# ANTENATAL FETAL ASSESSMENT



#### **OBJECTIVES:**

#### Describe how to test for each of the following

- Fetal well-being -
- Fetal growth -
- Fetal movement -
- Amniotic fluid -
- Fetal lung maturity

#### FETAL ASSESSMENT (FETAL WELL-BEING)

- Fetal assessment is to identify fetuses at risk of neurologic injury or death in order to prevent it
- To prevent prenatal mortality & morbidity

### FETAL AND NEONATAL COMPLICATIONS OF ANTEPARTUM ASPHYXIA

- Stillbirth (Mortality)
- Metabolic acidosis at birth
- Hypoxic renal damage
- Necrotizing enterocolitis
- Intracranial hemorrhage
- x Seizures
- Cerebral palsy

#### RATIONAL

- If fetal oxygenation challenged:
- blood flow directed to brain, heart & adrenal & blood flow away from the kidney → decrease fetal urine production → decrease AF volume.
- CNS hypoxia → Fetal movement decrease
- -chemoreceptor's → vegally-mediated reflex →

Fetal heart rate abnormality late deceleration.

## CONDITIONS ASSOCIATED WITH INCREASED PERINATAL MORBIDITY/MORTALITY

Small for gestational age fetus

Decreased fetal movement

Postdates pregnancy (>294 days)

Pre-eclampsia/chronic hypertension

Pre-pregnancy diabetes

Insulin requiring gestational diabetes

Preterm premature rupture of membranes

Chronic (stable) abruption

## WHEN TO START FETAL ASSESSMENT ANTENATALLY

- \*\* Risk assessed individually
- \*\*For D.M. fetal assessment should start from 32 weeks onward if uncomplicated
- \*\*\*If complicated D.M. start at 24 weeks onward
- \*\*For Post date pregnancy start at 40 weeks
- \*\*For any patient with decrease fetal movement start immediately
- \*\* Fetal assessment is done once or twice weekly

#### EARLY PREGNANCY ASSESSMENT

#### Fetal heart activity

\* fetal auscultation (special stethoscope or Doppler)

~12weeks



Nuchal translucency measurement for early screening for chromosomal abnormality Between 11-13+ weeks

#### \* fetal heart activity seen by USS

#### Can be seen from 6weeks



#### EARLY PREGNANCY ASSESSMENT

#### Fetal movement

- Fetal movement are usually first perceptible to mother ~17w-20w (quickening)
- 50% of isolated limb movements are perceived
- 80% of trunk and limb movements

#### Fetal growth

- Fundal height
- × USS

### FETAL GROWTH

- \* By fundal height measurement in the clinic
- \* By ultrasound

#### **Biometry**:

Biparietal diameter (BPD)

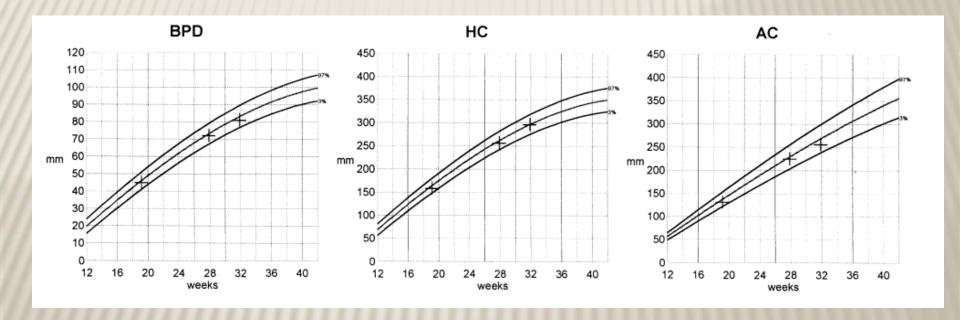
Abdominal Circumference (AC)

Femur Length (FL)

Head Circumference (HC)

**Amniotic fluid** 

#### **GROWTH CHART**



## ASSESSMENT OF FETAL GROWTH BY ULTRASOUND

**BPD** 





AC



FL



#### LATE PREGNANCY ASSESSMENT

- \* Fetal movement counting kick chart
- **×** Contraction stress test CST
- Non stress test NST
- Doppler Velocimetry UAV
- \* amniotic fluid index AFI

#### Indications for antepartum fetal surveillance

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Antiphospholipid syndrome

Poorly controlled hyperthyroidism

Hemoglobinopathies

Cyanotic heart disease

Systemic lupus erythematosis

Chronic renal disease

Type 1 diabetes mellitus

Hypertensive disorders

#### Pregnancy complications

Preeclampsia

Decreased fetal movement

Oligohydramnios

Polyhydramnios

Intrauterine growth restriction

Postterm pregnancy

Isoimmunization

Previous unexplained fetal demise

Multiple gestation

Adapted from data in American College of Obstetricians and Gynecologists. Antepartum fetal surveillance. Practice Bulletin #9, October 1999.



### FETAL MOVEMENT COUNTING

- It should be started ~28w in normal pregnancy
- &~24w in high risk pregnancy
- It can reduce avoidable stillbirth

#### **CARDIFF TECHNIQUE**

- -10 movement in 12 hours
- -If abnormal patient should get further assessment SADOVSKY TECHNIQUE
- -4 movement / hour if not felt another hour If not patient need more assessment

#### **CONTRACTION STRESS TEST (CST)**

- Causing uterine contraction over 20minutes
- \* At least 2 uterine contractions
- Uterine contraction restrict 02 delivery to the fetus
- Normal fetus will tolerate contraction
- \* Hypoxic fetus will have late deceleration
- High false positive rate ~50%
- x 100% true negative rate

## NON STRESS TEST (NST)

- Main advantage over CST is no need for contraction
- ★ False +ve & false -ve higher than CST
- \* done







## **NON STRESS TEST**

- The base line 120-160 beats/minute
- ★ Different criteria in fetuses <32w
  </p>

#### Reactive:

At least two accelerations from base line of 15 bpm for at least 15 sec within 20 minutes

#### Non reactive:

No acceleration after 20 minutes- proceed for another 20 minutes

## NON STRESS TEST (NST)

If non reactive in 40 minutes—proceed for contraction stress test or biophysical profile

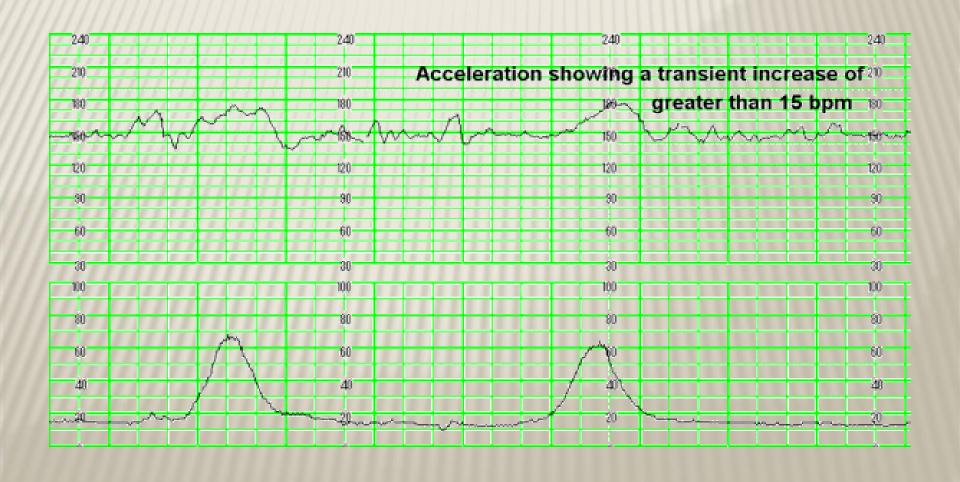
The positive predictive value of NST to predict fetal acidosis at birth is 55%

#### **INTERPRETATION OF CTG**

#### Normal Baseline FHR 110-160 bpm

- Moderate bradycardia 100–109 bpm
- Moderate tachycardia 161–180 bpm
- Abnormal bradycardia < 100 bpm</li>
- Abnormal tachycardia > 180 bpm

## **ACCELERATION**



#### **DECELERATION**

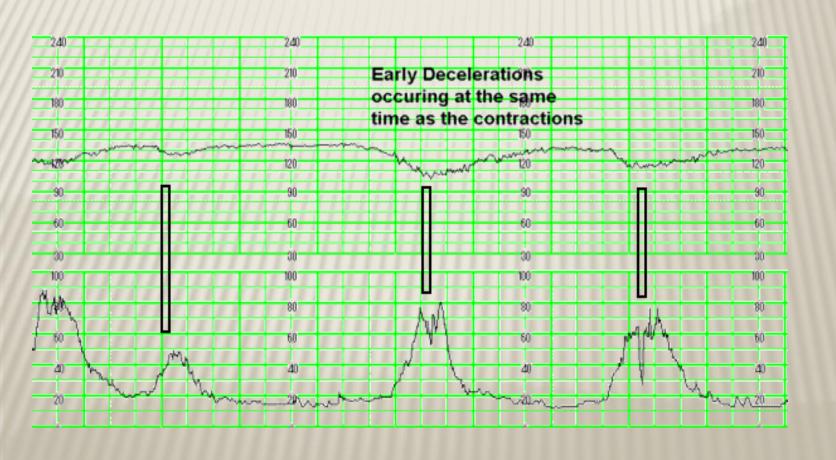
EARLY : Head compression

LATE: U-P Insufficiency

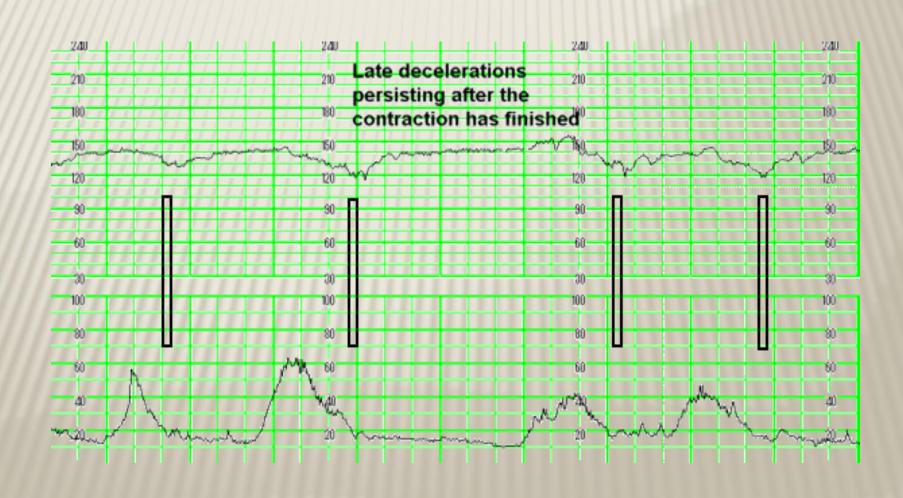
VARIABLE: Cord compression

Primary CNS dysfunction

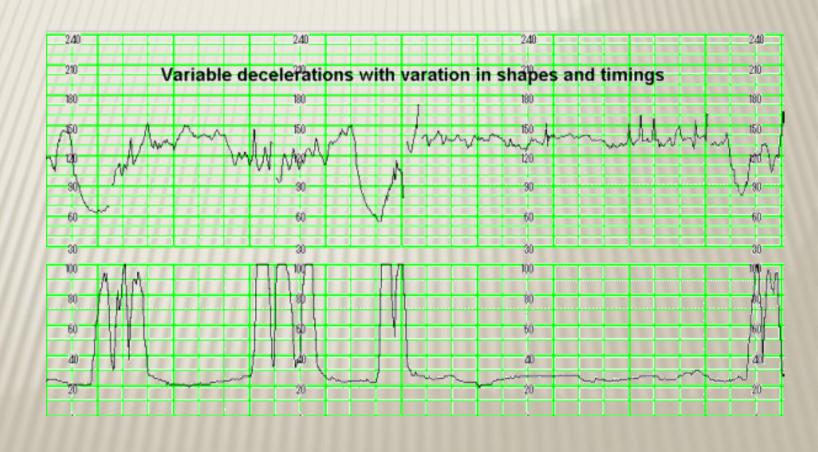
#### EARLY DECELERATION



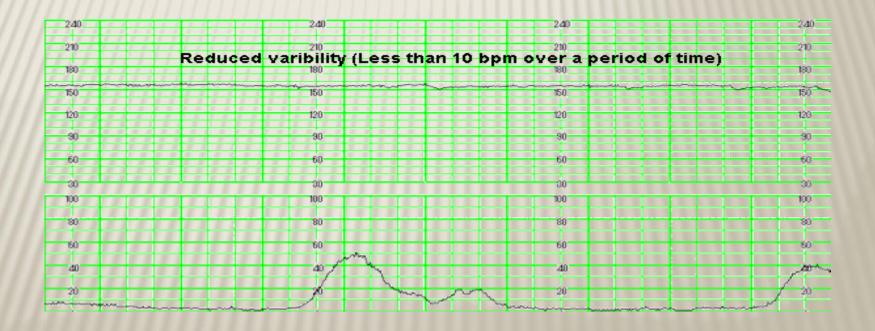
## LATE DECELERATION



## VARIABLE DECELERATION

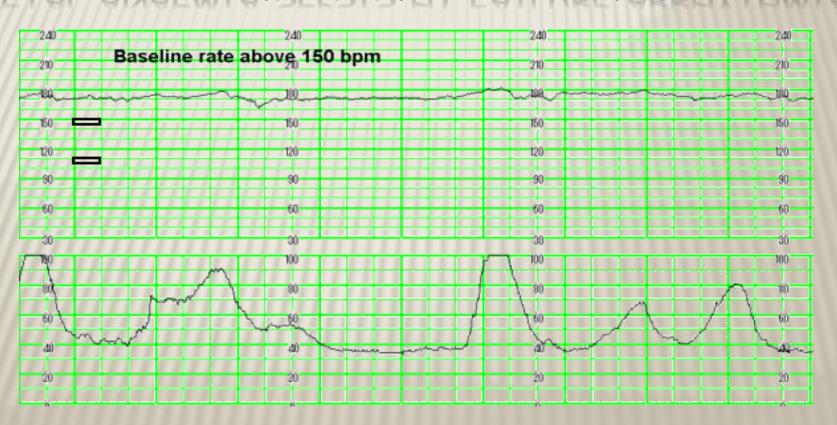


## REDUCED VARIABILITY



#### TACHYCARDIA HYPOXIA

## CHORIOAMNIONITIS MATERNAL FEVER B-MIMETIC DRUGS FETAL ANAEMIA, SEPSIS, HT FAILURE, ARRHYTHMIAS

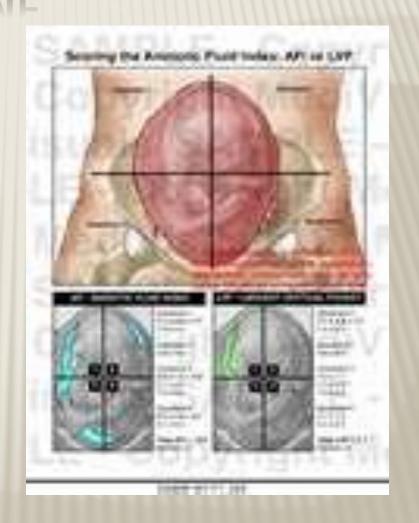


#### AMNIOTIC FLUID VOLUME ~AFI

- \* Amniotic fluid index AFI
- -the sum of the maximum vertical fluid pocket diameter in four quarters
- -the normal value 5-25cm
- -<5~ oligohydraminous
- ->24cm polyhydraminous

### AMNIOTIC FLUID VOLUME





#### **BIOPHYSICAL PROFILE (BPP)**

- Combines NST with USS estimation AFV, fetal breathing, body movement & reflex/tone/extension-flexion movement.
- it is a scoring system
- \* it is done over 30minute
- It measure acute hypoxia(NST, body mov. &breathing) & chronic hypoxia (AFI)

## FETAL BIOPHYSICAL PROFILE/NST+

Biophysical	Normal (score=2)	Abnormal (score=	
Variable		0)	
Fetal breathing movements	1 episode FBM of at least 30 s duration in 30 min	Absent FBM or no episode >30 s in 30 min	
Fetal movements	3 discrete body/limb movements in 30 min	2 or fewer body/limb movements in 30 min	
Fetal tone	1 episode of active extension with return to flexion of fetal limb(s) or trunk. Opening and closing of the hand considered normal tone	Either slow extension with return to partial flexion or movement of limb in full extension Absent fetal movement	
Amniotic fluid volume	1 pocket of AF that measures at least 2 cm in 2 perpendicular planes	Either no AF pockets or a pocket<2 cm in 2 perpendicular planes	

## BPP

- The risk of fetal death within 1 week if BPP is normal~ 1/1300
- Modified BPP (mBPP)
- -NST & AFI
- -low false negative 0.8/1000
- -high false positives ~60%

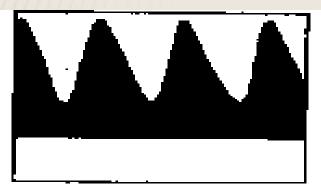
## DOPPLER VELOCIMETRY

- Measurement of blood flow velocities in maternal & fetal vessels
- Reflect feto-placental circulation
- Doppler indices from UA, Uterine A & MCA
- Doppler studies is mostly valuable IUGR
- In IUGR absent or reversed EDF (end diastolic flow) associated with fetal hypoxia

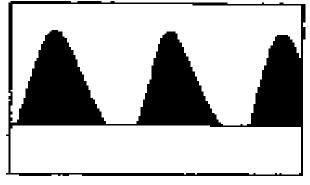
### **UMBILICAL ARTERY WAVEFORM**



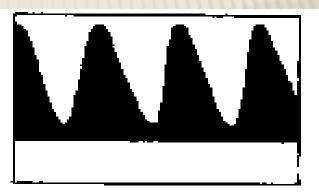
#### UMBILICAL ARTERY DOPPLER



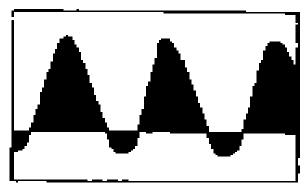
Normal pregnancy



Absent end diastolic velocity



Reduced end drastolic velocity

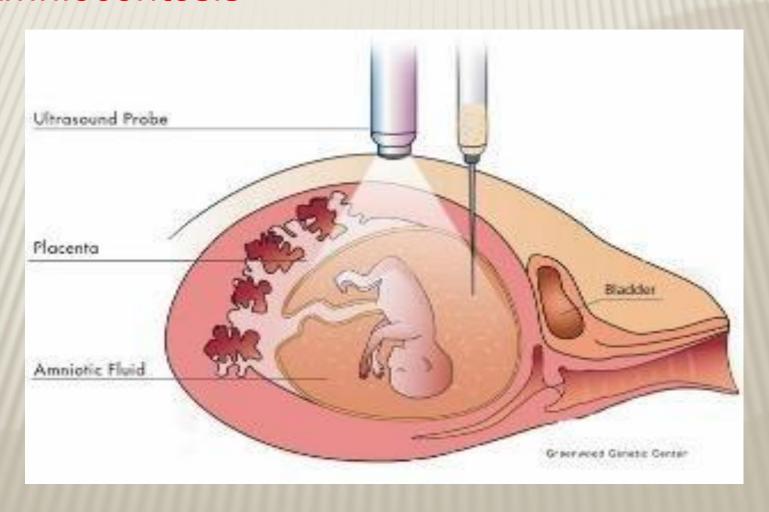


Reversed end diastolic velocity



## INVASIVE FETAL ASSESSMENT

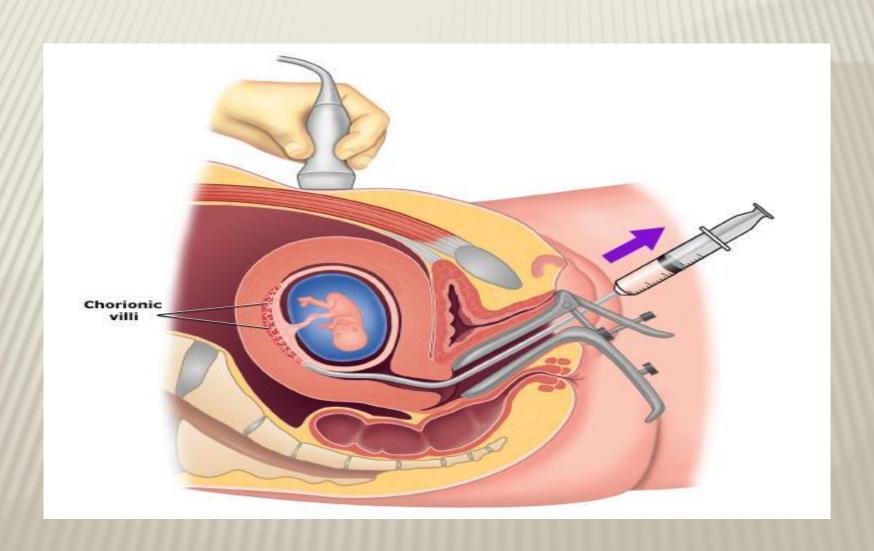
#### \* Amniocentesis



#### **AMNIOCENTESIS**

- Obtaining a sample of amniotic flui during pregnancy.
- Usullay done after 15w (can be done after 11w)
- Indication
  - -genitic (karyotype)
  - -billirubin level (RH-isimunisation)
  - -fetal lung maturity (L/S)
  - -therputic in polyhydramnios
- Risks: ROM ~1%, abortion 0.5%, infection 1/1000

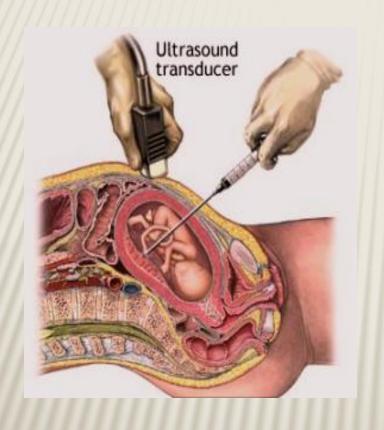
#### **× CVS** chorionic villus sampling



#### CVS CHORIONIC VILLUS SAMPLING

- ★ Usually done after 10w
- It is the procedure of choice for first trimester prenatal diagnosis of genetic disorders
- Complication: fetal loss (0.7 percent within 14 days of a TA CVS procedure and 1.3 percent within 30 days), Procedure-induced limb defects
- Second trimester amniocentesis is associated with the lowest risk of pregnancy loss; chorionic villus samplings safer than early (i.e, before 15 weeks) amniocentesis.

## CORDOCENTESIS





#### CORDOCENTESIS

- Indication: rapid karyotyping
  - -diagnosis of inherited disorders
  - -fetal HB assessment
  - -fetal plt level
  - -fetal blood transfusion

Complication: bleeding, bradycardia, infection....

#### FETAL DNA IN MATERNAL BLOOD

-Used if indication by ultrasound for testing or history of genetic disease

Result available in 2 weeks-

#### FETAL LUNG MATURITY

- A test for fetal lung maturity is performed before semi-elective but medically indicated births <39 weeks</p>
- \* Tests for fetal lung maturity are generally not performed before 32 weeks of gestation
- \* RDS develops as a consequence of surfactant deficiency and immature lung development.
- x L/S ratio is the most commonly used (ratio should be 2:1

## -FLM TESTING MAY HAVE VALUE IN THE FOLLOWING CLINICAL SITUATIONS:

- -Premature rupture of membranes (≥32 weeks) if FLM test is mature, delivery is likely safer than "wait and see" approach
- Assessment of need for NICU possible only if early delivery has medical mandate and time allows for FLM testing
- Other selected late preterm and early preterm pregnancy issues where FLM may guide management of at-risk pregnancy

### FETAL LUNG MATURITY FLM

All tests require amniocentesis for obtaining amniotic fluid

Comparison of FLM Laboratory Testing Options

Lamellar body count (LBC)

Phosphatidylglycerol (PG)

Lecithin-sphingomyelin ratio (L/S)

- Initial FLM of choice
- •Rapid, sensitive
- •New data indicates that one can estimate risk of respiratory distress syndrome (RDS) as a function of gestational age and LBC
- •Not useful unless gestational •Last test of choice
  - age ≥35 weeks
  - Limited availability
  - Sensitive

- Main role is in adjudication of immature LBC or PG
  - Labor intensive, imprecise
  - Limited availability
  - Results take >24 hrs unless performed at a local laboratory

## THANK YOU