In the Name of GOD

Anemia and Pregnancy



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Perinatalogist

First trimester – Hemoglobin  $\langle \mathcal{G} | g/dL$  (approximately equivalent to a hematocrit  $\langle \mathcal{G} | g \rangle$  percent)

Second trimester – Hemoglobin < $\cdot$ . g/dL (approximate hematocrit < $\forall$ ) or  $\forall$  percent)

Third trimester – Hemoglobin level  $< \mathcal{V} g/dL$  (approximate hematocrit  $< \mathcal{V} percent$ )





World Health Organization (WHO) estimates that over <sup>¢</sup> • percent of pregnant women globally have anemia



#### (WHO):

Severe antenatal or postnatal maternal anemia (of any type) was associated with an increased risk of maternal death . In other studies, maternal anemia was associated with several adverse pregnancy outcomes such as antenatal/postnatal sepsis, preterm birth, SGA and admission to a neonatal intensive care unit In mothers with iron deficient anemia, increasing severity of maternal anemia has been associated with decreasing neonatal hippocampal volume and production of brain-derived neurotrophic factor.

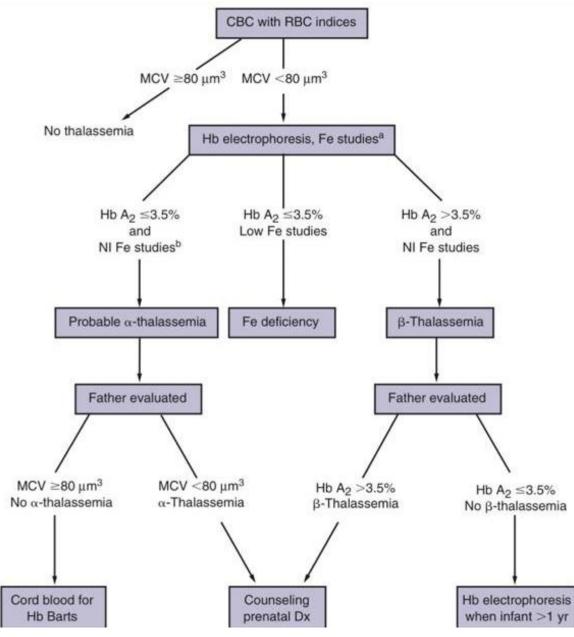






# EVALUATION OF ANEMIA







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MCV is an unreliable marker of iron deficiency in pregnancy. Stimulation of erythropoiesis leads to a physiologic increase in MCV during gestation that counterbalances the microcytosis of iron deficiency.

A low MCV, defined as an MCV ,^ fL, is highly sensitive, but not specific, for iron-deficiency anemia



Serum ferritin levels <'`` ngmL-` indicate established ID. A serum ferritin level<`` ngmL-`, with or without anaemia, indicates insufficient body iron reserves and should prompt treatment. However, as ferritin is an acute-phase reactant, a serum ferritin level within the normal range may not rule out ID in the presence of Inflammation, SLE, Chronic renal disease ,DM





### *Clinical judgment for the decrease and the need for (and aggressiveness of) further evaluation:*

As an example, in an individual with a baseline hemoglobin of  $\mathcal{V}$  g/dL that decreases to  $\mathcal{V}$  g/dL associated with macrocytosis, checking a reticulocyte count and testing for vitamin B $\mathcal{V}$  and folate deficiencies is reasonable. For an individual with a baseline hemoglobin of  $\mathcal{V}$  g/dL that decreases to  $\mathcal{V}$  g/dL without macrocytosis, testing for iron deficiency and vitamin B $\mathcal{V}$  and folate deficiencies is reasonable.



**Evaluate for other causes of anemia if :** 

Extreme microcytosis (eg, mean corpuscular volume [MCV]  $<^{\wedge}$  fL), suggestive of thalassemia

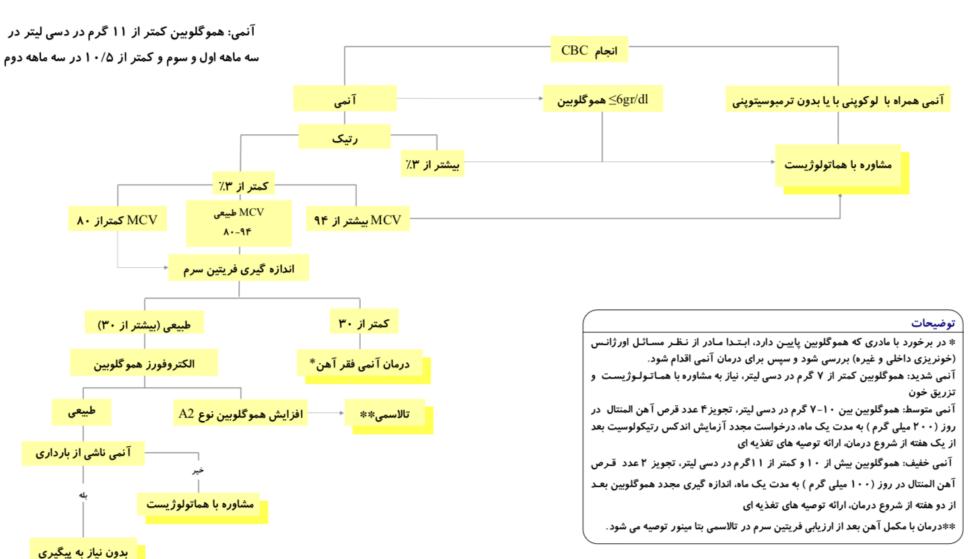
Macrocytosis (MCV >  $\cdot \cdot \cdot$  fL), suggestive of vitamin B $\cdot \cdot$  or folate deficiency or reticulocytosis due to hemolysis

Other cytopenias such as thrombocytopenia or neutropenia Abnormally high white blood cell (WBC) count or platelet count Abnormal RBC or WBC morphologies Failure of the anemia to correct with iron supplementation



أنمى





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ACOG: Screen all pregnant women for anemia at the first prenatal visit with a complete blood count (CBC), along with other appropriate prenatal testing, repeat screening with a CBC at week TY to TA.





All pregnant women should by typed for blood groups (ABO and D) and screened for the presence of red cell **antibodies early** in pregnancy (at booking) and at <sup>Y</sup><sup>A</sup>weeks gestation. In D-negative women, the <sup>YA</sup>-week sample should be taken before the administration of routine antenatal anti-D Ig prophylaxis





The direct Coombs test uses anti-human immunoglobulin to detect immunoglobulins attached to the surface of RBCs.

A positive test indicates an immune cause for a hemolytic anemia





In a woman with microcytic or normocytic anaemia, iron deficiency (ID) should be confirmed by a trial of oral iron.





Anaemic women with a known haemoglobinopathy should have their serum ferritin checked and should only be offered oral iron therapy if their serum ferritin level is <<sup>r</sup> • ngmL



#### Prevention

In areas with a high prevalence of anaemia in pregnancy, daily oral iron  $(\forall \cdot - ? \cdot mg)$  and folic acid  $(* \cdot \mu g)$  supplementation are as part of routine from first antenatal care In areas with a low prevalence of anaemia in pregnancy, non-anaemic women identified to be at an increased risk of ID should have their serum ferritin checked early in pregnancy and be offered oral supplements  $(\forall \cdot - ? \cdot mg/day)$  if serum ferritin is  $< \forall \cdot ngmL$ 



All pregnant women, especially those with antenatal anaemia, have an Hb determination prior to delivery

- Hb concentration be determined in all women after
- significant peri-partum bleeding

#### Prevention

Every effort be made to correct anaemia prior to delivery



Treatment of iron deficiency that is initiated after diagnosis of iron deficiency anemia may be **too late to prevent some adverse** outcomes.

Correction of iron deficiency before the third trimester is ideal, as iron-dependent neurogenesis is maximal during the third trimester and early neonatal life, and iron deficiency during this period has been associated with <u>deficits</u> <u>in neurocognitive development</u>





# Thank you for your attention

