

Lab results in primary health care

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Data station

- Hematology (CBC).
- Liver profile
- Lipid profile
- Hyperuricemia
- FBS/RBS/HbA1c/IGT/IFG
- Thyroid profile
- renal profile ,eGFR and UMA
- Hormonal therapy
- Vit d AND vit B12/hair fall
- Immunology tests/urine /stool analysis

ASK 7

- Why did you want it?
- Who ordered it for you?
 - Where did you do it?
 - When did you do it?
- What do you know about this test?
 - Was it first time?
 - How did you do it?

CBC

- 28 years old lady complains of easy fatigability, hair loss and palpitation since 3 month.
- Her CBC showed

RBC 4.28 LOW MCV 69.9 LOW

HGB 9.7 LOW HCT 29.9 LOW

MCH 22.6 LOW MCHC 32.2 LOW

RDW 18.4 HIGH

CASE 1

- What is the most likely Diagnosis?
 - a. Iron deficiency anemia
 - b. Thalassemia Trait
 - c. Both iron deficiency anemia on top of thalassemia trait
 - d. Chronic disease anemia

CASE 2

A 9-years old male

•	WBC	4.2	(3.5-10.5)
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• RBC
$$4.5 \times 10^9$$
 (4.0-5.2)

• Plt
$$360 \times 10^3$$
 (150-400)

- Serum iron 160 g/dl (50-120)
- **Serum ferritin 169ng/ml** (7-140)
- Iron saturation 50% (0-5)
- Q- what is the interpretation?
- Microcytic hypochromic anaemia with increased iron stores
- Diagnosis: Thallassemia

	Iron deficiency Anemia	Thalassemia	
Hb	Low	Low (100-115)	
MCV	< 85: Consider Fe deficiency <75: Fe deficiency	Low (60's)	
RBC Count	Low	High	
RDW	High	Normal	
Ferritin	Low (<30)		

MCV/RBC

- If more than 13 likely to be IDA.
- If less than 13 likely to be thalassemia trait.

	IDA	Anemia of chronic inflammation
Serum ferritin	Decreased	Normal or decreased
Serum iron	Normal or decreased	Normal or decreased
Total iron binding capacity or transferrin	Increased	Normal or decreased
% iron saturation	Decreased (<10%-15%)	Normal or decreased
MCV	Decreased	Normal or decreased
RDW	Increased	Normal
sTfR/log ferritin ratio	>2	<1
Hepcidin (not currently clinically available)	Suppressed	Increased

 $IDA = iron \ deficiency \ anemia; \ MCV = mean \ corpuscular \ volume; \ RDW = red \ blood \ cell \ distribution \ width; \ sTfR = serum-soluble \ transferrin \ receptor.$

Microcytic hypochromic anemia

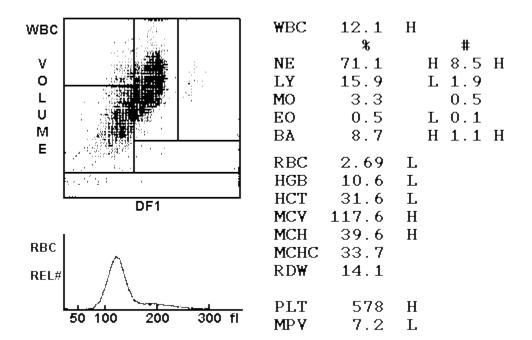
- Differential diagnosis:
- 1. Iron deficiency anemia.
- 2. Lead poisoning.
- 3. Thalassemia trait.
- 4. Sideroplastic anemia.

Iron deficiency anemia

- Look at ferritin level.
- Treatment:
- 1. Ferrous sulfate 200mg bd on empty stomach (2hrs prior and 4 hours after antacid).
- 2. If not tolerated \rightarrow decrease the dose or give ferrous gluconate.
- 3. Ascorbic acid(vit c) 250-500mg bd with iron.
- 4. Hgb should increase by 2g/100ml over3-4 wks
- 5. Confirm the response 2-4w after starting.
- 6. Continue treatment for 3 months after correction to replenish the iron store.
- 7. Once normal, monitor every 3 months for 1 year then annually.

CASE 3

- 48 years old Indian patient. Not known to have any previous medical problems.
 Presented with easy fatigability, weight loss, and numbness in both upper and lower limb.
- His CBC showed:



- What is the most likely diagnosis?
 - a. Multiple Myeloma
 - b. B12 Deficiency
 - c. Folate Deficiency
 - d. Acute leukemia

Macrocytic hyperchromic anemia

Differential diagnosis:

- 1. Vit B12 deficiency.
- 2. Falate deficiency.
- 3. Alcoholic liver disease.
- 4. Pregnancy.
- 5. Reticulocytosis.
- 6. Myelodysplasia.
- 7. Cytotoxic drugs.

Vitamin B12 deficiency DDx:

- 1. Dietary (vegans).
- 2. Low intrinsic factor (pernicious anemia/gastrectomy).
- 3. Blind loop syndrome.
- 4. Ilial resection.
- 5. Crohn's disease.
- 6. Metformin.
- 7. H.pylori.
- 8. PPI/Ranitidine.

B12 deficiency

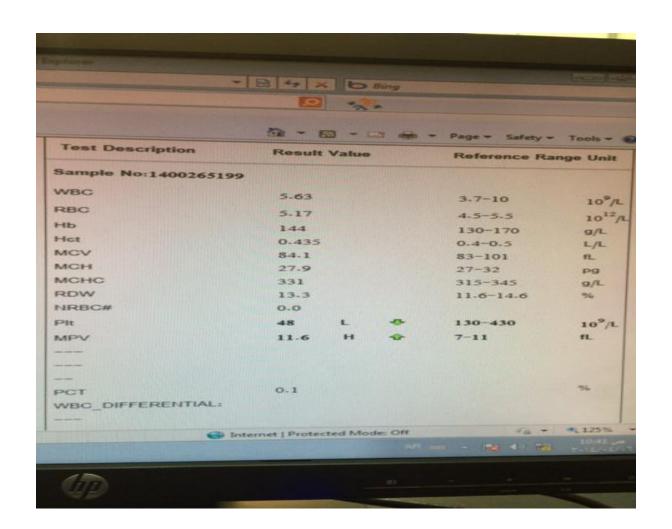
- Treatment:
- 1. Parenteral B12 (hydroxycobalamin), IM initially 1mg 3X/w for 2 weeks then every 2-3 m.
- 2. If dietary → oral B12 supplement.
- 3. Check response by repeating after 8 weeks.

Folate deficiency:

- 1. 5mg folic acid for 4 months.
- 2. If malabsorption \rightarrow increase the dose to 15mg.

CASE 4

- 20 years female complain on heavy periods and easy bruising in the last 1 month. These symptoms proceeded by a viral illness.
- Her CBC showed



- What is the most likely diagnosis?
 - a. ITP
 - b. TTP/HUS
 - c. Dilution thrombocytopenia
 - d. Inherit ant Thrombocytopenia

Manifestations of thrombocytopenia

- Petechial hemorrhage.
- Easy bruising.
- Mucosal bleeding
- e.g. _ epistaxis.
- _ gum bleeding

Causes of Thrombocytopenia

- Decreased production
- Congenital disorders
- Radiation or chemotherapy
- Vitamin B12 or folate deficiency
- Drugs
- Systemic lupus erythematosus
- Aplastic anemia
- Acute leukemia
- Lymphomas
- Alcohol abuse
- Viral infections, including HIV
- Splenic sequestration

CASE 5

- 50 years lady found on routine blood test to have a high platelet count. She is a symptomatic and history was not significant.
- What blood test you will initially ask for?
 - a. CRP
 - b. ESR
 - c. Ferritin
 - d. Blood film
 - e. All of the above

Conditions Associated with Thrombocytosis

- Iron deficiency (overactive reaction)
- Acute blood loss
- Inflammatory disorders (IBD)
- Malignancies
- Splenectomy
- Myeloproliferative disorders
- Essential thrombocythemia
- Polycythemia
- Myelofibrosis

Causes of pancytopenia

- cancers that affect the bone marrow, such as:
 - leukemia
 - multiple myeloma
 - Hodgkin's or non-Hodgkin's lymphoma
 - myelodysplastic syndromes
 - megaloblastic anemia, a condition in which your body produces larger-than-normal, immature red blood cells and you have a low red blood cell count
- <u>aplastic anemia</u>, a condition in which your body stops making enough new blood cells
- paroxysmal nocturnal hemoglobinuria, a rare blood disease that causes red blood cells to be destroyed
- viral infections, such as:
 - Epstein-Barr virus, which causes mononucleosis
 - cytomegalovirus
 - HIV
 - hepatitis
 - malaria
 - sepsis (a blood infection)

- diseases that damage bone marrow, such as <u>Gaucher disease</u>
- damage from <u>chemotherapy</u> or <u>radiation</u> treatments for cancer
- exposure to chemicals in the environment, such as radiation, arsenic, or benzene
- bone marrow disorders that run in families
- vitamin deficiencies, such as lack of <u>vitamin B-12</u> or <u>folate</u>
- enlargement of your spleen, known as <u>splenomegaly</u>
- liver disease
- excess alcohol use that damages your liver
- autoimmune diseases, such as <u>systemic lupus erythematosus</u>
- In about half of all cases, doctors can't find a cause for pancytopenia. This is called idiopathic pancytopenia.

CAUSES OF POLYCYTHEMIA

- Polycythaemia can cause <u>blood clots</u>. These put you at risk of life-threatening problems such as:
- <u>pulmonary embolisms</u> a blockage in the blood vessel that carries blood from the heart to the lungs
- deep vein thrombosis (DVT) a blockage that forms in the blood vessels in your leg before moving elsewhere in your body

 Polycythaemia also increases your risk of heart attack and stroke. Seek emergency medical help if you think that you or someone you're with is having a <u>heart attack</u> or <u>stroke</u>.

- Apparent polycythaemia" is where your red cell count is normal, but you have a reduced amount of a fluid called plasma in your blood, making it thicker.
- Apparent polycythaemia is often caused by being overweight, smoking, drinking too much alcohol or taking certain medicines – including diuretics (tablets for high blood pressure that make you pee more).
- Apparent polycythaemia may improve if the underlying cause is identified and managed. <u>Stopping smoking</u> or reducing your alcohol intake, for example, may help.

Relative polycythaemia

 This is similar to apparent polycythaemia. It can happen as a result of dehydration.

- Absolute polycythaemia
- "Absolute polycythaemia" is where your body produces too many red blood cells. There are 2 main types:
- primary polycythaemia there's a problem in the cells produced by the bone marrow that become red blood cells; the most common type is known as polycythaemia vera (PV)
- secondary polycythaemia too many red blood cells are produced as the result of an underlying condition

- Secondary polycythaemia
- Secondary polycythaemia is where an underlying condition causes more erythropoietin to be produced.
 This is a hormone produced by the kidneys that stimulates the bone marrow cells to produce red blood cells.
- Health conditions that can cause secondary polycythaemia include:
- chronic obstructive pulmonary disease (COPD) and sleep apnoea – these can cause an increase in erythropoietin, due to not enough oxygen reaching the body's tissues
- a problem with the kidneys such as a <u>kidney tumour</u> or narrowing of the arteries supplying blood to the kidneys

LFT

PETOR: SHERRILL AST PROCEDURES	RESULTS		REFERENCE RAN
TOTAL HEALTH PLUS			
CHEM 27		2073	
ALK. PHOSPHATASE -	99	(H)	0-62
ALT (SGPT)	241	(H)	28-75
AST (SCOT)	80	(H)	5-55
CK —	1058	(H)	64-440
GCT	2		1-7
AMYLASE	733		500-1500
LIPASE	118		10-195
ALBUMIN	2.8		2.4-4.1
TOTAL PROTEIN	6.9		5.9-8.5
GLOBUL IN	4.1		3.4-5.2
TOTAL BILIBUBIN	0.2		0.0-0.4
DIRECT BILIRUBIN	0.0		0.0-0.1
BUN	17		15-34
CREATININE	0.8		0.8-2.3
CHOLESTEROL	152		82-218
GLUCOSE	98		70-150
CALCIUM	10.0		7.5-10.8
PHOSPHORUS	5.9		3.0-7.0
TCO2 (BICARBONATE)	11	(L)	13-25
CHLORIDE	117		111-125
POTASSIUM	4.6		3.9-5.3
SODIUM	155		147-156
A/G RATIO	0.7		0.6-1.5
B/C RATIO	21.3		
INDIRECT BILIRUBIN	0.2		0-0.3
NA/K RATIO	34		
ANION GAP	32	(H)	13-27

- Abnormal LFT (AST fatty liver, ALT autoimmune/viral, GGT alcohol)
- Don't miss hormonal supplements in adults!!!!
- High LFT once → repeat
- >/= 3X upper limit of normal:

Repeat after 1 week

< 3X upper limit of normal:

Repeat after 1 month

- If still high LFT results → refer for US and hepatitis screen.
- If ALT < 2X ULN(upper limit of normal) + normal
 US + -ve Hep. Screen → repeat after 3-6 months.
- If US shows fatty liver
- a. Alcohol consumer → stop alcohol then repeat after 3-6 months.
- b. Non alcohol consumer → treat metabolic syndrome then repeat after 3-6 months.

ALP

- Elevation of alkaline phosphatase is seen in normal childhood and adolescence, as well as pregnancy.
- Nonspecific and can be seen with liver or bone disorders or can be related to medication.

GGT

 GGT is a microsomal enzyme that is inducible by alcohol and certain drugs, including warfarin and some anticonvulsants.

Hepatitis Serology

- Hepatitis A virus (HAV), hepatitis B virus (HBV) and, less often, hepatitis C virus (HCV) are the usual causes of acute viral hepatitis.
- A person with symptoms of acute hepatitis should have the following three hepatitis serologies performed
- IgM anti-HAV
- hepatitis B surface antigen (HBsAg)
- anti-HCV (PCR)
- Presented in microbiology section

Hyperurecemia

- Deposition of monosodium urate monohydrate in the synovium.
- Uric acid > 450 Mmol/l.
- Management (RAPRIOP):

R reassurance: resolves within 2 weeks.

A advice: reduce alcohol intake/loose weight if obese/avoid food rich in purines (liver, yeast, kidney, seafood, oily fish e.g. sardines)/Aspirin at low dose increase gout (safe dose 75-150mg).

P prescription:

- 1. NSAIDs.(It's not an option in uncontrolled hypertensive patients)
- 2. Intraarticular cortisone injection. .(It's not an option in uncontrolled hypertensive patients)
- 3. Colchicine (lower onset of action) → SE diarrhea 500Mcg bd max 6mg.(only can be used with uncontrolled hypertensive patients)
- 4. Oral steroids 15mg od.
- 5. If no response → allopurinol (zyloric 100mg)or febuxostat(adenouric 80mg od). (start after 2 weeks of settling of symptoms → increase dose of zyloric till uric acid < 300Mmol

R referral: if hyperurecemia with ureate stone and recurrent UTI \rightarrow urology.

investigations:

- 1. Uric acid.
- 2. ESR (high).
- 3. WBS (high).
- 4. Microscopy of synovial fluid (needle like crystals).
- 5. X-ray (soft tissue swelling/erosion if severe).
- O observation: after 2 weeks of NSAID to see response.
- P prevention: (indication for allopurinol)
- 1. Recurrent attacks >/= 1
- 2. Tophi.
- 3. Renal disease.
- 4. Uric acid renal stones.
- 5. Prophylaxis if on cytotoxic or diuretics.
- 6. Radiological findings of chronic disease.
- 7. Urine uric acid > 1100 mg/dl (6.5mmol).

FBS, RBS, HbA1c, IFG, IGT

- To diagnose diabetes (TWO READING OR ONE READING WITH SYMPTOMS)
- 1. HbA1c > /= 6.5%
- 2. FBS >/= 7mmol/l
- 3. RBS >/= 11.1 mmol/l

Prediabetes cut points:

- 1. IFG 5.6-6.9 mmol/l
- 2. IGT 7.8-11 mmol/l
- 3. HbA1c 5.7-6.4%

- GESTATONAL DM (OGTT)
- FBS 5.1
- AFTER 1 HR 10
- AFTER 2HR 8.5

Prostate specific antigen

- 3 Ng/ml from 50-59y
- 4 Ng/ml from 60-69
- 5 Ng/ml > 70 y

In older men the prevalence is:

50% at the age of 50 y.

80% at the age of 80 y.

Voiding symptoms/storage symptoms (irritation)/ post micturition.

Complications:

- 1. Recurrent UTI.
- 2. Retention of urine.
- 3. Obstructive uropathy.

Cases of raised PSA:

- 1. BPH.
- 2. Prostatitis and UTI (1 m after treatment).
- 3. Ejaculation (48hr).
- 4. Exercise (48hr).
- 5. Urinary retention.
- 6. Instrumentation of the urinary tract.

Management:

Watchful waiting

Alpha antagonist (tamsulosin (Omnic 0.4mg)/alfuzosin (xatral 10mg) → decrease symptoms.

SI: retrograde ejaculation, postural hypotension

5 alpha reductase (finasteride/Proscar 5mg)

decrease progression (needs 6 m to show proper effect, decrease PSA by 50%)

Subclinical hypothyroidism

- High TSH & normal T4.
- Presence or absence of mild symptoms of hypothyroidism.
- Risk of progression to overt hypothyroidism is 2-5% per year (higher in men).
- Risk increased by presence of thyroid autoantibodies (thyroid peroxidase AB) thyroglobulin AB used for thyroid cancer follow up
- Treat subclinical hypothyroidism patient if:
- 1. TSH > 10.
- 2. Thyroid autoantibodies +ve.
- 3. Other autoimmune disorders.
- 4. Previous treatment of Graves's disease.
- 5. Symptomatic.(wt gain, cold intolerance, bradycardia or constipation)
- 6. Hypercholesterolemia.

FT3	FT4	TSH	Comment
↑	1	↓	1ry Hyperthyroidism
↑	↑	1	2ry Hyperthyroidism
\	↓	1	1ry Hypothyroidism
\	↓	↓	2ry Hypothyroidism

Renal function test and eGFR

CR, UREA (hormonal supplements in adults!!

Table 10. Stages of Chronic Kidney Disease

Stage	Description	GFR (mL/min/1.73 m ²)
1	Kidney damage with normal or ↑ GFR	≥90
2	Kidney damage with mild ↓ GFR	60–89
3	Moderate ↓ GFR	30–59
4	Severe ↓ GFR	15–29
5	Kidney failure	<15 (or dialysis)

Chronic kidney disease is defined as either kidney damage or GFR <60 mL/min/1.73 m² for \geq 3 months. Kidney damage is defined as pathologic abnormalities or markers of damage, including abnormalities in blood or urine tests or imaging studies.

- Microalbuminuria (MA) is defined as a persistent elevation of albumin in the urine of (>20 to <200 microg/min).
- Other tests section
- In diabetic patients treat with ACE INH even if the patient is not hypertensive.

Autoimmune Diseases

Routine clinical tests:

- ESR and CRP; 个个 in inflammation
- CBC: anemia and ↓ platelet and ↓ WBCs
- routine chemistry panels which may reveal:
 - kidney involvement; ↑ ↑ BUN & creatinine
 - abnormalities of liver function tests
 - − ↑ muscle enzymes (CPK)

Lab Diagnosis of RA:

- Rheumatoid factor: -ve in 30 % early in illness; can repeat 6 - 12 ms; +ve in numerous other processes (e.g., lupus; scleroderma; Sjögren's syndrome; neoplastic disease); not an accurate measure of disease progression.
- Anticyclic citrullinated peptide (Anti-CCP) antibody: correlate well with disease progression; increases sensitivity when used in combination with RF; more specific than RF.
- Antinuclear antibody (ANA): Limited value for screening
- Complement levels: Normal or elevated
- <u>Urinalysis:</u> hematuria or proteinuria

Lab Diagnosis of SLE:

- Anti-nuclear antibody (ANA)
- Anti-DNA antibody: to determine Abs to the genetic material in the cell
- Anti-Sm antibody: a ribonucleo-protein found in the cell nucleus
- Complement proteins C3 and C4

Vit D def

- Management plan: (vit d 50,000)
- Less than 50 one tab every one week for 8 wks then one tab monthly for 4 mths then repeat test if becomes normal from(75 to 250) preventive dose every 2 mths better to be taken for long life
- Above 50-75 one tab taken monthly for 6 mths then repeat test and if it becomes normal preventive dose better to be taken every 2 mths

Vit B12 def

- Methylcobal injection (b12 injection)better than neurobion because it's not painful taken E.O.D for 2 wks 6 injection 1gm then tablets taken for 2 to 3 months.
- Stress on diet rich in b12

Roaccutane

- Some tests should be done monthly to follow patient taken roaccutane
- Liver profile , lipid profile(TG), renal profile
- SI: dryness, depression and high lipid and liver profile

Hormonal profile

- (FSH, LH, PROLACTIN AND ESTRAIDOL)
- Should be done 2nd or 3rd day of period look for follicular phase normal FSH higher than LH.
- If LH higher than FSH think in PCO refer for assessment(Rotterdam Criteria)
- High FSH and LH with LOW Estraidol in late thirty females think in premature ovarian failure after repeating the test in 4-6 wks

Stool analysis

- Entameba histolytica treated by flagyl 500mg tds for 5 days
- Round worms(Ascaris) treated by mebendazole(vermox) 100mg bd for 3 days
- Not given below 2yrs
- Thread worms(pin worms) caused by enterobius vermicularis treated by single dose of vermox the dose may need to be repeated in 2weeks if infection persists

 Consider treating all household contacts as threadworms are highly transmissible.

Hair fall

- Tests are recommended for hair fall are Ferritin should be more than 70 normal range from (11-250), CBC, Zinc and TSH.
- Normally 100 hairs fall out per day in females and about 40 in males.