	Classification due to morphology blood film
Anemia of acute blood loss	Normocytic, normochromic RBCs with reticulocytosis
Anemia of chronic blood loss	Microcytic, hypochromic RBCs with low reticulocytes
Iron deficiency anemia	Microcytic, hypochromic RBCs and target cells. • Poikelocytosis (small and normal) • Low reticulocytes • Thrombocytosis
Anemia of chronic disease	Normal morphology then microcytic, hypochromic RBCs with low reticulocytes.
Megaloblastic anemia	Macroovalocyte RBCs and hypersegmented neutrophils (>5 lobes)
Aplastic anemia	 Biopsy reveals an empty, fatty bone marrow. Pancytopenia on peripheral blood smear Anemia is normochromic or macrocytic Decreased hematopoietic cells
Myelophthisic anemia	Leukoerythroblastic anemia (Immature granulocytic and erythroid precursors commonly appear in peripheral blood)

	Note : Myelophthisic refers to the displacement of hematopoietic bone marrow tissue by fibrosis, tumors or granulomas. Features:Shift to left (high levels of immature cells), nucleated RBCs (peripheral blood), Hypercellular bone marrow (but not erythropoietic cells)
Anemia of renal disease	Echinocytes (Burr cells) appear.
Anemia of liver disease	Acanthocytes (spur cells) appear
Anemia of hypothyroidism	Most commonly normocytic but can be macrocytic.
Myelodysplastic syndrome	Macrocytic RBCs.

	Microcytic, hypochromic RBCs with target cell and nucleated
Thalassemia	RBCs.
	• Basophilic stippling (small dots in the cytoplasm of RBCs that are remnants of ribosomes).
	Reticulocytosis
Thalassemia major (Cooley's anemia)	Poikelocytosis, nucleated RBCs (peripheral blood). ↑↑ Normoblasts, filling bone marrow spaces and expanding into bone, hemosiderosis (iron overload).
	Sickle cells and target cells (secondary to abnormal hemoglobinization) are seen on blood smear in sickle cell disease, but not in sickle cell trait.
Sickle cell anemia	
	Oxidative stress precipitated
	hemoglobin as Heinz bodies.
	Heinz bodies are removed from
Intravascular hemolysis due to	RBCs by splenic macrophages.
G6PD deficiency	resulting in bite cells.
	Bite cells Heinz bodies

Immune hemolytic anemia	 IgG mediated disease (<u>Warm type</u>) can lead to extravascular hemolysis. Spherocytes (very small without pale center) and polychromasia (due to reticulocytosis, reticulocytes appear blue). Mild chronic anemia. IgM mediated disease (<u>Cold type</u>) can lead to intravascular hemolysis. RBCs agglutination; IgM binds 5 RBCs thus creating agglutination. Can be acute (associated with Mycoplasma pneumoniae and infectious mononucleosis) Or chronic (associated with B-cell lymphoma).
Heredity spherocytosis	Spherocytes with loss of central pallor. Treatment is splenectomy; anemia resolves, but spherocytes persist and Howell-Jolly bodies (fragments of nuclear material in RBCs) emerge on blood smear.

	Intravascular hemolysis occurs
	episodically, often at night during
Paroxysmal nocturnal	sleep.
hemoglobinuria	RBCs, WBCs, and platelets (Destroyed platelets release cytoplasmic contents into circulation, inducing thrombosis which is the main cause of death) are lysed.
	Schistocytes are seen in
	mechanical valve induced (prosthetic
	heart valves and aortic stenosis) and
	microangiopathic hemolytic
Traumatic hemolysis	anemia which is intravascular
	anemia results from vascular
	pathology; RBCs are destroyed as
	they pass through the circulation.
	Microthrombi produces
	schistocytes on blood smear.

Notes :

- * Microcytic anemia reflects hemoglobinization problem.
- ✤ Macrocytic anemia reflects stem cell disease and maturation.
- * Target cells are found in iron deficiency anemia, thalassemia and sickle cell anemia.