

Obstetrics & Gynecology



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Anemia in Pregnancy

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This sheet contains:

- lecture slides
- Doctors notes
- additional notes and pictures from OBS & GYN books

Introduction

Anemia is common in pregnancy (from 5.4% in developed countries to more than 80% in developing countries)

Anemia in pregnancy has been associated with maternal and fetal adverse outcomes -Hb used to define anemia in pregnancy is lower than in non-pregnant patients An increased requirement of iron in pregnancy so iron deficiency remains the most common cause of anemia in pregnancy and warrants a preemptive approach to prevent a further reduction in Hb

Anemia secondary to sickle cell disease and autoimmune hemolytic anemia merit special attention because there are risks secondary to red blood cell transfusion and risks to withholding transfusion

we should screen for anemia to avoid its complications to the baby & mother

Normal physiological changes in pregnancy

Alterations of hematological parameters particularly in a reduction of hemoglobin (Hb) concentration:

-Total blood volume increases by 50%

Plasma volume rises from 6 weeks gestation progressively increasing to a peak at 32 weeks gestation (due to hormone-mediated vasodilation subsequent activation of the reninangiotensin-aldosterone system)

During pregnancy (Physiological anemia) :

Angiotensinogen rises with estrogen production Increased vasopressin leads to salt and water retention RBC mass increases from the early 2nd trimester around 30–35% a <u>lesser degree</u> compared with plasma volume The increase in RBC mass results from an increased erythropoietin level in response to circulating progesterone and placental lactogen The <u>disproportionate</u> increase in plasma volume to RBC mass leads to decreased Hb and

hematocrit levels

Definition of anemia in pregnancy by WHO

Anemia in pregnancy is defined as Hb:

11>g/dl in the 1st trimester 10.5> in the 2nd & 3rd trimester 10> in the postpartum period

General approach : Preconceptional monitoring of Hb Treatment of low Hb Keep in mind that the most common cause is IDA The goal is To avoid post-partum complications

Choice of contraception : Copper-IUD is not good for IDA due to menorrhagia Sick cell crisis: Progesterone-containing methods are better Types :

Microcytic anemia \rightarrow Iron or Thalassemia Megaloblastic anemia \rightarrow B12 or Folate deficiency Hemoglobinopathy \rightarrow Sickle cell anemia

Complications of anemia in pregnancy





Thalassemia

Adult Hb consists predominantly of HbA composed of α and β globin. Adult Hb also consists of a small proportion of HbA2 ($\alpha 2\delta 2$) and HbF ($\alpha 2\gamma 2$) Thalassemia refers to the reduced production of α and/or β globin chains due to mutations in the α and/or β globin gene Worldwide commonest inherited single-gene disorders Absent or decreased normal α and B globulin chains <u>Autosomal recessive</u> condition -Heterozygous called trait -Homozygous called disease Women with trait status <u>don't need</u> special care Women with HbH may have a successful pregnancy Close medical evaluation and follow-up Increased risk of neural tube defect due to folic acid deficiency **The laboratory parameters suggestive of thalassemia consist of :** Microcytic, hypochromic anemia.

Management and counseling:

All women should be offered hemoglobinopathy screening To : Identify early those at risk of the affected baby Offer prenatal diagnosis To prevent infant morbidity and mortality Counsel : Maternal and fetal risk If the fetus is affected involve pediatricians Folic acid 5 mg before and during pregnancy No specific intrapartum or postpartum care

Beta thalassemia:

Effects of pregnancy on B thalassemia:

Trait \rightarrow Mild anemia Major \rightarrow Risk of blood transfusion increases Maternal complications of iron overload: \rightarrow If short stature with pelvic bone deformity CPD \rightarrow risk of CS increasing

Effects of B thalassemia on pregnancy:

Beta Trait has a normal outcome Beta Major Fetal hypoxia due to maternal anemia IUGR Preterm birth

Alpha Thalassemia:

Alpha major is incompatible with life **HbH disease:** Mild to moderate hemolytic anemia Adult: worsened in pregnancy Hepatosplenomegaly

Effects of pregnancy on alpha thalassemia:

Alpha trait has Normal outcome Alpha major Maternal risks: Gestational hypertension (50%) Preeclampsia (30%) Placental abruption Obstructed labor (large baby) DIC APH PPH

Effects of alpha thalassemia on pregnancy:

Alpha trait has normal outcome **Alpha major:** Incompatible with life baby Severe anemia Hydrops fetalis Abnormal organogenesis Polyhydramnios Placentomegaly Stillbirth

Iron deficiency anemia (IDA)

The most common type of anemia during pregnancy (75% of cases).

Is caused by blood loss, insufficient dietary intake, or poor absorption of iron from food Diagnosis: if microcytic do an iron study Ferritin level has the greatest sensitivity and specificity Signs & symptoms : -Irritability -Angina (chest pain) -Palpitations (feeling that the heart is skipping beats or fluttering) -Breathlessness -Tingling, numbness, or burning sensations -Glossitis (inflammation or infection of the tongue) -Angular cheilitis (inflammatory lesions at the mouth's corners) -Koilonychias (spoon-shaped nails) or brittle nails -Poor appetite -Dysphagia (difficulty swallowing) due to the formation of esophageal webs (Plummer (Vinson syndrome -Restless leg syndrome

Parameters:

- \downarrow Ferritin Hemoglobin MCV MCH
- **†** Total iron-binding capacity Transferrin RDW

IRON

OB Triad

Iron Deficiency Anemia

- Hemoglobin < 10 g
- MCV <80 μm^3
- RDW >15%
- Iron requirements ↑ in pregnancy to about 1000 mg/day.
- Most of the iron is used for hematopoiesis, especially in the last half of pregnancy.
- The amount of iron from the diet is insufficient to meet the needs of the pregnancy, so many patients will need to take supplemental iron. The most common side effect of this supplementation is constipation.

Management:

RCOG guidelines recommendations :

Iron supplement for all women after 12 weeks if there are no contraindications Daily elemental iron 30 mg prophylaxis 60–120mg treatment

Dosage forms :

Ferrous sulfate 325 mg (65 mg iron)

Ferrous gluconate 300 mg (34 mg iron)

Ferrous fumarate is a large compound compared to ferrous sulfate 300 mg (98.6 mg iron) Foods rich in ascorbic acid (vitamin C) enhance iron absorption

If the patient is taking a Calcium supplement, Iron should not be taken at the same time should be 12 hours apart ;(Calcium in the morning & Iron in the evening and so on)

Indications for IV iron therapy: -Can't take Iron by mouth Can't absorb Iron adequately through the gut- have inflammatory bowel disease or other intestinal illnesses that are aggravated by oral iron supplements -Can't absorb enough iron due to blood loss -Need to increase iron levels fast to avoid medical complications or a blood transfusion All types of treatment can increase Hb by 0.8 g/dl/week IV iron has minimal side effects but should be monitored for : Gastrointestinal pain such as nausea and cramping Difficulty breathing Skin irritations/rash Chest pain Low blood pressure Anaphylaxis can include difficulty breathing, itching, and rash

Precautions before giving IV iron :

Prepare IV Hydrocortisone + Anti-histamine Dilute Iron in 250-500 ml of N/S given over 2 hours IM route should be avoided as it leads to : Pain Abscess formation

Megaloblastic anemia

Impaired DNA synthesis \rightarrow Ineffective erythropoiesis. Folic acid deficiency 2nd most common during pregnancy Less common B12 deficiency which is difficult to detect (folic acid supplements masking B12 (deficiency Slowly progressive Tend to occur mostly in 3rd trimester **Usual symptoms :** Weight loss Anorexia Glossitis Bleeding due to thrombocytopenia Leads to poor outcomes : Placenta abruption Preeclampsia IUGR PTL Folic acid deficiency may lead to open neural tube defects Labs: Macrocytic normochromic anemia -Peripheral blood smear hypersegmented neutrophils & oval macrocytes and Howell Jolly bodies Erythrocyte folate level is the best indicator of the serum level

Management :

Folate deficiency: treated with folic acid 1 mg/day within 10 days WBC and Platelet normalize, Hb increases after several weeks B12 deficiency: IM cobalamin 1 mg monthly or sublingual

OB Triad

Folate Deficiency Anemia

- Hemoglobin < 10 g
- MCV > 100 μm^3
- RDW >15%

Sickle cell anemia

Autosomal recessive. Sickle-shaped RBCs Common in African Americans (8%), Middle East & India Hb S: Sickle cell anemia Hb SC: Sickle cell hemoglobin C Hb S -Thal: Sickle cell -Thalassemia Hb SS: Homozygosis is the most common phenotype (Saudi Arabia), Less soluble, and tends to polymerize and distort RBCs

Risk of sickling increased during pregnancy (metabolic requirements) Risk of vascular stasis + Hypercoagulable status

Pathogenesis :

Hemolytic anemia \rightarrow Extravascular hemolysis \rightarrow Chronic anemia \rightarrow Microvascular obstruction \rightarrow ischemia \rightarrow Infraction(Vaso-occlusive crisis)

Points:

Pregnant with sickle cell trait have twice the frequency of UTIs SCA patients should be screened for UTIs each trimester Blood pressure checked every visit One in four children will be affected if parents have SC trait Clear care for those women

Complications :

Spontaneous miscarriage IUGR IUFD SGA Preeclampsia Preterm labor UTI more than 2 time

Labs:

Normocytic normochromic anemia Reticulocyte count (3-15 %) ↑ LDH ↑ Haptoglobin ↓ Peripheral blood: sickle cell, target cell, Howell-Jolly bodies <u>Screening and diagnosis by Hb electrophoresis :</u> Hb S (85-100%) Absent Hb A Normal Hb A2 Hb F elevated more than 15%

Management (OSCE):

Hydroxyurea is not recommended in pregnancy (should be stopped 3 months before (pregnancy BP & Urinalysis (UTI & Proteinuria) for each visit Infections should be treated with antibiotics Severe anemia needs blood transfusion (in more severe plasma exchange) Pain crises are managed with O2, hydration (vomiting and nausea are common), analgesia Should receive pneumococcal vaccine before Pregnancy Folate supplements 4 mg/day Low-dose Aspirin as prophylaxis for PET Give Iron supplements only by indication Fetal well-being twice weekly since 32 weeks Low-molecular-weight heparin any antenatal hospital period if there are no contraindications Fetal growth weekly in 3rd trimester Avoid dehydration and stress during the intrapartum period Avoid pethidine as it increases the risk of seizures Early ambulation after delivery or wearing stockings to prevent thromboembolism **Contraception** : **Excellent options are Mirena and Progesterone only pills COC** are contraindicated Medroxyprogesterone acetate decreases pain crises Blood transfusion in sickling patients: May precipitate a crisis if a sudden increase in HCT Hb 6–8 g/dl is typical for HbSS Consider transfusion if : Severe anemia Multiple pregnancy Preeclampsia Acute chest syndrome Acute renal failure Target level < 30 % of sickle cells in circulation : Partial exchange transfusion Time and mode of delivery SCD with normal growing fetus : Prepare cross-matched blood before delivery the hematologist should be consulted Induction of labor or CS (by indication as SCD is not an indication for CS) at 38 weeks Epidural analgesia is preferred Continuous intrapartum fetal monitoring

Key points

Offer screening for anemia at booking and 28 weeks as this allows time for treatment. Hb < 11 or 10.5 in 2nd trimester need investigation

Anemia increases the risk for Preterm labor

The parenteral iron should only be considered for intolerant women

At term, IDA should be treated with blood transfusion

