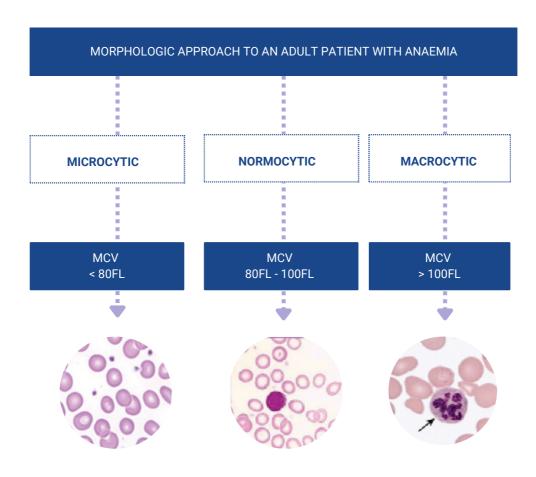
# Newsletter <

## APPROACH TO ADULT PATIENT WITH ANEMIA

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Anaemia is defined as a reduction in one or more of the red blood cell indices of Hemoglobin (Hb) concentration, Hematocrit, and Red blood cell (RBC) count. However, Hemoglobin concentration is the most widely used parameter.

- Normal lower limits of "Hb" varies with age, sex and race.
- As a Clinician, one is often familiar with the Kinetic Approach to a patient with Anemia.
- This approach classifies Anemia by the mechanisms causing the Anemia. These mechanisms are decreased red blood cell production, increased red blood cell destruction and blood loss.



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#### MICROCYTIC ANEMIA

CONDITION	DIAGNOSIS/ CAUSES	MECHANISM	CONDITION CLINICAL SIGNS	PERIPHERAL BLOOD SMEAR	ADDITIONAL INVESTIGATION
IRON DEFICIENCY	Blood Loss:  Menorrhagia  GIT bleed (Hemorrhoids, Peptic Ulcer, etc)  Bleeding Disorders	Inadequate supply of iron for Erythropoiesis	Koilonychia Angular Cheilosis Glossitis	Hypochromia Poikilocytoses Elliptocytes	Iron studies: Serum iron ↓ Serum Ferritin ↓ Transferrin ↑
	Malabsorption:     Gluten Enteropathy     Pregnancy     Poor dietary intake				
THALASSEMIA'S	Alpha Thalassemia  Beta Thalassemia	Disorders of globin production	Hepatosplenomegaly Thalassemic facies:	Hypochromia Tear drops Target cells Basophilic stippling	Hemoglobin electrophoresis HPLC Hb quantification
	Acquired primary: Refractory Anemia with ring sideroblasts (MDS)	Clonal disorder with impaired Haem synthesis	Hypochromia Target cells Basophilic stippling Poikilocytosis Pappenheimer bodies Red cells can be microcytic or macrocytic		Iron studies: Serum Iron ↑ Ferritin ↑ Bone marrow Aspirate
SIDEROBLASTIC ANEMIA	Acquired Secondary:  Drugs Isoniazid, Pyrazinamide, Chloramphenicol, Alcohol, Lead toxicity, Copper deficiency.	Underlying molecular defects affecting synthesis of the haem group	Mitochondrial toxicity/impaired haem synthesis	Red cells can be microcytic or macrocy	rtic
	<ul><li>Congenital:</li><li>·X-linked Autosomal</li></ul>	Underlying molecular defects affecting synthesis of the Haem group	Syndromic features	Hypochromia Target cells Basophilic stippling Poikilocytosis	Biochemical assay of enzymes
	Severe cases of Anemia or inflammation or chronic disease	See normocytic Anemias		Pappenheimer bodies	

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#### NORMOCYTIC ANEMIA

CONDITION	DIAGNOSIS/ CAUSES	MECHANISM	CONDITION CLINICAL SIGNS	PERIPHERAL BLOOD SMEAR	ADDITIONAL INVESTIGATION
APLASTIC ANEMIA (AA)	Acquired  • Drugs:     chloramphenicol,     Sulphonamides,     linezolid, diclofenac,     phenytoin,     carbamazepine,     thiouracil     phenothiazides,     chloroquine,     allopurinol,     mebendazole,     thiazides etc.  • Environmental:     benzene, pesticides     cutting oils     lubricating agents     recreational drugs  Inherited  • Congenital bone     marrow failure     syndromes	Inadequate supply of Iron for Erythropoiesis.	Koilonychia Angular chelosis Glossitis	Hypochromia Poikilocytes Elliptocytes	Iron studies: Serum iron ↓ Serum Ferritin↓ Transferrin↑
PURE RED CELL APLASIA	Transient Infections:  Parvovirus B19 CMV HIV Mumps Drugs Chronic Idiopathic  Congenital syndromes: Immunological Thymoma Haematological malignancies Autoimmune diseases	Serum antibodies with selective cytotoxicity for marrow Erythroid cells. Immunological suppression of Erythropoiesis		Normocytosis or Macrocytosis	↓ Reticulocyte count Bone marrow aspirate & trephine biopsy PCR for Parvovirus B19 Screening for auto antibodies
SYSTEMIC DISORDERS CHRONIC DISEASE	Anemia of inflammation (previously called Anemia of chronic disorders) Infections (bacterial, viral and fungal)  1. Acute/Chronic Autoimmune conditions 2. Chronic diseases Ageing	Changes in iron handling/systematic iron distribution under control of hepcidin, red blood cell production due to a "blunted" Erythropoietin (EPO) response and reduced red blood cell lifespan.		Mild-normocytic red cells, severe "Microcytic" red cells Rouleaux formation Neutrophilia or Neutrophilia with toxic granulation and left shift lymphopaenia Thrombocytopaenia	Iron Studies: Serum Iron ↓ Ferritin N/ ↑ Infective markers: ↑ CRP ↑ PCT ↑ ESR Autoimmune screen Viral studies Blood Culture



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### MICROCYTIC ANEMIA

CONDITION	DIAGNOSIS/ CAUSES	MECHANISM	CONDITION CLINICAL SIGNS	PERIPHERAL BLOOD SMEAR	ADDITIONAL INVESTIGATION
CHRONIC RENAL DISEASE		Deficiency of EPO production by the Rena Interstitial cells	al		U&E and eGFR EPO levels
MALIGNANCY		Anemia of inflammatic therapy related bone marrow infiltration	on		Bone marrow biopsy
ACQUIRED ANEMIA IN HOSPITALISED PATIENTS		Blood loss during procedures: Anemia o inflammation changes in nutrition especially i ICU	•		





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