Assessment of Fetal Health

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Assessment of Normal Fetal Physiology

- Maternal evaluation of fetal activity
- Non stress testing
- BPP
- Doppler



Fetal heart rate changes during development

- Gradual decrease in heart rate
- Increase in variability
- Increasing responsiveness to acute changes in fetal status (including accelerations and decelerations)

The ability of the fetus to accelerate its heart rate in response to movement is related to fetal oxygenation and metabolic state

This provides the basis for nonstress testing.



Fetal Behavior

The presence of normal fetal muscle tone, gross body movements, and breathing have been reliably tied to the absence of fetal hypoxemia and academia

This is the basis for the BPP

Near term, periods of inactivity that correspond to episodes spent in the quiet sleep state, as illustrated by the nonreactive nonstress test (NST), can be confused with fetal compromise.



This illustrates the importance of using multiple modalities of fetal assessment to decrease the false positive rate of any one test.



Fetal Heart Rate Monitoring

Many studies have shown it to be the most sensitive short term predictor of worsening hypoxemia or acidosis, it has become part of fetal monitoring of labor and delivery.

However, preterm fetuses, fetuses with intrauterine growth restriction (IUGR), or fetuses exposed to maternal medication such as narcotics or magnesium sulfate frequently have paired FHR acceleration and fetal movements that do not meet these criteria. Modification of these criteria based on gestational age.



False reassuring NST results

Defined by fetal death with in one week, occurred at a rate of 1.9:1000 fetuses in the largest study.

Programs incorporating maternal counting of fetal movements and other published protocols for fetal assessment (in most cases, the addition of an assessment of amniotic fluid volume) may help to identify additional pregnancies at risk. About 10% to 12% of fetuses in the third trimester do not meet these criteria at 30 minutes but this number falls below 6% by 40 minutes.

The most common explanation for a non reactive NST result is a sleep cycle in a normal fetus that is longer than average.

Ultrasound evaluation with a BPP should be available as the back up test. Ultrasound provides additional fetal evaluation that may help to diagnose the reason for an apparently abnormal NST result.



Bradycardia

Congenital heart block and serious fetal compromise

Maternal hypothermia under general anesthesia

□ Sustained fetal bradycardia in severe pyelonephritis

□ Baseline FHR < 110 bpm



Tachycardia

□ Baseline FHR > 160 bpm

Imaternal fever from chorioamnionitis (most common) with no fetal compromise unless there are associated periodic heart rate changes or fetal sepsis

🗆 fetal compromise

□ cardiac arrhythmias

maternal administration of parasympathetic (atropine) or sympathomimetic (terbutaline)

maternal hypotension caused by epidural analgesia

The key feature to distinguish fetal compromise in association with tachycardia seems to be concomitant heart rate decelerations (tachycardia + deceleration)



Beat-to-Beat Variability

- an important index of cardiovascular function
- regulated largely by the autonomic nervous system
- Short-term variability
- Long-term variability

most clinical interpretation is based on visual analysis with subjective judgment of the smoothness or flatness of the baseline









The normal frequency of such waves is 3-5 cycles per minute



Causes of decreased variability

Prematurity

- Fetal metabolic acidosis
- •CNS depressants
- Fetal sleep cycles (after 30 wk)
- Fetal tachycardia
- Neurologic abnormality
- Betamethasone
- Analgesic
- Mgso4





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Decreased Variability

- Diminished beat-to-beat variability can be an ominous sign indicating a seriously compromised fetus
- loss of variability + decelerations

Cause: Maternal acidemia



fetal acidemia

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- beat-to-beat variability is affected by a variety of pathological and physiological mechanisms
- Variability has considerably different meaning depending on the clinical setting
- The development of decreased variability in the absence of decelerations is *unlikely to be due to fetal hypoxia*
- persistently flat fetal heart rate baseline—absent variability—within the normal baseline rate range and without decelerations may reflect a previous insult to the fetus that has resulted in neurological damage



Biophysical Profile

The BPP relies on the premise that multiple parameters of well being are better predictors of outcome than any single parameter.

Amniotic fluid measurement

Fetal breathing movements Rhythmic fetal diaphragm contractions or hiccups lasting more than 30 seconds.

Fetal movement and tone

Factors that influence Biophysical Profile scoring performance

Fetal Effect

TABLE Factors That Influence Biophysical Profile 34.3 Scoring Performance

Agent

Drugs Sedatives or sedative side effects (e.g., Aldomet) Excitatory (e.g., theophylline) Street drugs (e.g., crack cocaine) Indomethacin Maternal cigarette smoking

Maternal hyperglycemia (iatrogenic or unregulated)

Maternal hypoglycemia (e.g., poor nutrition, insulin excess) Single parameter removed by perinatal condition

Persistent fetal arrhythmia Spontaneous premature rupture of membranes Acute change in status (e.g., eclampsia, abruptio placentae, ketoacidosis) Corticosteroids Diminished activity of all varieties; abolition of none Continuous, "picket fence" **FBMs** Rachitic, rigid, furious, bizarre **FMs** Oligohydramnios Various observations; FBMs abolished or attenuated but some report no change; FMs reduced Sustained FBMs or acidosis, diminution or abolition of FMs or FT or NST reactivity Abnormal paucity of all behaviors, normal AFV

Uninterpretable NST Obligatory oligohydramnios

May invalidate predictive accuracy

Transient decrease in FBMs

Vibroacoustic Stimulation

Contraction stress test and oxytocin challenge test



Doppler ultrasound

Umbilical artery





Middle Cerebral artery





Ductus venosus





Umbilical Vein



Practical Aspects of fetal testing

 Impacts of monitoring on perinatal mortality and long term outcomes

