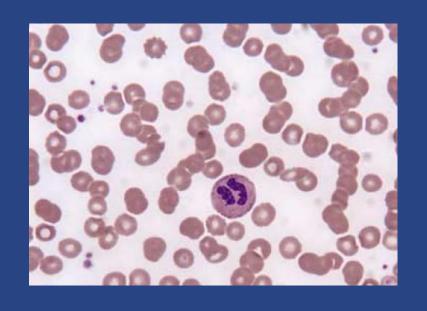


## **Evaluation of the Anemic Patient**



Brenda Shinar, MD, FACP September 14, 2021

# Objectives:

Understand the basics of hematopoiesis.



- Define anemia and understand how to classify it based on the kinetic model and the morphologic model.
- Understand how to calculate the absolute reticulocyte count and absolute reticulocyte index and what these values mean.
- Understand the differentials of microcytic, normocytic, and macrocytic anemias and how to appropriately evaluate them to make a diagnosis.

# Red Blood Cell Factory: What is Required?



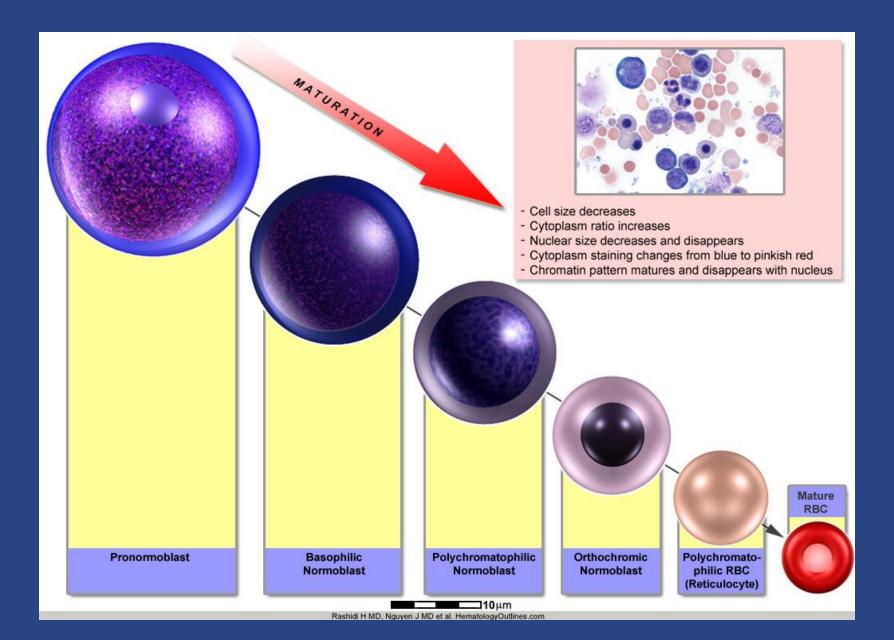
- Work space
- Task master
- Building blocks
- Instructions
- Distribution
- Survival

- Bone marrow
- Erythropoietin
- Precursor cell, heme, iron, B12, folate
- Cytokines
- Exit out of bone marrow
- No lysis, sequestration, or loss

#### Lymphoid Myeloid stem cell stem cell Pluripotential stem cell BFU-E CFU-GEMM V Pre-B V Pre-T CFU-MEG CFU-GM CFU-EO CFU-BASO CFU-E CFU-G CFU-M Pronormoblast Megakaryoblast Myeloblast Monoblast Myeloblast Myeloblast B lymphoblast T lymphoblast Basophilic Megakaryocyte Promyelocyte Promonocyte Promyelocyte Promyelocyte B lymphocyte T lymphocyte normoblast Myelocyte Monocyte Eosinophilic Basophilic Plasma cell Polychromatic myelocyte myelocyte Platelets normoblast Metamyelocyte Eosinophilic Macrophage Basophilic Orthochromic myelocyte myelocyte normoblast Band Basophilic Eosinophilic band Polychromatic erythrocyte (reticulocyte) Polymorphonuclear neutrophil Eosinophil Basophil Erythrocyte Mast cell Figure 2-1 Chart of hematopoiesis.

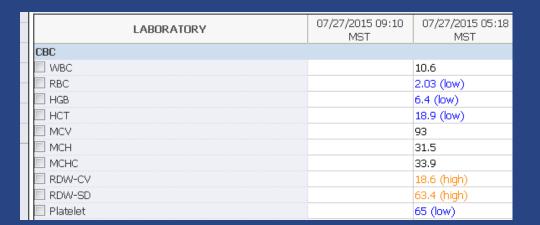
# The Red Cell Family Tree

### The Red Cell Maturation: Benjamin Button



# The Complete Blood Count (CBC)

- Hemoglobin (g/dL)
- Hematocrit (percent)
- Red Blood Cell Count (RBC) (million/mm3)
- Mean Cell Volume (MCV) (fL)
- Red Cell Distribution Width
  - Standard deviation of MCV/Mean MCV x 100
  - CV: coefficient variation
  - SD: standard deviation
- MCH: mean cell hemoglobin (pg)
- MCHC: mean cell hemoglobin concentration (g/dL)



## Normal values for red blood cell parameters in men and women

Red cell parameter	Adult men		Adult women
Hemoglobin, g/dL	15.7 ± 1.7		13.8 ± 1.5
Hematocrit, percent	46.0 ± 4.0		40.0 ± 4.0
RBC count, million/µL	$5.2 \pm 0.7$		4.6 ± 0.5
Reticulocytes, percent	$1.6 \pm 0.5$		1.4 ± 0.5
Mean corpuscular volume, fL		88.0 ± 8.0	
Mean cell hemoglobin, pg/RBC		30.4 ± 2.8	
Mean cell hemoglobin concentration, g/dL of RBC		34.4 ± 1.1	
Red cell volume distribution width, percent (RDW)		13.1 ± 1.4	

Values are mean ± 2 standard deviations. Adapted from Williams' Hematology, 6th ed, Beutler, E, Lichtman, MA, Coller, BS, et al (Eds), McGraw-Hill, New York, 2001.

# Ratio of RBCs to Plasma Determines Hgb, Hct, and RBC count

- Acute severe bleeding may cause orthostatic changes but gives falsely normal values of Hgb, Hct, and RBC count because the ratio of RBCs and plasma lost is the same
- Pregnancy increases plasma volume by 20-25% and therefore lowers the Hgb, Hct, and RBC count
- Dehydration decreases the plasma volume and artificially increases the Hgb, Hct, and RBC count

# Evaluating the Anemic Patient: Kinetic Model:

How the bone marrow is responding to the anemia

# **Decreased Production**Absolute Reticulocyte Index < 2

- Bone marrow disorders or suppression
- Lack of erythropoietin
- Lack of iron, B12, or folate
- Anemia of chronic disease

# Increased Destruction or Blood Loss

Absolute Reticulocyte Index > 2

- Blood Loss
- Congenital Destruction
  - Membrane defects
  - Hemoglobinopathies
  - Enzyme deficiencies
- Acquired Destruction
  - Autoimmune HA
  - Microangiopathic HA
  - Infections (Malaria)
  - Paroxysmal Nocturnal Hemoglobinuria

### **Kinetic Model:**

How is the bone marrow responding to the anemia?

Pt Hct: 30% Normal Hct 45%

- 1) Reticulocyte Count= 2.8%
- 2) Absolute Retic Count=
  - 2.8 x pt's hct/nl hct=
  - $2.8 \times 30/45 = 1.24$
- 3) Absolute Retic Index=

Absolute Retic Ct / Maturation factor



Maturation Factor corrects for the length of time a reticulocyte is in circulation so as not to count it twice or more. This factor varies with the degree of anemia.

$$HCT \ge 35\% = 1.0$$
  
 $25-35\% = 1.5$   
 $20-25\% = 2.0$   
 $< 20\% = 2.5$ 

# Evaluating the Anemic Patient: The Morphologic Model

# Microcytic Anemia: MCV <80 fL

- Reduced iron availability
  - Iron deficiency
  - Anemia of chronic disease
- Reduced heme synthesis
  - Lead poisoning
  - Congenital or acquired sideroblastic anemia
- Reduced globin production
  - Thalassemias
  - Other hemoglobinopathies





### ID and CC:

22-year-old previously healthy white female with fatigue and dyspnea with exertion. She is on an OCP and has light, regular periods once a month. She has never been pregnant. In the ED, she has been given 2 units of pRBC and admitted to your service.

### Labs:

WBC 5.0 g/dL Hemoglobin 5.8 g/dL Platelets 450K MCV 70 fL RDW 18 (11-16) Which of the following is the most appropriate next step to evaluate the cause of her anemia?

- A. Order ferritin, TIBC, and % sat
- B. Order bone marrow biopsy
- C. Order hemoglobin electrophoresis
- D. None of the above are correct choices

> Transfusion. 1992 Oct;32(8):764-5. doi: 10.1046/j.1537-2995.1992.32893032107.x.

Transfusion

# The effects of blood transfusion on serum ferritin, folic acid, and cobalamin levels

C H Ho 1

Affiliations + expand

PMID: 1412686 DOI: 10.1046/j.1537-2995.1992.32893032107.x

### Abstract

Thirty patients with anemia of various causes received packed red cell transfusions. To evaluate the influence of blood transfusion on the serum levels of different nutrients, serum ferritin, folate, and cobalamin levels were determined before and 2, 2 to 4, 5 to 7, 8 to 10, and 11 to 14 days after blood transfusion. No significant change was found in these levels before or at any time after blood transfusion. Blood transfusion exerted little effect on the serum levels of ferritin, cobalamin, or folate. Moreover, if the blood for testing was drawn 1 to 14 days after transfusion, nearly all cases of nutritional deficiency anemia could have been diagnosed without the influence of blood transfusion.

# What are the positive predictors of GI lesions in iron deficient premenopausal women?

- Symptoms of heartburn, regurgitation, or dyspepsia (OR 3.76 p=.002)
- 2. MCV < 70 fL (OR 1.88 p=.04
- 3. Hemoglobin < 10 g/dL(OR 1.7 p=.05)

NEGATIVE risk factor: Heavy menstrual blood loss

 $(OR_{0.46} p=.002)$ 



- ID and CC:

   45-year-old man admitted
   with SIRS due to a UTI and
   chronic stage 4 sacral decub
- PMH
  Paraplegic due to GSW
  Noncompliance with urinary self-catheterization
  Sacral decubs
  Bipolar disorder
  Anemia

Labs:
 Hgb 8.0 g/dL
 MCV 75 fL
 RDW 12.0 (11-16)
 Ferritin 250 ng/mL
 Transferrin 100 mg/dL (188-341 mg/dL)
 Percent saturation 8%

## What is the most likely cause of his anemia?

- A. High hepcidin, Anemia of Inflammation
- B. Low hepcidin, Anemia of Inflammation
- C. Iron deficiency anemia
- D. Undiagnosed thalassemia

# Laboratory Tests in Iron Deficiency of Increasing Severity

	Normal	Fe deficiency without anemia	Fe deficiency with mild anemia	Severe Fe deficiency with severe anemia
Marrow reticulo- endothelial iron	2+ to 3+	None	None	None
Serum iron, µg/dL	60 to 150	60 to 150	<60	<40
Iron binding capacity (transferrin), μg/dL	300 to 360	300 to 390	350 to 400	>410
Saturation (SI/TIBC), percent	20 to 50	30	<15	<10
Hemoglobin, g/dL	Normal	Normal	9 to 12	6 to 7
Red cell morphology	Normal	Normal	Normal or slight hypochromia	Hypochromia and microcytosis
Plasma or serum ferritin, ng/mL	40 to 200	<40	<20	<10

Serum ferritin  $\leq 30$  ng/dL = Iron deficient (PPV 83%, PLR= 11) Serum ferritin  $\geq 100$  ng/dL = Iron sufficient (NLR .08)

What about if the ferritin is between 30 and 100?

# Other Tests to help distinguish IDA from ACD...

- Transferrin: (TRUCKS)
  - High in IDA
  - Low or low normal in ACD
- Low reticulocyte-hemoglobin concentration (RET-He) increase with IV iron
  - Does not distinguish
     between ACD and IDA but
     can improve in 2-3 days
     after IV iron
     supplementation to prove
     response to iron therapy

Retic %	1.4
Retic #	51
Immature Retic Fraction (IRF)	21.2 * H
Retic Hgb Equivalent [RET-He]	21.0 * L
🔲 Iron	23 L
Transferrin	185 L
🔲 Trans % Sat	9.8 L

# Distinguishing between ACD and IDA in NON-dialysis patients

- A ferritin of > 100 makes iron deficiency statistically unlikely even in patients with chronic inflammation!
- A transferrin level that is low gives further evidence of anemia of chronic disease.
- A low serum iron level and percent saturation cannot distinguish between ACD and IDA!
- The reticulocyte-Hgb content may be a new way to determine response to IV iron therapy.



- ID and CC:
  - A 42-year-old black woman with a history of iron deficiency on iron therapy comes to see you as a new patient.
- She was told that she was iron deficient 10 years ago and has been on iron supplementation for as many years.
- Labs:
   Hemoglobin 11.0 g/dL
   MCV 72 fL
   RDW 11 (11-16)
   Ferritin 800 ng/mL

You suspect a thalassemia and order a hemoglobin electrophoresis which comes back normal.

What is the most likely diagnosis?

- A. She was iron deficient but is now iron replete
- B. She has B-thalassemia trait
- C. She has A-thalassemia trait
- D. She has a myelodysplastic syndrome

## Thalassemias

 Found most frequently in the Mediterranean, Africa, Western and Southeast Asia, India, and Burma

 Distribution parallels that of Plasmodium Falciparum



### The thalassemias: Genetic, clinical, and laboratory findings

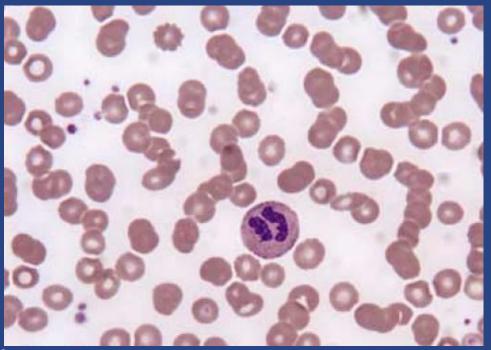
Disorder	Genotype	MCV	Anemia	Hb electrophoresis
Alpha thalassemia				
Silent carrier	a a / a -	NL	None	Normal <3 percent Hb Barts at birth
Trait	a a / or a - / a -	Low	Mild	Normal 3-8 percent Hb Barts at birth
Hb H disease (deletional)	a - /	Low	Moderate	5-30 percent Hb H present in adults 20-40 percent Hb Barts at birth
Major (fetal hydrops)	/	Low	Fatal	Hb Barts, Hb Portland, and Hb H present Hb A, Hb F, and Hb A2 are absent
Beta thalassemia			•	
Trait	β / β °	Low	Mild	Hb A2 increased (3.5-7 percent)
Intermedia	$_{\beta}$ $^{+}$ / $_{\beta}$ $^{+}$ and others*	Low	Moderate	Hb F increased in about 50 percent of patients
Major	β°/β°	Low	Severe	Hb A absent Only Hb A2 and Hb F are present

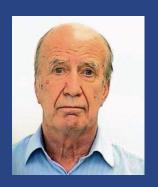
MCV: mean corpuscular volume; Hb: hemoglobin; NL: normal;  $_{\beta}$  <sup>+</sup>: thalassemic gene producing some  $_{\beta}$  -chain;  $_{\beta}$  °: thalassemic gene producing no  $_{\beta}$  -chain. \* See text for multiple other genotypes. Courtesy of Stephen A Landaw, MD, PhD.

# Evaluating the Anemic Patient: The Morphologic Model

# Normocytic Anemia: MCV 80-100

- Early iron deficiency
- Anemia of chronic disease
- Bone marrow disease
  - Invasion, aplasia
  - Myeloma, MGUS
- Chronic renal insufficiency
- Endocrine dysfunction
  - Hypothyroid
  - Hypopituitarism
- Combined disorders in patients with high RDW
  - ACD or IDA +B12 or folate deficiency





ID and CC

75-year-old man with fatigue and palpitations admitted with anemia and oliguric renal failure. He has had some severe back pain for a month and physical therapy visits have not helped.

• Labs:

Hemoglobin 7.2 g/dL

MCV 84 fL

**RDW 12** 

Creatinine 3.2 mg/dL Total Protein 9.1 g/dL Albumin 2.5 g/dL

Urine protein/creatinine ratio: 4.5

Which of the following laboratory tests should be ordered next to help make the diagnosis?

- A. Serum protein electrophoresis (SPEP)
- B. Serum immunofixation (IFE)
- C. Serum free light chains
- D. B2-glycoprotein I
- E. A,B, and C

# Diagnosis of Multiple Myeloma

### Table 1. Findings on Presentation for Patients with Multiple Myeloma

Symptom or laboratory finding	Percentage of patients
Anemia (hemoglobin < 12 g per dL [120 g per L])	73
Bone pain	58
Elevated creatinine (> 1.3 mg per dL [115 µmol per L])	48
Fatigue or generalized weakness	32
Hypercalcemia (calcium > 10.1 mg per dL [2.52 mmol per L])	28
Weight loss	24

### **Table 2. Diagnostic Tests for Multiple Myeloma**

Initial tests	Confirmatory tests
Complete blood count	24-hour urine protein
with differential	Beta <sub>2</sub> -microglobulin
Serum albumin, calcium, creatinine, electrolytes, and urea nitrogen	Lactate dehydrogenase
	Serum free light chain assay
	Serum immunofixation electrophoresis
	Serum protein electrophoresis*
	Serum quantitative immunoglobulins
	Skeletal survey†
	<b>Orine immunofixation electrophoresis</b>
	Urine protein electrophoresis



### ID and CC:

35-year-old woman with Crohn's disease is admitted with abdominal pain and bloody stools consistent with a flare of her disease for the past week. She has had several abdominal surgeries for her disease.

### Labs:

Hemoglobin: 7.9 g/dL

MCV: 85 fL RDW: 19

Ferritin: 8

B12: 211

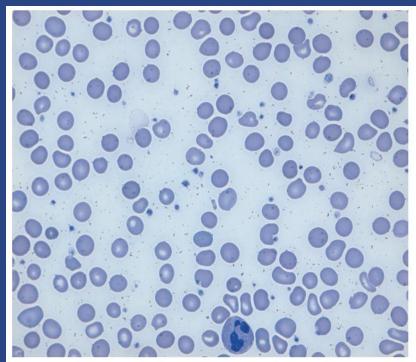
Absolute retic index: 1.2

What is the most likely cause of this patient's anemia?

- A. Iron deficiency anemia
- B. B12 deficiency
- C. A and B
- D. Bone marrow suppression

# Multifactorial Causes of Anemia Resulting in Normocytic MCV with Elevated RDW

- MCV normal + RDW high
- Check retic count, iron studies, ferritin, B12 and folate
- Peripheral blood smear may give clues as well



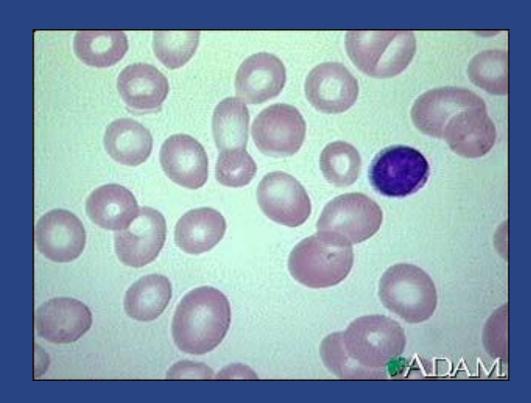
**Fig. 1.** Peripheral blood smear shows hypersegmented neutrophils and normocytic hypochromic red blood cells (Wright stain, × 1,000 magnification).

Spivak JL. Masked megaloblastic anemia. Arch Intern Med. 1982 Nov;142(12):2111-4. PMID: 7138159.

# Evaluating the Anemic Patient: The Morphologic Model

# Macrocytic Anemia MCV > 100

- Ethanol abuse
- Drug-induced
  - AZT, hydroxyurea
- Liver disease
- Hypothyroidism
- Reticulocytosis
  - Hemolysis
  - Blood loss
- Myelodysplastic syndromes
- Folate deficiency
- B12 deficiency





### ID and CC:

42-year-old man with Hep C and decompensated liver cirrhosis is admitted for encephalopathy. He has no signs of GI bleeding.

What is the most appropriate lab test to order next?

### • Labs:

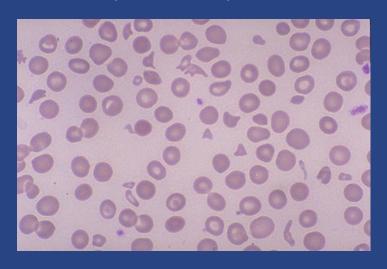
Hemoglobin 7.0 g/dL
Platelets 52K
MCV 107 fL
RDW 20 (11-16)
Absolute retic index: 2.5

Ferritin 1700

- A. Peripheral smear
- B. B12 and folate
- C. Bone marrow biopsy
- D. LDH and haptoglobin
- E. Direct Coomb's test

# Hemolytic Anemia

- Combination of normal LDH and haptoglobin rules OUT hemolysis with 92% sensitivity.
- Combination of elevated LDH and low haptoglobin rules IN hemolysis with 90% specificity.



- Peripheral smear can help to guide the work up once hemolysis is confirmed:
- Schistocyte
  - (mechanical shear)
- Spherocyte
  - (autoimmune)
- Acanthocyte
  - (liver disease)
- Bite cell
  - (oxidant damage)

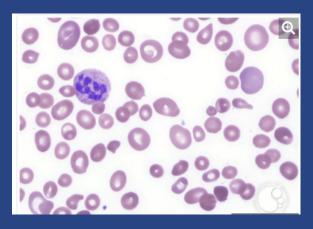


### ID and CC:

59-year-old man with progressive dyspnea with exertion over 3 years

### Labs:

Hemoglobin 4.7 g/dL MCV 118 fL Creatinine 1.6 mg/dL Platelets 97 K WBC 2.3 K Retic count 7.5 LDH 450 Haptoglobin <10 A peripheral smear is shown:



What do you want to order next to evaluate the cause of his anemia?

- A. B12
- B. ADAMSTS13
- C. Methylmalonic acid
- D. Bone Marrow Biopsy

### Causes of vitamin B12 deficiency

- · ·		
Gastric	abnor	malifies

Pernicious anemia

Gastrectomy/Bariatric surgery

Gastritis

Autoimmune metaplastic atrophic gastritis

#### Small bowel disease

Malabsorption syndrome

Ileal resection or bypass

Crohn's disease

Blind loops

#### **Pancreatitis**

Pancreatic insufficiency

#### Diet

Strict vegans

Vegetarian diet in pregnancy

### Agents that block absorption

Neomycin

Biguanides (eg, metformin)

Proton pump inhibitors (eg, omeprazole)

N2O anesthesia inhibits methionine synthase

Inherited transcobalamin II deficiency



## Practice Question #1

### 61 year old AA man

	WBC	14.8 H
	RBC	1.88 L
	HGB	5.4 ' C
	HCT	15.1 ' C
	MCV	80
	MCH	28.7
	MCHC	35.8
	RDW-CV	17.8 H
	RDW-SD	51.3 H
	Platelet	162
	Immature Platelet Fraction (IPF)	
	MPV	11.6
Pla	atelet Estimate	
	Retic Hgb Equivalent (RET-He)	32.7 *
Di	ff Type	AUTO
	Immature Granulocyte % (IG%)	1.0 *
	Immature Granulocyte # (IG#)	0.2 H
	Segs %	40
	Lymphs %	38
	Monos %	11
	Eos %	7
	Basos %	3 H
	Neut#	5.9
	Lymph#	5.6 H

- Retic count: 4.7%
- Retic index: 0.7 %
- B12: 910 pg/mL
- RBC Folate: 971 ng/mL
- Ferritin: 7,650 ng/mL

HEMOGLOBINOPATHY EVALUATION		
Helec RBC	2.84 L	
Helec HGB	8.3 L	
Helec MCV	84	
Helec MCH	29.2	
Helec RDW-CV	17.1 H	
Helec RDW-SD	51.8	
Hgb A	92.4 L	
Hgb S	5.0 H	
Hgb A2	2.6	
Helec Interp	Helec Interp	
GENERAL COAGULATION		

Hemoglobin SS disease

## Practice Question #1

 Differential diagnosis for a patient with sickle cell disease who presents with Hgb 5 and low retic index?

- Parvo B19
- Aplastic crisis from folate deficiency
- ? Epopoietin deficiency

CHEM GENER	
Specimen Integrity  Glucose Level	120 * H
BUN	41 H
Creatinine	2.47 * H
Estimated Glomerular Filtration Rate	28 * L
BUN/Creat Ratio	17
Sodium	143
Potassium	4.1
Chloride	111 H
C02	22
Anion Gap	10
Magnesium	
Calcium	8.2 L
Phosphorus	
Protein, Total	6.5
Albumin Albumin	2.3 L
Alb/Glob Ratio	0.5 L
Bilirubin Total	3.4 H
AST	98 H
ALT	49
Alkaline Phos	252 H

 Epo level: 5.0 mIU/mL (2.6-18.5)

# Objectives:

Understand the basics of hematopoiesis.



- Define anemia and understand how to classify it based on the kinetic model and the morphologic model.
- Understand how to calculate the absolute reticulocyte count and absolute reticulocyte index and what these values mean.
- Understand the differentials of microcytic, normocytic, and macrocytic anemia and how to appropriately evaluate them to make a diagnosis.