



Antenatal Fetal Assessment

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References: 436 doctor's slides and notes, Kaplan

Color code: Notes | Important | Extra | Kaplan

Editing file:

Objectives:

Describe how to test for each of the following:

- 1. Fetal well-being.
- 2. Fetal growth.
- 3. Fetal movements.
- 4. Amniotic fluid volume.
- 5. Fetal lung maturity







Definition

- Fetal assessment is to identify fetuses at risk of neurologic injury or death in order to prevent prenatal mortality & morbidity.
- The most common reasons for fetal testing are decreased fetal movements, diabetes, postdates, chronic hypertension, and IUGR.
- It can be It can be divided into: early pregnancy fetal assessment, late pregnancy fetal assessment
- OR can be divided into: assessment of low risk pregnancy, assessment of high risk pregnancy

Rationale

If fetal oxygenation challenged:

- Blood flow directed to brain, heart & adrenal & blood flow away from the kidney → decrease fetal urine production → Decrease amniotic fluid (AF) volume (so AF volume is our 1st marker if the fetus is at risk)
- CNS hypoxia → Decreased Fetal movement.
- Ochemoreceptors \rightarrow vagally-mediated reflex \rightarrow Fetal heart rate abnormality (present as late deceleration).

Indication for antepartum fetal surveillance

Maternal	Pregnancy complication	
 Antiphospholipid syndrome Poorly controlled HTN Hemoglobinopathies (e.g. skill cell) Cyanotic heart disease Chronic renal disease Type 1 diabetes mellitus Hypertensive disorders 	 Preeclampsia Decreased fetal movement Oligohydramnios Polyhydramnios Intrauterine growth restriction (IUGR) Postterm pregnancy Isoimmunization Previous unexplained fetal demise Multiple gestation 	







Pregnancy Assessment

Early Pregnancy Assessment:

1) Fetal heart activity:

- Fetal auscultation (special stethoscope or Doppler) 12 weeks¹.
- Fetal heart activity seen by USS from 6 weeks.

2) Fetal movement

- Fetal movement are usually first perceptible to mother ~17w-20w.
 (Quickening) it takes time to feel the movement bc the uterus is small compare to the abdominal wall at 1st trimester
- 50% of isolated limb movements are perceived
- 80% of trunk and limb movements.

3) Fetal growth:

- Symphysial Fundal Height (SFH)
- UltraSound Scan (USS)

Late Pregnancy Assessment:

1) Fetal movement counting (kick chart):

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

¹ You need 12 weeks to listen bc there is the symphysis pubis of the mother, so when the uterus goes above the symphysis pubis then you can listen to the heart





Fetal movement counting technique			
CARDIFF TECHNIQUE	SADOVSKY TECHNIQUE		
 10 movements in 12 hours at least the mother felt more than 10 movements in 12 hours this is nor the abnormal is less than 10 in 12 If abnormal patient should count at then get further assessment (the 0) 	hour → still not felt → patient need more assessment. This technique is helpful bc some patients get busy so they cannot feel the movements.		

2) Contraction stress test (CST):2

- We Cause (induce) at least 2 uterine contractions over 20 minutes.
- Uterine contraction restricts O₂ delivery to the fetus.
 - Normal fetus will tolerate contraction. Negative CST requires absence of any late decelerations with contractions or sometimes acceleration
 - Hypoxic fetus will have late deceleration bc the blood vessels will be closed.
 Positive CST is worrisome. This requires the presence of late decelerations associated with at least 50% of contractions.
- High false positive rate ~50%.
- 100% true negative rate. That's why we don't use it these days unless you don't have US
- Contraindications: include previous classical uterine incision, previous myomectomy, placenta previa, incompetent cervix, preterm membrane rupture, and preterm labor.

3) Non stress test (NST)

- The first step in the assessment of fetal well-being is the NST.
- Main advantage over CST is no need for contraction.
- False +ve & false –ve higher than CST.
- The baseline 120-160 beats/minute.
- Different criteria in fetuses <32w.

The criteria vary by gestational age:

- <32 weeks, the increase should be ≥10 beats/min lasting ≥10 s
- >32 weeks, the increase should be ≥15 beats/min lasting ≥15 s

² sometimes called oxytocin challenge test bc we give oxytocin to contract the uterus.





•				
	Reactive		Non-reactive	
•	At least two accelerations from baseline of 15 bpm for at least 15 sec within 20 minutes Assessment: reassuring of fetal	another 20 min If non reactive istress test or bi	in 40 minutes proceed for cont ophysical profile (US). edictive value of NST to predict	raction
	well-being.		eping, immature, or sedated fetus	;
•	Follow-up: repeat	acidotic, compro	mised fetus?	
	weekly.	Follow-up: VAS "If still NR do CST	Vibratory Acoustic Stimulation".	
	FHH 100	(A) Reactive NST (B) Nonreactive NST	(A) Normal baseline range, and no UCs are present. Thus, only the NST component can be assessed. Because 3 accelerations are present, the assessment is reactive NST. This is a reassuring tracing. (B) Normal baseline range and no UCs are present. Thus, only the NST component can be assessed. Because no accelerations are present, the assessment is nonreactive NST. Because this is not a reassuring tracing, the next step should be a vibroacoustic fetal stimulation.	
	F441 190 100 100 100 100 100 100 100 100 10	Reactive (NST) Negative CST	(C) Normal baseline range and 4 UCs are present in 10 minutes. Thus, both the NST and CST components can be assessed. Because 3 accelerations are present, and no late decelerations are present, the assessment is reactive NST, negative CST. This is a reassuring tracing.	
	FHI 100 (D) No	a a a a a a a a a a a a a a a a a a a	(D) Normal baseline range and 4 UCs are present in 10 minutes. Thus, both the NST and CST components can be assessed. Even though no accelerations can be seen, no late decelerations are present. The assessment is nonreactive NST, negative CST. This suggests fetal sleep, sedation, or central nervous system (CNS) abnormality.	
	200 180 FHR 150 5pm 20		(E) Elevated baseline range and 4 UCs are present in 10 minutes. Thus, both the NST and CST components can be assessed. No accelerations can be seen. but	

4) Amniotic fluid index (AFI) very important

- The sum of the maximum vertical fluid pocket diameter in four quarters. You will put the probe in the 4 abdominal quarts and measure the vertical line in each quarter and sum them up
 - o The normal value: 5-25 cm
 - oligohydraminous: AFI < 5 cm. Consideration must be given to problems with urinary tract anomalies or renal perfusion.
 - polyhydraminous: AFI > 24 cm Consideration must be given to problems with decreased fetal swallowing or GI tract anomalies.







5) Biophysical profile (BPP) very important

Combines NST with USS estimation AFV, fetal breathing, body movement & reflex/tone/extension-flexion movement.

- It is a scoring system.
- It is done over 30 minutes.
- It measures acute hypoxia through (NST, body mov. and breathing) & chronic hypoxia (AFI).
- The risk of fetal death within 1 week if BPP is normal~ 1/1300.
- Modified BPP (mBPP): includes only the NST and AFI.
- Low false negative 0.8/1000, High false positives ~60%.

Fetal Biophysical profile/NST + AFI (extremely important):

Biophysical Variable (all seen by US)	Normal (score=2)	Abnormal (score= 0)
Fetal breathing Movements (FBM)	1 episode FBM of at least 30s duration in 30 min	Absent FBM or no episode >30s 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.

- Score of 8 or 10 —highly reassuring of fetal well-being. Management is to repeat the test weekly or as indicated.
- Score of 4 or 6 —worrisome. Management is delivery if the fetus is ≥36 weeks or repeat the BPP in 12–24 h if <36 weeks. An alternative is to perform a CST.
- Score of 0 or 2 —highly predictive of fetal hypoxia. Management is prompt delivery regardless of gestational age.





6) Doppler Velocimetry (UAV)

- Measurement of blood flow velocities in maternal & fetal vessels.
- Reflect fetoplacental circulation.
- Doppler indices from UA (Umbilical a.), Uterine a. & MCA (middle cerebral a. Of fetus).
- Doppler studies are mostly valuable for IUGR, as well as fetal anemia in alloimmunized pregnancies.

Normal pregnancy Reduce end diastolic velocity Systole Obastole (thick) Baseline (thick) If you see this you have to repeat it every 3-4 days to make sure if the baby going to the normal side or deteriorating Reversed end diastolic velocity Systole Systole Systole Systole Diastole (reversed)

which means the blood go from fetus to the

mother. this happen bc the fetus is hypoxic → asphyxia → more resistance → the blood can't go to the fetal circulation → return to the mother. This is very dangerous (it is the stage before fetal death) E.g. IUGR associated with fetal hypoxia.

UMBILICAL ARTERY DOPPLER:

during diastole there is no flow between the mother and

the fetus. E.g. IUGR associated with fetal hypoxia.

- This test measures the ratio of systolic and diastolic blood flow in the umbilical artery.
- The umbilical circulation normally has low resistance, so significant diastolic blood flow is expected.
- The systolic/diastolic (S/D) ratio normally decreases throughout pregnancy.
- This test is predictive of poor perinatal outcome only in IUGR fetuses.
- Nonreassuring findings, which may indicate need for delivery, are absent diastolic flow and reversed diastolic flow.





Interpretation of CTG			
FHR Baseline	 Normal Baseline FHR 110–160 bpm Moderate bradycardia 100–109 bpm Moderate tachycardia 161–180 bpm Abnormal "Severe" bradycardia < 100 bpm Abnormal tachycardia > 180 bpm 		
Variability	 Reduced variability. Baseline variability: Fluctuations in the baseline FHR that are irregular in amplitude and frequency. It is a reflection of the autonomic interplay between the sympathetic and parasympathetic nervous system. 	Reduced coming (Less Than 10 type over a period of time)	
Acceleration	Accelerations are always reassuring.	Assertion browing it transport receive of - greater han 16 fgm.	
Deceleration	 Early → Head compression. Late → Uteroplacental Insufficiency. Variable → Cord compression, Primary CNS dysfunction. Mixed → characteristics of any of the aforementioned patterns. 	180 PHR 100 C Early Decelerations – head compression 180 PHR 100 C Late Decelerations – uteroplacental insufficiency 180 PHR 190 Voriable Decelerations – uteroplacental insufficiency 180 Voriable Decelerations – uteroplacental insufficiency Voriable Decelerations – uteroplacental insufficiency Voriable Decelerations – uteroplacental insufficiency	
Tachycardia	 Severe Hypoxia Chorioamnionitis Maternal fever Mimetic drugs Fetal anemia, sepsis, HT failure, Arrhythmias. 		
Bradycardia	When FHR baseline is <110 beats/min Non-hypoxic explanations include:		

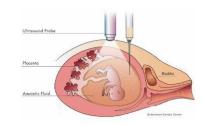




Invasive Fetal Assessment

1. AMNIOCENTESIS

- Definition: Obtaining a sample of amniotic fluid during pregnancy.
- When it's done? Usually done after 15w under ultrasound guidance without anesthesia (can be done after 11w)

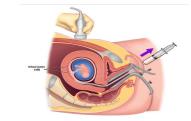


Indications:

- Genetic (karyotype). NTD (neural tube defect) screening is performed on amniotic fluid with biochemical analysis (AFP and acetylcholinesterase).
- Bilirubin level (in RH-isoimmunisation)
- Fetal lung maturity (L/S)
- Therapeutic in polyhydramnios to decrease the pressure tension on the mother
- Risks: ROM ~1%, abortion 0.5%, infection 1/1000.

2. CHORIONIC VILLUS SAMPLING (CVS)³

- Definition: It is the procedure of choice for first trimester prenatal diagnosis of genetic disorders. The procedure can be performed transcervically or transabdominally.
- When it's done? Usually done after 10w. (between 10 and 12w of gestation), why? bc if there is genetic disease the mother can terminate the pregnancy early which has less complication than if done later



- It is the procedure of choice for first trimester prenatal diagnosis of genetic disorders we do it for high risk mothers: previous history or family history of genetic diseases
- Second trimester amniocentesis is associated with the lowest risk of pregnancy loss; chorionic villus samplings safer than early (i.e, before 15 weeks) amniocentesis. so before 11 weeks we do CVS but after 11 weeks we do amniocentesis
- Complication:
 - Procedure-induced limb defects.
 - Fetal loss (0.7 % within 14 days of a TA CVS procedure and 1.3 % within 30 days).

³ The catheter is placed directly into placental tissue without entering the amniotic cavity.





3. CORDOCENTESIS

- usually done after the 1st trimester (after 13 weeks) but can be done at any time
- Indication:
 - Rapid karyotyping.
 - Diagnosis of inherited disorders.
 - Fetal HB assessment.
 - Fetal plt level.
 - Fetal blood transfusion
- Complication: Bleeding, Bradycardia, Infection.



MCQs

- 1) A 32-year old pregnant at 33-weeks came to the clinic with reduced fetal movements. Which one of the following is the best initial management for this patient?
- A- Caesarean section. B- Induction of labor. C- Non-stress test D. Ultrasound
- 2) During monitoring fetal heart rate on labor, you noticed repetitive deceleration after uterine contraction. What is the most likely diagnose?
- A- Congenital fetal cardiac block B- Cord compression
- C- Head compression D- Fetal hypoxia
- 3) 10 weeks of gestation presented to antenatal clinics for the first time. you did US which is useful for what in this stage:
- A- Gestational age.

 B- Congenital anomalies
- C- Locate placenta D- Amniotic fluid index
- 4) A 42-year old presented in early pregnancy. She is worried that her baby might have down syndrome. Which one of the following should be performed as screening method?
- A- Amniocentesis B- Chorionic villus sampling
- C- Ultrasound scan D- Umbilical blood sampling
- 5) Which one of the following will be obtained by doing a first trimester ultrasound scan for twin pregnancy?
- A- Determination of fetal presentation. B- Localization of cord insertion.
- C- Placental Localization. D- Determine chorionicity.

Answers: 1- C. 2- D. 3- A. 4-C. 5- D.