Infections in Pregnancy and puerperium **Clinical aspects**

Dr. Nadisha Badanasinghe Senior Lecturer Dept. of Medical Microbiology

Objectives

- Why infections are problematic in pregnancy?
- What are the common infections which pose problems in pregnancy and pueperium?
- How do you investigate and manage these infections?

What are the problems with pregnancy?

- Low immune status in mother
 - · To prevent rejection of semi-allogeneic foetus
 - Adaptive immune responses are weakened

 - Boosted innate response
 Some infections are more severe in pregnancy (ex. Malaria, influenza, etc)
 - Some infections are reactivated in pregnancy (HSV, CMV)
- Foetus has poorly developed immune system
 - IgM andNo IgG IgM and IgA - Very low

 - Low CMI

Problematic infections

- · Infections which cause congenital infections
- Varicella
- Rubella
 CMV
- Syphilis
 Toxoplamsma
- UTI
- PROM/ intra-amniotic infections

- STI (Hep B, HIV, Chlamydia, Candida etc)
- · Infections which can be severe in pregnancy
 - Chickenpox
 Influenza
 - Malaria
 - Hep E
- - Endometritis/ puerperal sepsis
 Mastitis/ Breast abscess
 Episiotomy infections
 Septic abortion

Case I

- POA 20 weeks, uneventful
- No significant past history
- Urinary culture report
- Coliform> 10⁵ C.F.U
- No symptoms

UTI

- The most common medical complications during pregnancy
- asymptomatic bacteriuria 2-10 %
- acute cystitis- I-4 %
- Pyelonephritis 0.5-2%
- · Risk begins in week 6 and peaks during weeks 22 to 24
- Asymptomatic bacteriuria \rightarrow 20-40% develop symptomatic pyelonephritis later if untreated
 - Also lead to premature delivery/IUGR/ LBW





UTI in Pregnancy - Pathogenesis

- Relaxation of muscles in ureters and bladder → dilatation → stasis and reflux
- · Dilatation of renal pelvis
- physiologic increase in plasma volume during pregnancy decreases urine concentration
- Up to 70 % of pregnant women develop glycosuria, which encourages bacterial growth in the urine

UTI pathogens

- Similar to a non pregnant female
- E. coli 80-90%
- Other coliforms
- GPC
- S. saprophyticus, enterococci, Grp. B Streptococci

Management

- Screening urine culture in first visit
- Repeat culture in T3
- Treat with a sensitive antibiotic
 - Cephalexin, Nitrofurantoin, Co-amoxiclav
 - 5 days for asymptomatic bacteriuria
 - 5-7 days for cystitis
 - Pyelonephritis Admit and IV (Amp/ Gent/ Cefotaxime) x at least 3ds \Rightarrow oral 10 days
- Repeat UC 48 hours after stopping AB

Case 2

- POA 32 weeks
- Dribbling for ~ 24 hrs
- Back pain
- Fever

PPROM

- ROP before 37 weeks of gestation without labour contractions
- Prolonged ROM ROM for greater than 24 hours
- · Associated with an increased risk of ascending infection
 - \rightarrow infection complications in foetus
 - \rightarrow infection complications in mother

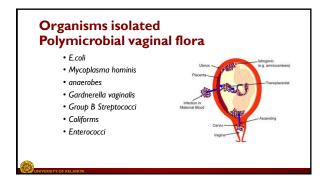
Intra-amniotic Infection (Chorioamnionitis)

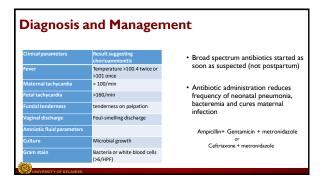
- Infection of the membranes and chorion of the placenta
- Typically due to ascending polymicrobial bacterial infection in the setting of membrane rupture

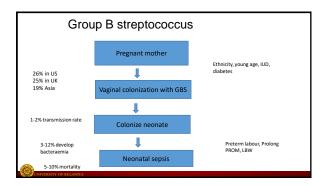
 - Rare cases after diagnostic amniocentesis
- Risk factors: PROM, young age, nullparity and bacterial vaginosis



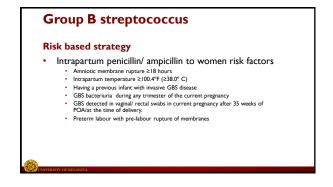
Presentation Maternal Foetal Fever Tachycardia Uterine tenderness Foul smelling or grossly purulent amniotic fluid Foetal Foetal Heart rate abnormalities Preterm labour Arrest of progress of labour Neonatal sepsis Neonatal death/ stillbirth







Group B streptococcus Important pathogen in early and late onset neonatal sepsis Management Screening-based strategy Universal screening vaginal and rectal swab specimens at 35–37 weeks POA Screening in high risk mothers



Case 3

- POA 12 weeks
- Exposed to chicken pox ~ I day
- Uncertain past history of chicken pox

Chicken pox

- ~ 10% of women in childbearing age are susceptible
- Transplacental infection in T1 < 3%
- Can affect the foetus if it occurs in early (foetal varicella syndrome) or very late pregnancy (neonatal varicella)

Chicken pox in pregnancy

- Effects in Foetus
 - No damage
 - Still birth
 - Foetal Varicella Syndrome (2-3% if within 20 weeks of POA) skin scarring, limb hypoplasia, visceral, neurological and eye lesions
- · Effects in mother
 - Severe chicken pox

Chicken pox significant exposure

- Check IgG Abs
 - Seropositive No action required
 - Seronegative/ Serology not available

 - Give VZIG as soon as possible preferably within 48 hours of contact (maximum, 96 hours) Advice to seek medical attention immediately if chickenpox develops
 - VZIG is indicated after significant exposure to VZV upto 20 weeks of POA

Case 4

- POA 20 weeks
- Fever, chickenpox like rash ~ 2 days
- No history of childhood chickenpox

Management of Chickenpox in pregnancy

- Medical review is essential
- <24 hrs since onset of rash → oral aciclovir
- > 24 hrs since onset of rash > No aciclovir
- No benefit from VZIG once the rash develops
- If low risk of complications \rightarrow manage at home
- High risk group → monitor in hospital
- Isolate from other pregnant mothers, babies and nonimmuned staff
- If complications IV aciclovir



Chickenpox just before delivery

- Rash develops 6 days or less before delivery or upto 7 days after delivery
 - · Neonates should receive prophylactic VZIG soon after delivery
 - Virus cross placenta

 - No protective Abs to foetus

 Baby develops severe disseminated varicella with pneumonia and encephalitis
- Rash develops 7 days before delivery

 - Abs cross placenta
 Foetus cope remarkable well
 - Baby does not develop severe disease

STI

- I. Syphilis
 - pregnant women with untreated early syphilis,
 - 70% chance of transmitting the infection to her fetus
 - 25% of pregnancies result in stillbirth
 14% in neonatal death

 - overall perinatal mortality of about 40%

 - Treatment with single IM benzathine penicillin for primary, secondary, and early latent syphilis in pregnancy, and three doses one week apart for late

STI

- - Transmitted to foetus/ neonate through placenta, birth canal, breastfeeding
 - Increased risk with high viral load, young maternal age, low CD4 count, Prolonged PROM, vaginal delivery, other STIs

 Maternal anti retroviral therapy significantly reduce the mother-child transmission

 - zidovudine, nevirapine, or a combination of the two, are recommended during pregnancy to reduce mother-to-child transmission of ${\sf HIV}$
 - Other interventions
 - Screening
 - Counseling
 EL-LSCS before onset off labour
 - Neonatal post exposure prophylaxis with ARV

STI

- 3. HBV infection
- Transplacental and perinatal transmission
 - Acute HBV infection during pregnancy usually is mild and not associated with teratogenicity or mortality
- Unless the patient has acute liver failure, antiviral therapy is usually unnecessary Treatment is mainly supportive, with monitoring of liver biochemical tests
- To reduce transmission to baby:
 Screening and monitoring of viral load
- All babies born to positive mothers should receive Hep B Immunoglobulin and Hep B vaccine within 12 hours of delivery

STI

- 4. Gonorrhoea/ Chlamydia infection
 - Pre-term labour, PPROM, Chorioamnionitis, Endometritis
 - opthalmia neonatorum (40-50%), Pneumonia (18%)
 - Most are asymptomatic
 - · Screening in high risk population and treatment

Other severe infections

- I. Influenza
 - Pregnant women are at high risk for severe complications of influenza during seasonal influenza periods and pandemics
 - some studies suggest an increased risk for adverse outcomes among infants born to mothers infected with influenza during pregnancy

 - influenza vaccine for all pregnant women in any trimester during flu season.
 - Pregnant women with symptoms of influenza should be tested and treated immediately with oseltamivir



Other Infections

- 3. Zika virus
 - · Febrile illness ~ dengue Transmitted by Aedes mosquitoes
 - Zika virus is a cause of microcephaly and other congenital anomalies
 - Risk higher with infection in TI
 - Outbreak in Brazil started in 2015 → spread to Americas, Caribbean, Pacific, and SE Asia
 - Diagnosis Pregnant mothers with recent travel + symptoms
 - Serum for PCR (within 2 weeks of travel)
 4 weekly USS

 - · Treatment supportive

Case 5

- EM-LSCS performed in a 36 weeker with prolonged ROM, and lack of progress.
- Developed fever following LSCS with high WBC and CRP

Post-partum Endometritis

- Postpartum infection of the uterus
- Most common cause of puerperal fever \Rightarrow puerperal sepsis
- Predominant predictor: Caesearan section particularly after labour or premature rupture of membranes
 - Rates vaginal delivery 0.9-3.9%
- Caesearan section rate: 10-50%
- Secondary risk factor BV
- Presents with
- Fever on 1st or 2nd day postpartum Lower abdominal pain
- Uterine tenderness
- Leucocytosis

PP Endometritis

- It is a polymicrobial Infection
- **Pathogens**
- Group B Streptococci
 - G.vaginalis
- Enterococci
- E.coli, Bacteroides spp
- · Diagnosis: Blood culture, Culture of high vaginal swab
- Treatment: intravenous antibiotics

(Ampicillin/ Co-amoxiclav/ Ceftriaxone + gentamycin + metronidazole)

Case 6

- 2 weeks post delivery
- Breast feeding
- Pain and swelling in L/ breast
- Localized inflammed area

Mastitis/ Breast abscess

- Present with Inflammation+/- infection +/- abscess (3%)
- Management
 - Physiological correct breastfeeding, expressing, massage, apply warmth
 - Treatment analgesics
 - Antibiotics If symptoms are not resolving within 12 to 24 hours with physiological methods or if presenting symptoms are moderate or severe
 - Most common organisms- S. aureus (less commonly Streptococcus or Escherichia coli)
 - Oral flucloxacillin/ cephalexin x 5 days
 - If not responding to first line AB/ severe mastitis hand expressed mid stream breast milk culture and ABST



Episiotomy infections

- Presents with significant pain and a delay in healing of episiotomy, oedema, redness, ecchymosis, and wound discharge, dehiscence, +/- fever, +/- abscess, +/- myonecrosis, +/- sepsis
- $\label{lem:causative agents} \textbf{Causative agents} \textbf{Streptococcus pyogenes}, \textbf{Staphylococcus aureus (MRSA)}, \\ \textbf{Clostridium perfringens}$
- $\label{thm:continuity} \mbox{Treatment surgical debridement + antibiotics (depends on superficial/ deep infection)}$

Septic abortion

- Usually result from induced abortions done by untrained practitioners using nonsterile techniques \Rightarrow high mortality
- Infection of the placenta and fetus, or products of conception, of a pre-viable pregnancy
- Polymicrobial vaginal flora, bacteria from contaminated equipment, toxin
- Diagnosis Blood, evacuation products/ curettings, high vaginal swab for culture
- $\label{eq:Management-Broad spectrum IV antibiotics to cover all possible organisms + prompt evacuation of retained products and devitalized tissue$

Summary

- Infections in pregnancy
- Risk to baby
- Risk to mother
- · Clinical suspicion
- Adequate investigations/ screening
- Adequate treatment
 - Antibiotics/ antivirals should be safe to use in pregnancy

