin CI: eGFR < 30 → INSULIN: 80% Calculated dose



Basal Bolus regimen
TYPE 1 DM



Skin involvement in DM

MC skin lesion in type 2 DM **DIABETIC DERMOPATHY**
(BINKEY SPOTS)

MC skin lesion in type 1 DM **NECROBIOSIS LIPOIDA DIABETICORUM**

DM, DEPRESSION, DVT: PNET
MC skin lesion in glucagonoma: **NECROLYTIC MIGRATORY**
DERMATITIS = **ERYTHEMA**

Causes of Acanthosis nigricans (R) — TYPE II DM, PCOD
METABOLIC SYNDROME
C PANCREAS, C stomach



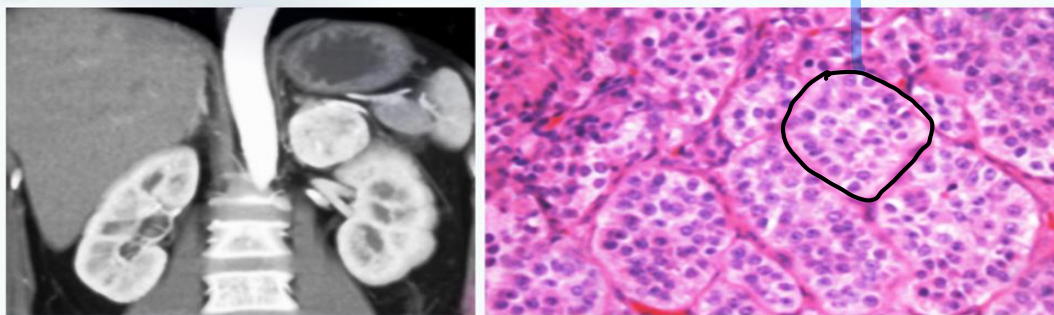
- Syndrome - X**
metabolic syn
1. Acanthosis nigricans: DORSUM Axilla > 90cm
 2. CENTRIPETAL obesity: Abdominal ○ > 80cm
 3. HTN: > 130/85 mmHg
 3. IGT FBG: 100-125, 2hr: 140-195
 4. TRIGLYCERIDES > 150
 5. HDL: < 40 / < 50
- Syndrome - 2**
Syndrome X + OSA

* NESTS of ZELERBILLEN



Pheochromocytoma

Fight or flight phenomenon



Leading catecholamine produced by normal adrenal medulla

↳ EPINEPHRINE *

Leading catecholamine produced by pheochromocytoma

↳ NOREPINEPHRINE

Leading catecholamine produced by pheochromocytoma in MEN-2 is SIPPLE

↳ EPINEPHRINE

Extra adrenal pheochromocytoma is called

↳ PARAGANGLIOMA
(location: sym. ganglie)

* NF-1 is associated with Pheochromocytoma

Palpitations, headache, diaphoresis

PHD PAROXYSMAL HTN : Early phase
EPISODIC HTN

Clinical Buzz words

It can mimic Anxiety neurosis/ Panic attacks

B2D: ETILOZAM, SSRI: PAROTEXINE

POSTURAL HYPOTENSION : late phase illness
IGT, wt loss

Screening test

24 HOUR Urinary fractionated Metanephrine

IOC

→ PLASMA FREE Metanephrine levels

↳ LAP. U/L ADRENALECTOMY

Rx of benign pheochromocytoma

PRE-OP: PHENOLXY BENZAMINE : 7AM

PROPRANOLOL : Post B.F

$\alpha + \beta$ 😊

$\beta + \alpha$ 😞

~~$\alpha = VC$ HTN~~ ~~$\beta = HTN$~~
 Unopposed

RX of malignant pheochromocytoma

SURGERY ci ↑

Use MIBG containing iodine-125 and Chemotherapy

methy iodo-benzyl-guanidine scan

- * benign vs malignant pho
- * location medulla: sym. chain

Gallium - 68

DOTATE-PET/CT

> MRI Abdomen

1. PNET
 2. pheochromocytoma

↓
 Ga-68, DOTATE-PET/CT

- * 18-F-FDG, distant Mets → *fluorodeoxyglucose*
 ↳ FLUORIDE

* INTRA-OP HTN, PRE-OP HTN

↑ both ↓
 drugs works in

: Nicardipine
labetalol

* INTRA-OP HTN dlt Tumor manipulation

SOD NITROPRUSSIDE

* Pseudocushing app = alcohol intake



Cushing Syndrome



MOON FACIES



BROAD PURPLE STRIPE

DORSOCERVICAL FAT PAD

Clinical features

1. weight gain
 2. Lemon on sticks appearance
 3. Dorsocervical fat pad
 4. Purple striae
 5. Thin skin with increasing bruising
 6. Hypokalaemic alkalosis
 7. Hirsutism and oligomenorrhea
 8. Hyperpigmentation of knuckles and IP joints
- partial MSH like action

CORTISOL ↑: ENac ++

↑ ACTH

Causes

Exogenous: STEROIDS: CORTISOL ↑, ACTH ↓

O: Oat cell carcinoma lung: (Ectopic) ACTH ↑, CORTISOL ↑

A: Adrenal Adenoma: (autonomous) CORTISOL ↑, ACTH ↓

P: PITUITARY Adenoma: (endogenous) ACTH ↑, CORTISOL ↑

Screening test

24 hr urinary CORTISOL ↑↑

IOC

MIDNIGHT salivary CORTISOL ↑

PIT: Oat cell carcinoma lung, Carcinoid syn

Endogenous vs ectopic source

ACTH ↑↑ ACTH ↑↑ = PRE

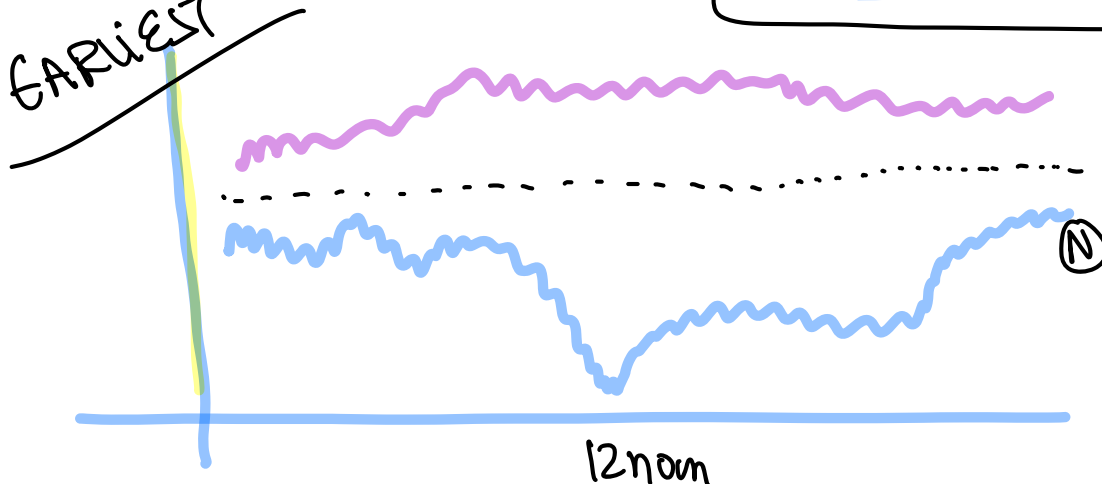
High dose dexamethasone suppression test: ACTH independent Cushing syn

ACTH ↑ ACTH ↑↑ = POS

Partial suppression of ACTH: Pit. Adenoma = Cushing disease

Non suppression of ACTH: Oat cell carcinoma lung = Ectopic

EARLIEST

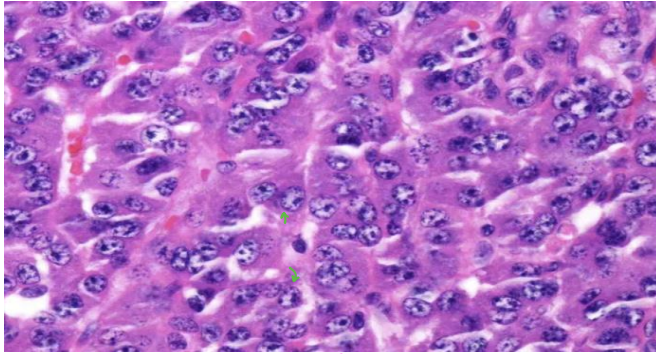


LOSS OF DIURNAL variation of CORTISOL production

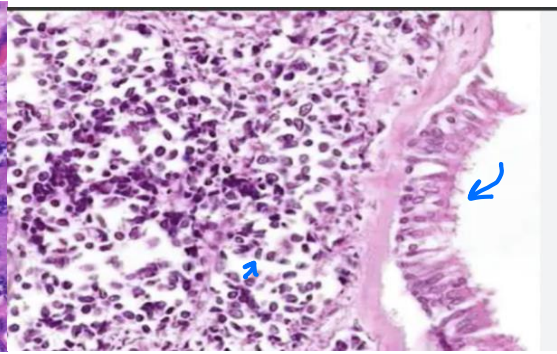
(A)

ECTOPIC CAUSES

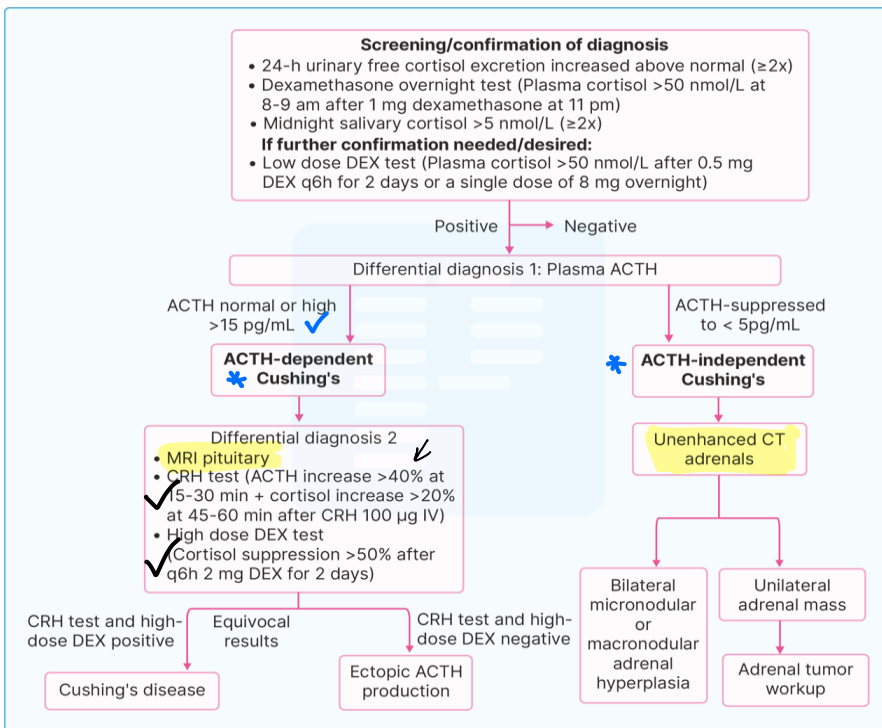
(B)



SALT & PEPPER
Carcinoid Tumor



CRUSHED nuclei
Oat cell carcinoma



CRH TEST: ACTH ↑↑
Pit. CORTICOTROPH Adenoma

Disorders of Calcium Metabolism



CHVOSTEK
facial N irritability
Twitching

DOC for tetany iv 10% calcium gluconate

Leading cause of Hypocalcaemia

- ↓
1. Inadvertent removal of parathyroid gland during total thyroidectomy
 2. AUTOIMMUNE

DOC for management of surgically induced hypoparathyroidism **TERIPARATIDE**

Calcium & 1/25 OH vit D3 ↓ PTH^{SC} ↓ PTH levels: ↑

	Primary Hyper PTH	Secondary Hyper PTH	Hypoparathyroidism <u>AUTOIMMUNE</u>	PHP <u>PTH receptor #</u>
Calcium	↑	↓	↓	↓
Phosphate	↓	↑	↑	↑
SAP	↑	↑	NORMAL	NORMAL

Bone

Tc99 - SESTAMI Bi
SPECT/CT

Calcium
vit D3
CINACALCET

TERIPARATIDE

?

Pseudo

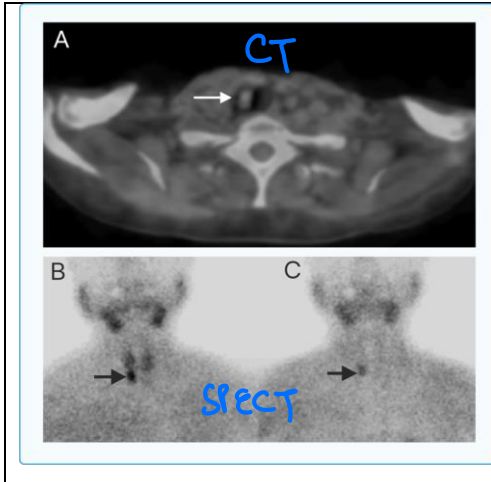
Tc99 PERTECHNATE scan = MECKEL DIVERTICULUM

Pg. 101 Tc99 STRESS SESTAMI Bi = CHRONIC STABLE Angina

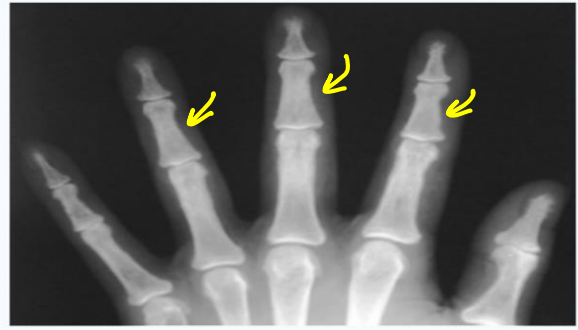


Renal colic
 OSTEITIS CYSTICA FIBROSA
 BROWN TUMOR, CONSTIPATION (T₄↓)

Primary hyperparathyroidism



Stones - bones - moans and psychic moans



SUB-PERIOSTEAL Resorption of phalanges

Leading cause of hypercalcemia in hospitalized patients

↳ Malignancy: Squamous cell ca lung, Ce breast
 PTH-rp

Management of hypercalcaemic crisis

NS + FUROSEMIDE (loop Ca⁺)

iv ibandronate

nasal calcitonin spray

DENOSUMAB

ORAL PHOSPHATE

(Refractory) Hemodialysis

do CONFUSION, ALTERED MENTATION

OLE: dehydration +

ECG ⇒ QT shortening

Na ↓ K ↓ Ca ↑ = 12.5 mg/dL

(Risk of SYSTOLIC cardiac arrest)

OSTEOCLAST Activity ↓

Tachyphrys

Doc 2

1. NS + FUROSEMIDE (loop Ca⁺)
2. iv ibandronate
3. nasal calcitonin spray
4. DENOSUMAB
5. ORAL PHOSPHATE
6. (Refractory) Hemodialysis

* SARCOIDOSIS: vit D₃ production; non caseating granuloma
 : STEROIDS

* vitamin D₃ intoxication → STEROIDS

* M. Myeloma: CRAB
 L Calcium ↑ : STEROIDS



Additional Notes

Q. 8yr child: admitted c pneumonia, iv levofloxacin X 2 days

day 3: vomiting, abdo pain, drowsy

O/E: **THREADY** pulse BP = 70/50 mmHg
cold clammy extremities

RBS = 300 mg/dL

PH ↓ pO₂ ↓ HCO₃ ↓

URINE KETOSTIX: 4+

1. NS Bolus 1hr → insulin infusion
2. NS infusion " " → " "
3. NS " + KCl " " → " "
4. insulin infusion → NS

Q. 60yr Kldo **Hashimoto's**: levothyronine poor compliance
 do DYSURIA, FEVER
 midstream urine culture
 Rx: NORFLOX

> 48 HOURS: **UNCONCIOUS, UNRESPONSIVE** state
 pulse: 60/min BP: $\frac{130}{110}$ mmHg Temp: **35°C**

1. MYXEDEMA coma, Hashimoto encephalopathy
 ↓
 iv T₄, T₃
2. ↓ Temp ⊕, seizures ⊕
 iv methylprednisolone

Additional Notes

Q.

GRAVE : non compliant \bar{c} Medication
during Sx: HR \uparrow BP \uparrow Temp $\uparrow > 40.5^{\circ}\text{C}$

? THYROID STORM



Additional Notes



Additional Notes



Additional Notes



Additional Notes



Additional Notes

Pulmonology



7.35-7.45

ABG analysis Hacks 35-45 mm 22-26 meq

10 change

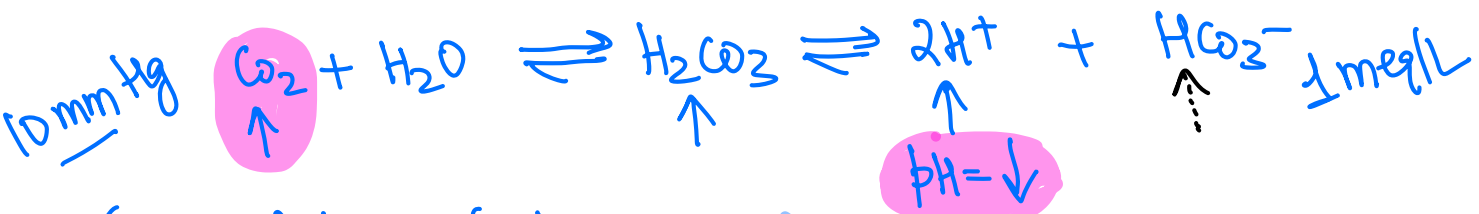
pH 1.	pCO2 2	HCO3 3	PC:	Interpretation
↓	↓	↓	PC:	<u>Metabolic</u> acidosis
↑	↑	↑	PC:	Metabolic alkalosis
<u>↓</u>	↑	↑	PC:	<u>Respiratory</u> acidosis
<u>↑</u>	↓	↓	PC:	Respiratory alkalosis

ROME

Respi opposite pH pCO2
Metabolic equivalent + "

Extra mile

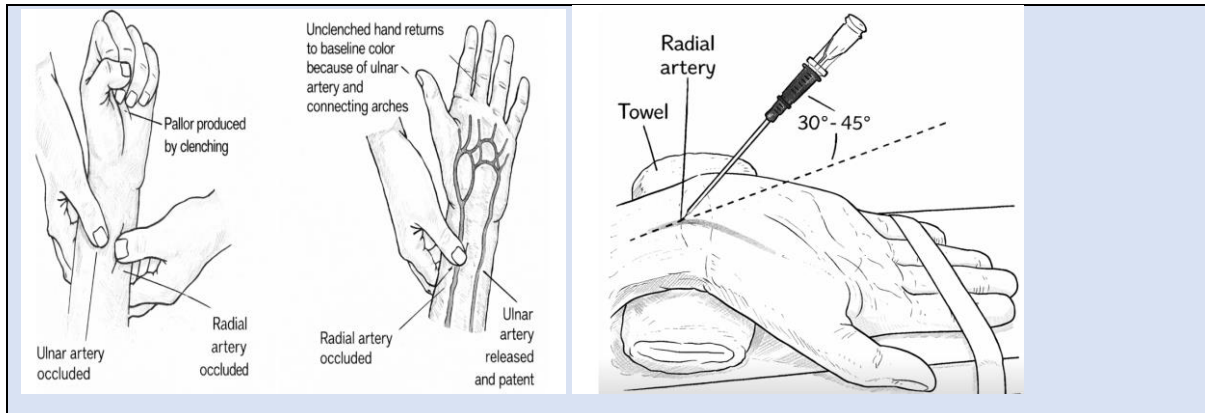
Aldosterone has inverse relation with potassium
pH has inverse relation with Potassium



(STATUS Asthmaticus) **Respi Acidosis**

COMPENSATION: KIDNEY: PCT: ULTRAFILTRATE
Metabolic alkalosis HCO3 Reabsorption ++

Modified Allen test is done for integrity of palmar arch before doing ABG



P02, pCO2 values are by default in mm Hg

How to interpret in 30 seconds

- Acute exacerbation of COPD
- Status asthmaticus

	Acute exacerbation of COPD	Status asthmaticus	Respi acidosis H ⁺ UNCOMPENSATED	Respi acidosis H ⁺ PARTIALLY COMPENSATED	Respi acidosis Fully compensated
1. pH ↓	7.2	7.22 ↓	7.36 NORMAL		
2. pCO2 ↑	60 ← 40	58 ↑	50 ↑		
3. HCO3 ⁻ (n)	26 → 28	28 ↑↑	32 ↑↑↑↑		

* Acute Exacⁿ of COPD ⇒ NIV (Bi-PAP)

NIV 1 A Acute pulm edema: AHF

2. B Acute exaⁿ of ch. bronchitis

3 C Covid-19 SEVERE pneumonia

Scenario 1:

	Parameters	Values	Interpretation
1.	pH	7.25 ↓	<p>Metabolic acidosis PC</p> <p>COMPENSATION VS MIXED</p> <p>Expected $pCO_2 = HCO_3 + 15$ $= 10 + 15$ $= 25 \text{ mmHg}$</p> <p>Expected $pCO_2 = \text{actual } pCO_2$</p>
2.	PCO2 <i>actual</i>	25 ↓	
3.	HCO3	10 ↓	

Expected PCO2= HCO3 plus 15 = 10 + 15 = 25 mm Hg

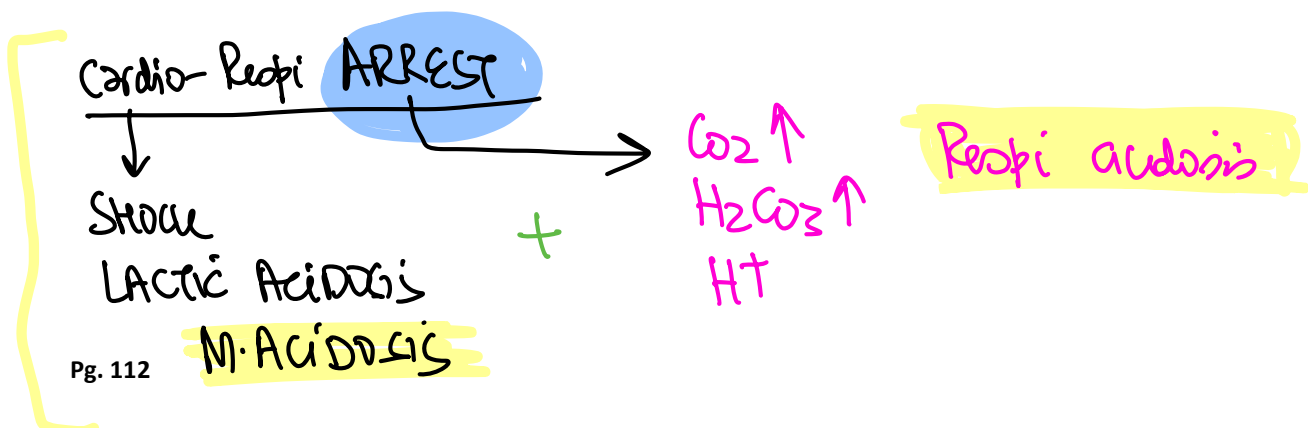
Optional for FMGE

Scenario 1.1

	Parameters	Values	Interpretation
	pH	7.25 ↓	<p>M-ACIDOSIS</p> <p>* expected $pCO_2 = HCO_3 + 15$ $= 10 + 15 = 25 \text{ mmHg}$</p> <p>Actual $pCO_2 >$ expected pCO_2</p> <p>Mixed disorder : Respi acidosis</p>
	PCO2 <i>actual</i>	30 ↓	
	HCO3	10 ↓	

Hypoventilation causing decreased CO2 washout

Expected PCO2= HCO3 plus 15 = 10 + 15 = 25 mm Hg and actual PCO2=



Scenario 1.2

Parameters	Values	Interpretation
pH	7.1 ↓	<p>M. Acidosis PC</p> <p><u>Compensation vs mixed</u></p> <p>expected $p_{CO_2} = HCO_3 + 15$ $= 6 + 15$ $= 21 \text{ mmHg}$</p> <p>expected $p_{CO_2} > \text{actual } p_{CO_2}$</p> <p>PC M. Acidosis = M. Acidosis + compensatory R. alkalosis</p> <p>$(1.5 \times HCO_3) + 8 \pm 2$</p>
PCO ₂ actual	16 ↓	
HCO ₃	6 ↓	
pO ₂	90	

Modified winter formula

$$HCO_3 + 15 = \text{expected } p_{CO_2}$$

* expected $p_{CO_2} = \text{actual } p_{CO_2}$

* // $> \text{actual } p_{CO_2}$

Compensation (PC)

actual $p_{CO_2} > \text{expected } p_{CO_2}$

Mixed

Scenario 2: Acute exacerbation of chronic bronchitis

Parameters	Values	Interpretation
pH	7.2 ↓	<p>R. acidosis</p> <p><u>Partially compensated</u></p> <p>CO₂ narcosis</p> <p>- ASTERIXIS</p> <p>- Metabolic encephalopathy</p>
PCO ₂ *	60 ↑	
HCO ₃	28 ↑↑	
pO ₂	90	<p><u>longest acting β₂ agonist</u></p> <p>INDACATEROL + <u>ICS</u></p> <p><u>LAMA</u> budesonide</p> <p>Tiotropium</p>

Management =



NIV

- ① NIV
1. UNCONSCIOUS, UNRESPONSIVE
 2. SBP ↓: CS, POST-MI, extensive GI bleeding
 3. MAXIOFACIAL Trauma
UPPER Airway burn

Scenario 3: COVID Moderate/ Severe case R. Rate: ↑ CO₂ ↓

Parameters	Values	Interpretation
pH ↑	7.5	Respiratory alkalosis Partially compensated Uncompensated R. alkalosis R. alkalosis kick out HCO ₃ HCO ₃ ↓
PCO ₂ ↓	30	
HCO ₃ ↓	20	

$HCO_3 = n$ UC
 $HCO_3 = \text{change}$ PC
 R. Acidosis
 alkalosis

Scenario 4: Chronic Vomiting in case of CHPS neonate

Parameters	Values	Interpretation
pH ↑	7.5	ALKALOSIS Respir Metabolic Uncompensated partially compensated
PCO ₂ (n)	45 / 48	
HCO ₃ ↑	30 (22-26) 33	

Loss of Acid
 Loss of K⁺
 Loss of Cl⁻

dehydration: aldosterone ↑
 ENac
 salt H₂O
 ↓
 K⁺/H⁺

M. ALKALOSIS: kick out HCO₃⁻

PARADOXICAL ACIDURIA

Optional for FMG

Scenario 5

CKD patient with multiple episodes of vomiting

inability to K^+/H^+ → Loss of Acid

Parameters	Values	Interpretation
pH	7.4	M. ACIDOSIS + M. ALKALOSIS = <u>mixed disorder</u>
PCO2	40	
HCO3	24	
Na	130	Anion gap = $(Na^+) - (Cl^- + HCO_3^-)$ $(140) - (106 + 24)$ 6-12 meq/l $(140) - (130)$
Chloride	80	

$$26 = 130 - (80 + 24)$$

*

Must know facts

Fluid of choice for metabolic alkalosis → Normal saline
 Metabolic alkalosis: RL: lactate: → HCO3 generation
 H+ excess

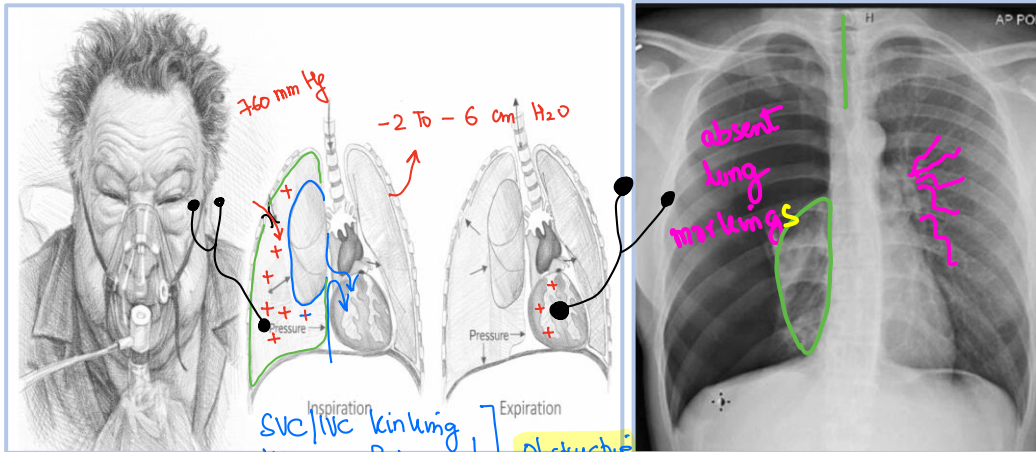
Side of Excess normal saline is → M. ACIDOSIS,
 O.g.f. NaCl Cl- excess
 To maintain electroneutrality HCO3- go inside cells.

Fluid of choice for Metabolic acidosis
 RL

Best for management of Acute exacerbation of chronic bronchitis
 Niv/Bi-PAP

* Best for management of Respiratory alkalosis → Paper bag Rebreathing
 Panic attacks, Pneumonia
 ARDS: RRate ↑ CO2 ↓ H2O3 ↓ pH ↑
 HAFE

Pneumothorax



When to say Tension Pneumothorax

Key words

any 1 findings

1. - Hypotension ✓
2. - Mediastinal shift/ Apex beat displaced away ✓
3. - Elevated JVP ✓
4. - Absent air entry
5. - Absent Breath sounds
6. - Hyper-resonant percussion note

Obstructive SHOCK

heart is beating!

pulse ⊖

PEA

NON SHOCKABLE

rhythm

HAMMAN CRUNCH

Sign = pneumo mediastinum

walking on snow c leather boots

Rx: 1. WIDE BORE Needle THORACOSTOMY

* 5 ICS, Ant To Mid-Axillary line

* 2 ICS, lateral To MCL

2. I.C.D intercostal drainage Tube TDC
5 ICS, Δ of safety

3. UNDER WATER SEAL DRAINAGE

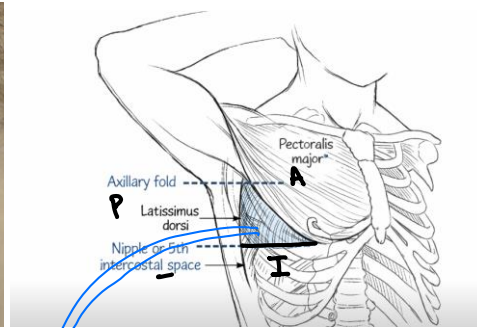
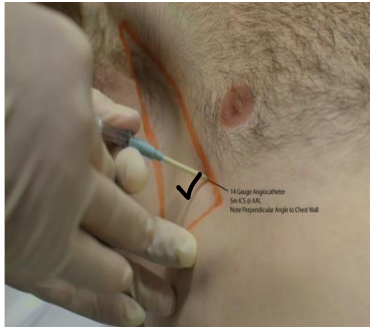
adult

child/adult →

36-40 fr

OBSTRUCTIVE SHOCK	DISTRIBUTIVE SHOCK <u>SVR ↓</u>
1. Cardiac Tamponade	S septic shock
2. Tension pneumothorax	A Anaphylaxis, Addisonian CRISIS
3. MASSIVE PE	N Neurogenic shock

First line intervention



AIR collection chamber

blood collection chamber

Air collection chamber should exhibit intermittent bubbling

-Continuous bubbling in air collection chamber indicates

- ↳ AIR LEAK / POOR CONNECTION
- ↳ BRONCHOPLEURAL FISTULA

-Intermittent bubbling stops

- > 1 HOUR ↳ TUBE KINKING
- ↳ DISLODGEMENT OF TUBE

😊 ~ 7 day GRADUAL Re expansion

EMERGENCY THORACOTOMY & HEMOSTASIS

Hemothorax

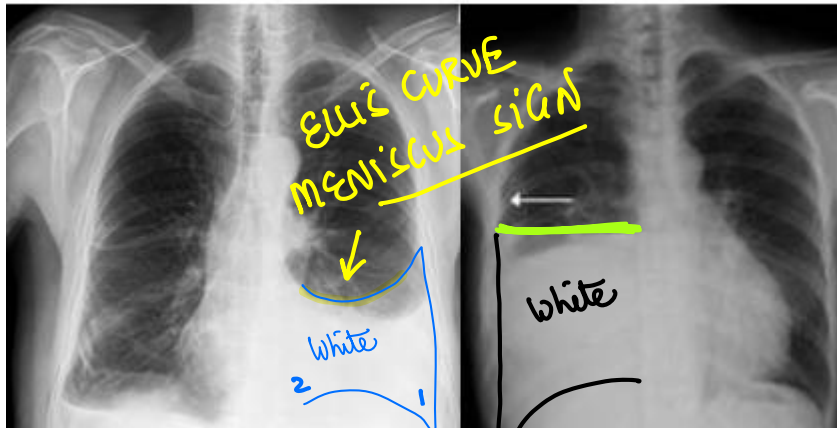
When to inform surgeon

1. Drain amount > 200 ml per hour for 2 hours
2. Drain amount > 1.5L single reading

EMERGENCY THORACOTOMY

1. Cardiac Tamponade due to Hemopericardium
2. Traumatic pneumothorax : 200 x 2 , > 1.5L

Pleural effusion

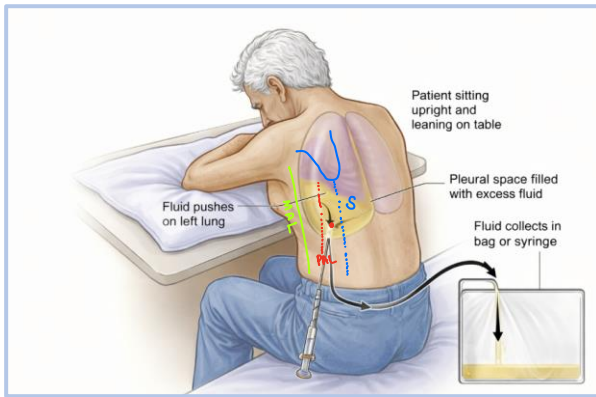


HYDROTHORAX

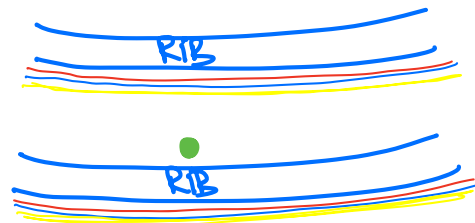
HYDRO-PNEUMO-THORAX

COSTOPHRENIC angle: 1st To obliterated in hydrothorax

Site for thoracocentesis



8th ICS midway between PAL & Scapular line



Maximum amount removed per sitting

< 1.5 litres sitting

Risk of Reexpansion pulm oedema

Site from perspective of avoiding injury to Neurovascular bundle

✓ SUPERIOR BORDER of INF RIB *

✓ **Transudative**: PRESSURE ISSUES: always BL

* HF: (MC)

PLEURAL space (n): 5-15ml

Oncotic pressure: albumin ↓
 * CIRRHOSIS
 * Nephrotic syn
 * Peritoneal dialysis

✓ **Exudative**: FIBRIN Rich fluid : Clots on standing
TB, Ca lung

Above the upper border of lower rib

Lights criteria

CIRRHOSIS, Nephrotic syn

* **Transudative**: CHAP: CHF, Hypoalbuminemia, Ascites and peritoneal dialysis

PF protein \downarrow : < 0.5 g S. Protein

> 0.5 **Exudate**

PF LDH \downarrow : < 0.6 g S. PROTEIN

> 0.6

PF Adenosine deaminase levels elevated are seen in

\rightarrow TB pleural effusion (CBNAAT)
slo

Right sided pleural effusion is seen in

MEIG SYNDROME

FIBROMA/THECOMA + ASCITES + Rt sided pleural effusion

Bloody pleural effusion is seen in

Trauma | warfarin | TB | Ca lung

Low sugar with cholesterol crystals are seen in

\rightarrow Rheumatoid A

Methotrexate	
PLEURAL PLAUVE	-
Calcification	o

Low sugar in Pleural fluid

\rightarrow ? malignancy: CYTOLOGY/ IHC of PF

Left sided effusion

BOER HAAVE

esophageal malignancy

Acute pancreatitis (Tail)

* Ca ovary

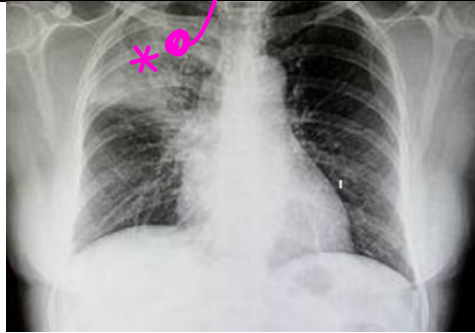
SEROUS CYSTADENOCARCINOMA



malignant Ascites

* Adenocarcinoma lung \rightarrow malignant pleural effusion
Rx: PLEURODESIS (doxycycline)
 \rightarrow obliteration of pleural surface

Pneumonia



Bronchial B / Tubular B

Most reliable auscultatory finding of lobar pneumonia

legionelle pneumophile

Organism that can cause both typical and atypical pneumonia

CAP

Pneumococcus / Strep. pneumoniae
 Pyogenic M: > 3mths To upto 55yrs

VAP

Pseudomonas AERUGINOSA > Acinetobacter
 * CETRIMIDE AGAR
 * MEROPENEM > PIPERACILIN - TAZOBACTAM

Leading cause in children with Cystic Fibrosis/ Mucoviscidosis (F508 mutation, del phenylalanine, Chromosome 7), AR

1480 amino acids
CFTR protein

< 10 years

S. Aureus

> 10 years

PSEUDOMONAS

CURB 65 Score or CRB-65 score

BUN

update

The CURB-65 criteria include five variables: confusion (C); urea >7 mmol/L (U); respiratory rate ≥30/min (R); blood pressure—systolic ≤90 mmHg or diastolic ≤60 mmHg (B); and an age of ≥65 years. Patients with a score of 0 (a 30-day mortality rate of 1.5%) can be treated as outpatients. With a score of 1 or 2, the patient should be hospitalized unless the score is entirely or in part attributable to an age of ≥65 years; in such cases, hospitalization may not be necessary. Among patients with scores of ≥3, mortality rates are 22% overall; these patients may require ICU admission. The PSI has greater efficacy than CURB-65 but is more difficult to calculate.

CONFUSION = 1 point
 UREA ↑ = "
 RR > 30/min = "
 BP ≤ 90/60 = "
 age > 65yr = "

0-1 (only for age) = OPD BASIC = oral A/B

1-2 = IPD " = IV "

3 = ICU " = IV "

(except age)

DECIDE need for HOSPITALIZATION: CURB 65



RX: SCORE 0-1

Outpatients oral Amoxicillin + Azithromycin

Comorbidities, previous hospitalization

Stents, CKD, UD

oral amoxicillin + Clavulanic acid + Azithromycin

SCORE 1-2-3



Inpatient / ICU

IV CEFTRIAZONE + Azithromycin

or
or

IV Ampicillin + Sulbactam + Azithromycin

IV Respiratory FQ L levofloxacin
M Moxifloxacin
G gemifloxacin

VRSA

Add coverage for S. Aureus

iv linezolid

* Central line

iv MEROPENEM

*

Add coverage for Pseudomonas Aeruginosa

