Renal & thyroid disease in pregnancy

Objectives

a-Know Physiological changes of these two organs during pregnancy .

b- Discuss types of their disorders during pregnancy , their C/F , diagnosis & management .

C-Discuss fetal & maternal complications of these disorders .

Physiological changes in pregnancy

Ureters and renal calyces dilatation (remembered in U/S).

renal plasma flow + glomerular filtraon urinary protein excretion and creanine dearance. So: -

The upper limit of serum creanine dearance fails 65 µmol/L.

The upper limit for proteinuria throughout pregnancy is 300mg/24 hours.

Urinary tract infection

It is more common in pregnancy due to physiological dilatation of the upper renal tract.

Asymptomac bacteriuria 4-7%, 40% of them will develop symptomatic UTI.

Cyss: 1% of pregnancies

Pyelonephris: 1 to 2% of pregnancies

Predisposing factors:

- previous history of UTI.

- Diabetes millets, polycystic kidneys, urinary tract calculi, renal tract abnormalities (duplex kidney or ureter)

- Neuropathic bladder(spina bifida or multiple sclerosis).

- Drugs: steroids or immunosuppression.

Presentation

Asymptomatic: Asymptomatic bacteriuria + patients with predisposing factors: midstream urine specimens (antenatal screening).

Clinical features include:

Cystitis: urinary frequency, dysuria, haematuria, protienuria and - suprapubic pain.

Pyelonephritis: fever, loin pain and/or abdominal pain, vomiting and rigors.

Diagnosis

Dipstick for proteinuria.

MSU for analysis. Bacteriuria: 100000 organisms/ml of urine or <a>o more

MSU for culture and sensitivity. It should be repeated if it is non- significant or with mixed growth.

<u>management</u>

<u>Asymptomatic bacteriuria</u>: a 3-day course of antibiotics (oral) to environment pyelonephritis + preterm labour.

<u>Acute cystitis</u>: a 7-day course of antibiotics (oral). 🍩

- Urine culture following treatment to ensure eradication of organisms. Recurrent bacteriuria occurs in 15% of women in pregnancy and requires a second course of antibiotics. - U/S: in paents with 2 or nor e UTIs (+ve culture).

<u>Pyelonephritis:</u>

- antibiotics for 10-14 days.
- IV antibiotics for patients with vomiting or pyrexia.
- IV fluids may be required.
- renal function should be checked.

- U/S to exclude hydronephrosis, renal calculi and congenital abnormalities (*risk factors*).

prophylactic antibiotics: two or more UTIs (positive culture) i.e. recurrent UTI or one of the above risk factors



Renal impairment

Aetiology:

- 1. reflux nephropathy
- 2. diabetes

3. systemic lupus erythromatosus (SLE)

4.Glomerulonephris.

5. polycysc kichey disease.

<u>Classification</u>: mild, moderate or severe depending on the serum creatinine.

creatinine depends on the muscle mass i.e. a figure representing moderate impairment in an 85-kg may represent severe impairment for a 50-kg woman

Presentation:

hypertension and protienuria ± haematuria in early pregnancy. Blood tests for urea and creatinine must be done.

Effect of pregnancy on renal impairment:

- mild impairment (creani $re < 125 \mu mol/l$): tolerate pregnancy well with no renal function deterioration.

- severe renal impairment (creani $re > 250 \mu mol/l$): at increased risk of permanent loss of function during and after pregnancy and even end stage of renal failure.

Effect of renal impairment on pregnancy :

1. PE, IUGR, spontaneous and iatrogenic premature delivery.

- severe renal impairment + hypertension have < 50 % chance of successful pregnancy because of severe, early-onset of PE with severe IUGR.

- premature delivery is justified in rapidly worsening renal function to avoid dialysis even in the absence of PE.

2. severe renal impairment polyhydramnios and risk of cord prolapse due to fetal polyuria in response to high osmotic load from increased maternal urea.

3. nephroc syndrome and heavy protienuria severe hypoalbuminria with associated risks of pulmonary oedema and thrombosis.

management of renal impairment

prepregnancy counseling and multidisciplinary care.

Documenting baseline values (prepregnancy & early pregnancy) • for creatinine, uric acid, albumin and protein.

Tight control of even mild hypertension with antihypertensive • agents (the choice is no different in women with renal disease).

discontinue angiotensin-converting enzyme (ACE) inhibitors prior • to pregnancy or once pregnancy is confirmed.

Discontinue: diuretics unless there is severe hypoalbuminaemia • and insipient pulmonary oedema.

Admission: in worsening hypertension, increasing creatinine, and • large increase in proteinuria because of high risk of PE with difficult diagnosis in the present of BP + proteinuria.--

Diagnosis of PE is supported by: IUGR, thrombocytopenia and abnormal liver function.

Prophylactic low-dose(75 mg/day) aspirin to decrease the risk of • PE.

Serial scans for fetal growth and liquor volume. •

Serial haematology and biochemistry. •

Post partum: continue close monitoring. ACE inhibitors are safely used in breastfeeding

Renal transplants

Pregnancy outcome in well functioning renal transplants is similar to the general population.

Pregnancy should be delayed for 1-2 years to allow graft function to stabilize and immunosuppression to reach maintenance levels.

Risks in pregnancy: is related to pre-pregnancy renal function and to the presence of hypertension.

Women are immunosuppressed and prone to infection.

Immunosuppressive drugs used in pregnancy: prednisolone, azathioprine, & cyclosporine.

Dialysis

pregnancy on dialysis is unusual: end-stage renal failure reduces fertility.

Patients on dialysis should be advised not to get pregnant.

Common risks: anaemia and haemorrhage.

Increased risks of:

miscarriage, fetal death, pre-eclampsia, pre-term labour, PROM, polyhydramnios and placental abruption.

Pregnant women require increasing dialysis to maintain the predialysis urea < 15-20 mmol/l.

Poor obstetric outcome is similar with both haemodialysis and peritoneal dialysis.

Thyroid Disease

- Physiological changes :-

1- slight increase in size of T.G.

2- Thyroid binding globulin double by liver due to estrogen stimulation so increase total T4 &T3.

3- no significant change in free T3,T4 level.

4 - renal clearance of iodine increased due to increased GFR & increased consumption of fetus lead to decreased plasma inorganic iodine .

Maternal Hyperthyroidism

- incidence: 2: 1000 of pregnancies.

- **causes**: Graves disease (90%) autoimmune disease with presence of circulating TSAB, toxic nodule, Hashimotos thyroiditis, multiple nodular goiter, trophoplastic disease (rare).

- Diagnosis :-

1- In early pregnancy , the diagnosis c/o ly difficult

2- in uncontrolled hyperth.: maternal arrhythmias, vomiting, abd. Pain, diarrhea.

- Fetal complications : IUGR , stillbirth, fetal tachy cardia, preterm labor , abortion, congenital abn., fetal thyrotoxicosis.

-Treatment :-

1- drug therapy ; the aim is to maintain maternal T3,T4 level in high normal range ex.: carbimazole ,propyl thiouracil, B-blockers.

2- Radioactive iodine.

Maternal hypothyroidism

- Incidence :9:1000 of pregnancies.
- Causes : iodine deficiency , over treated hyperthyroidism.

- Diagnosis :-

A- C/F : easy fatigue ability, cold intolerance, slow speech, oligomenorrhea, infertility.

B- investigation: decreased free T3&T4, increased TSH.

- **Fetal complication** : congenital hypothyroidism & cretinism of newborn if the cause is maternal iodine deficiency, abortion, preterm labor, still birth.

- **Treatment** : restore TSH to normal range by thyroxine replacement therapy.

REFERENCE

OBSTETRIC BY TEN TEACHERS

DEWHURST, S TEXT BOOK OF OBSTETRIC & GYNECOLOGY