Intrapartum Fetal Monitoring (CTG)

- •5th year Medical Students
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OBJECTIVES

- •Discuss the modality used to monitor fetal wellbeing in labor
- How to read a normal CTG?
- List causes of abnormal CTG in labor

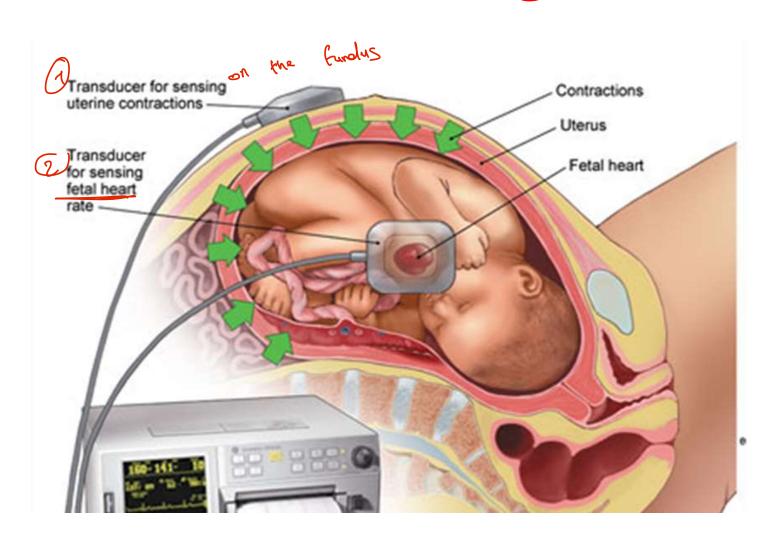
Fetal Heart Rate (FHR) Monitoring

• Fetal heart rate (FHR) monitoring can be done by:



- 1) External mostly used
- 2) Internal to the scalp of the baby

External Electronic Continuous Fetal Monitoring



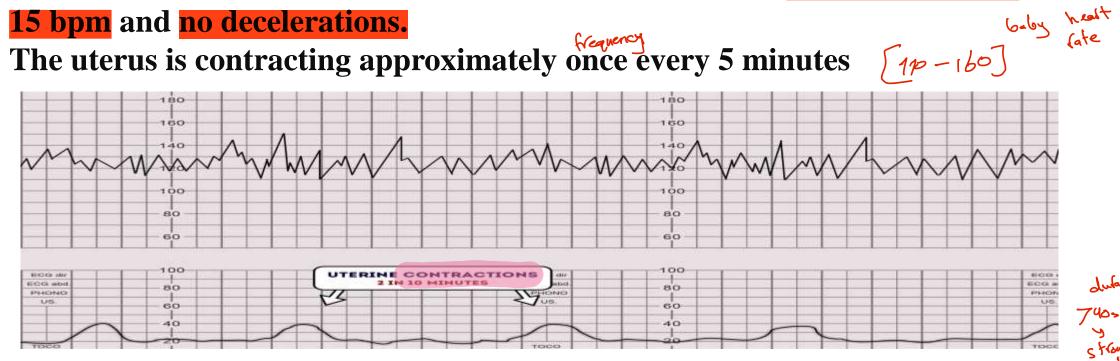
CTG

• The Cardio Toco Graph (CTG) has two components:

- Contractions
- •Fetal Heart

Normal CTG

A normal cardiotocograph (CTG), showing a baseline fetal heart rate of approximately 120 bpm, frequent accelerations, baseline variability of 10-



How to read a CTG?

- Name of the patient & Date
- Define the risk
- Contractions (frequency & duration)
- Baseline FHR (normal 110 bpm 160 bpm)
- FHR variability (beat to beat variation) [5- 25]
- Acceleration (present or absent)
- Deceleration (Early, Late & Variable, Prolonged)
- Assessment & plan of management
- The overall impression can be described as either reassuring, suspicious or abnormal.

Assessment of uterine contractions

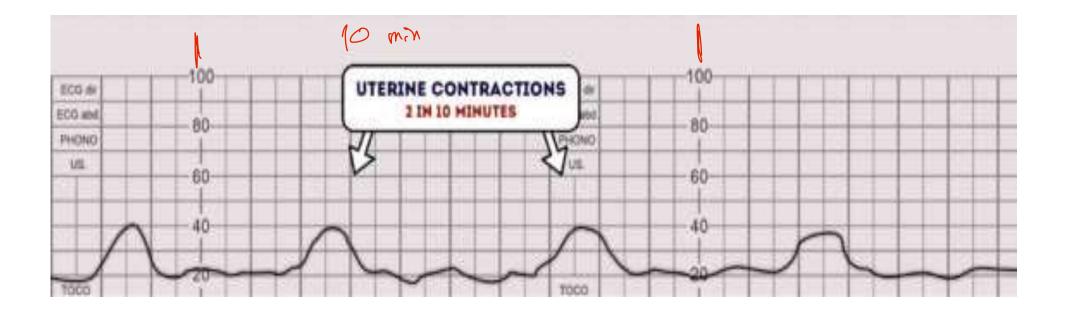
- The lowest intrauterine pressure between contractions is called resting tone.
- Normal resting tone is 5-10 mmHg, but during labor it may rise to 10-15 mmHg, in Labor
- Pressure during contractions rises to ~25-100 mmHg (varies with stage).
- A resting pressure above 20 mmHg causes decreased uterine perfusion.

Montevideo Units

- Montevideo Units (MVUs) are a standardized way to quantify the strength of uterine contractions during labor.
- It is calculated by internally (not externally) measuring peak uterine pressure amplitude (in mmHg).
- MVUs = The sum of contraction intensities (in mmHg) above baseline tone during a 10-minute window.
- Subtracting the resting tone of the contraction, and adding up the numbers in a 10-minute period.
- Generally, above 200 MVUs is considered necessary for adequate labor during the active phase

Contractions

- The number & duration of contractions in a 10-minute period is recorded.
- Hyperstimulation: More than 5 contractions in 10 minutes.
- Hyperstimulation can result in hypoxia, acidosis & abnormal fetal heart rate



CTG

10 week - we see it

- Comment on:
- Baseline (110-160 bpm)
- Variability (5-25 bpm)
- Acceleration
- Deceleration (Early (I), Variable (II), Late (III), Prolonged

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BASELINE RATE

• Normal fetal heart rate: Between 110-160 bpm

Accelerations: Abrupt increase in the fetal heart rate above the baseline.

Gestational Age Specific:

≥32 weeks: 15 bpm for ≥15 seconds

<32 weeks. 10 bpm for \geq 10 second

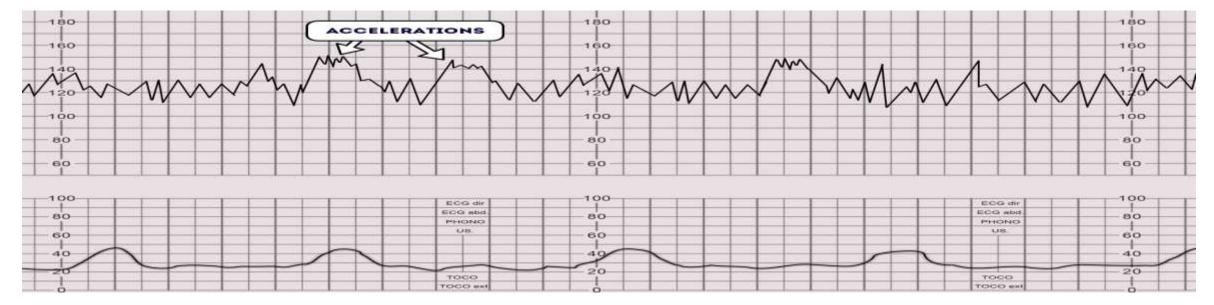
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The presence of accelerations is reassuring

The absence of accelerations with an otherwise normal CTG is of uncertain significance

we must have a least 2 accept than within 20 minutes

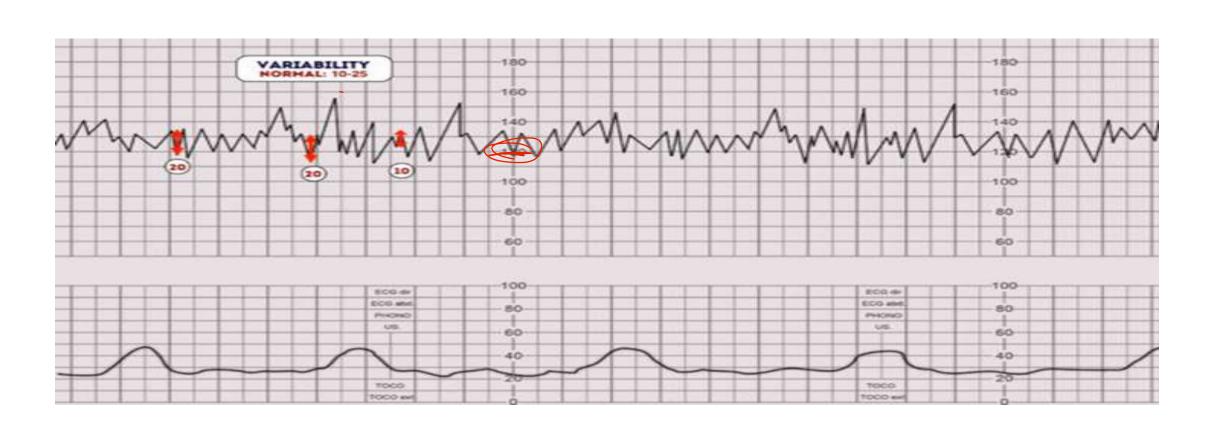




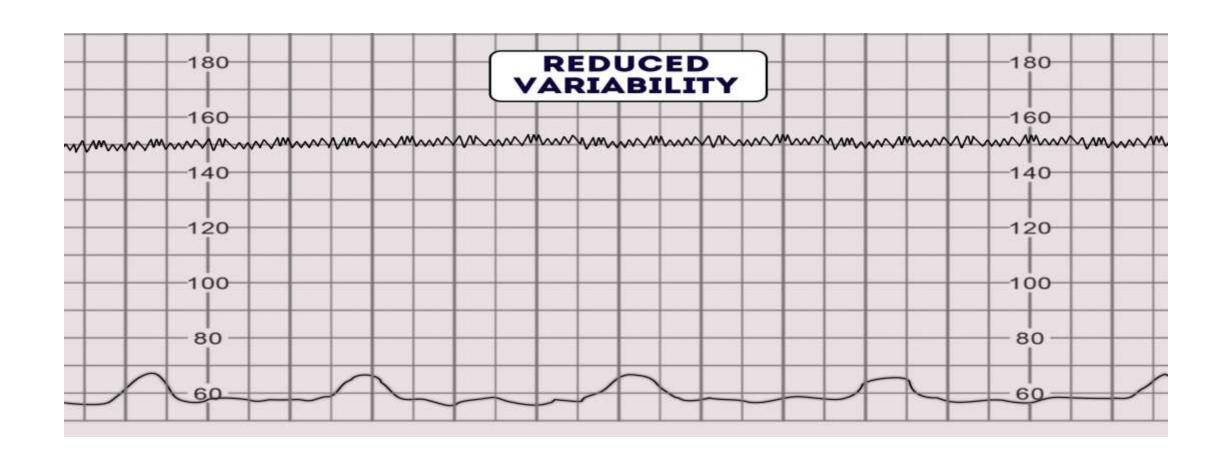
Variability

- It refers to the fluctuations in the baseline FHR
- Variability occurs as a result of the interaction between the nervous system, chemoreceptors, baroreceptors and cardiac responsiveness.
- Normal variability indicates an intact neurological system in the fetus
- Persistently minimal or absent FHR variability appears to be the most significant intrapartum sign of fetal compromise. Unless the baby is seeling (40 min
- Normal variability is between 5-25 bpm.
- Absent variability = Amplitude range undetectable
- Minimal = < 5 BPM \checkmark
- Moderate = 6 to 25 BPM
- Marked = > 25 BPM

Normal Variability Normal: (5-25 bpm)



Reduced variability

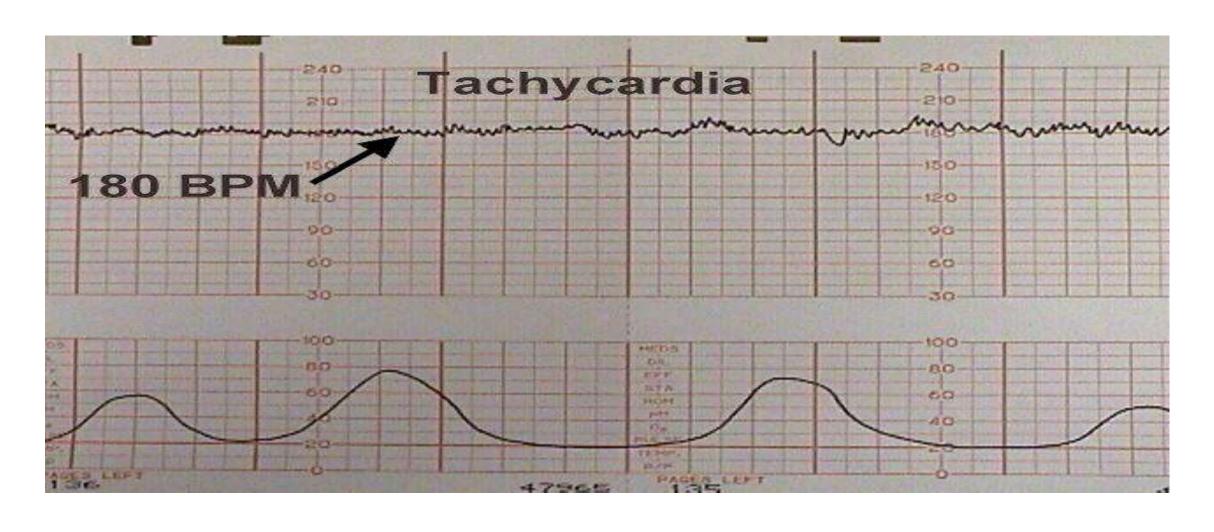


Reduced variability

- Reduced variability can be caused by any of the following:
- Fetal sleeping: this should last no longer than 40 minutes (this is the most common cause)
- Fetal acidosis (due to hypoxia): more likely if late decelerations are also present
- Fetal tachycardia
- Drugs: opiates, benzodiazepines, methyldopa, dexamethasone and magnesium sulphate.

 Mesent and West eclambia + used in pre-term
 - **Prematurity:** variability is reduced at earlier gestation (<28 weeks)
 - Congenital heart abnormalities

Fetal Tachycardia



Fetal Tachycardia

- Tachycardia: A baseline value above 160 bpm lasting more than 10 minutes.

 Chorioamnionitis (also called intra-ar
- Causes of fetal tachycardia include:
- Maternal pyrexia (infection, chorioamnionitis)///

• Fetal hypoxia.

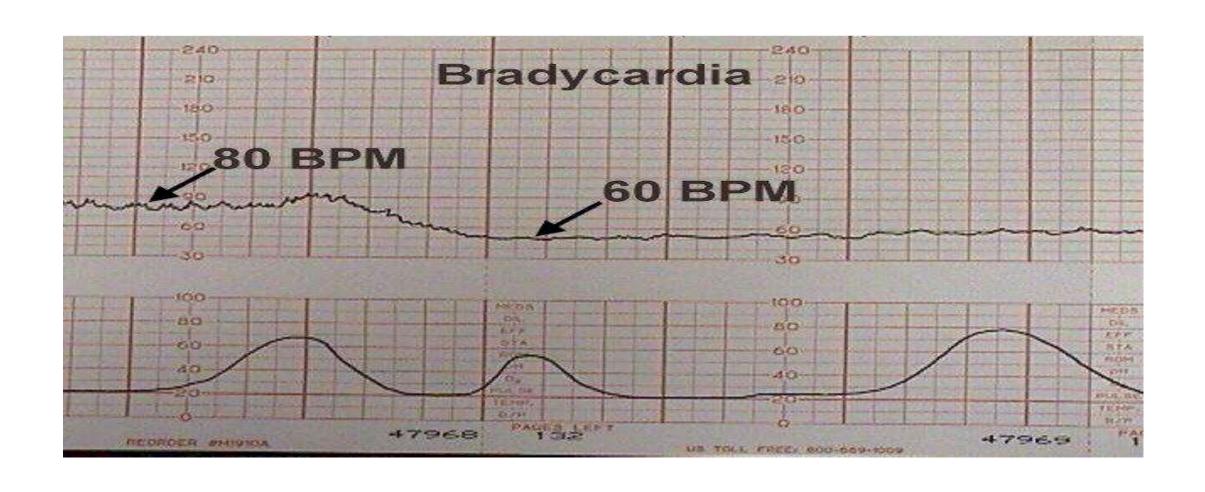
• Medications: (beta-agonist (Ritodrine (Yutonar) Terbutaline (Bricanvl) (drugs used for tocolysis) (Atropine) (Sympathomimetic drugs) (Parasympatholytic drugs)

- Fetal arrhythmias (SVT).
- Fetal or maternal anemia.
- Fetal hypoxia
- Hyperthyroidism
- Prematurity

Chorioamnionitis (also called intra-amniotic infection) is a bacterial infection of the fetal membranes (chorion and amnion), amniotic fluid, and sometimes the placenta. It typically occurs before or during labor, especially with prolonged rupture of membranes.

tocolytic agents

Fetal Bradycardia



Fetal Bradycardia

- Bradycardia: A baseline value below 110 bpm lasting more than 10 minutes.
- Values between 100 and 110 bpm may occur in normal fetuses, especially in postdate pregnancies.
- Causes of fetal tachycardia include:
- Maternal hypothermia, Hypotension, Hypoglycemia
- Administration of beta-blockers
- Fetal arrhythmias such as atrioventricular block
- (SSA/Ro positive pregnancies (systemic lupus erythematosus (SLE) and Sjögren's syndrome (SS)) associated with heart block)
- Fetal metabolic acidosis late hyperia

AJ block

Decelerations

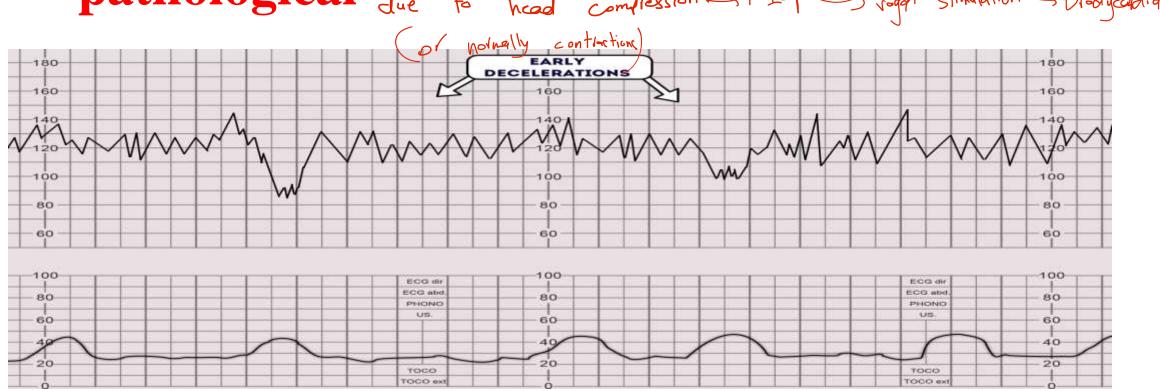
• Decelerations are an abrupt decrease in the baseline fetal heart rate of greater than 15 bpm for greater than 15 seconds.

- Early (Type I)
- Variable (Type II)
- Late (Type III)
- Prolonged decelerations

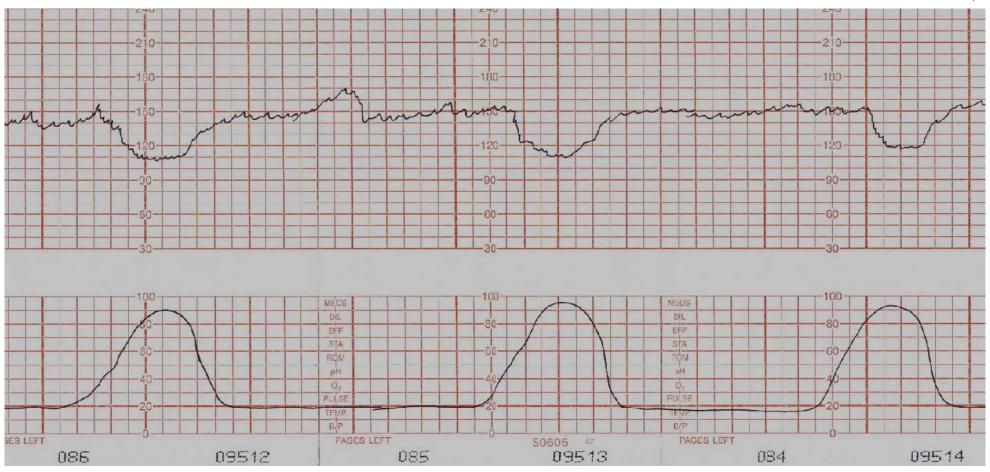
• Early deceleration caused by head compression

It is considered to be physiological and not pathological

| The proof struction - blodycadig



Early decelerations -good best to best vorsion -no accularior (not abroing)

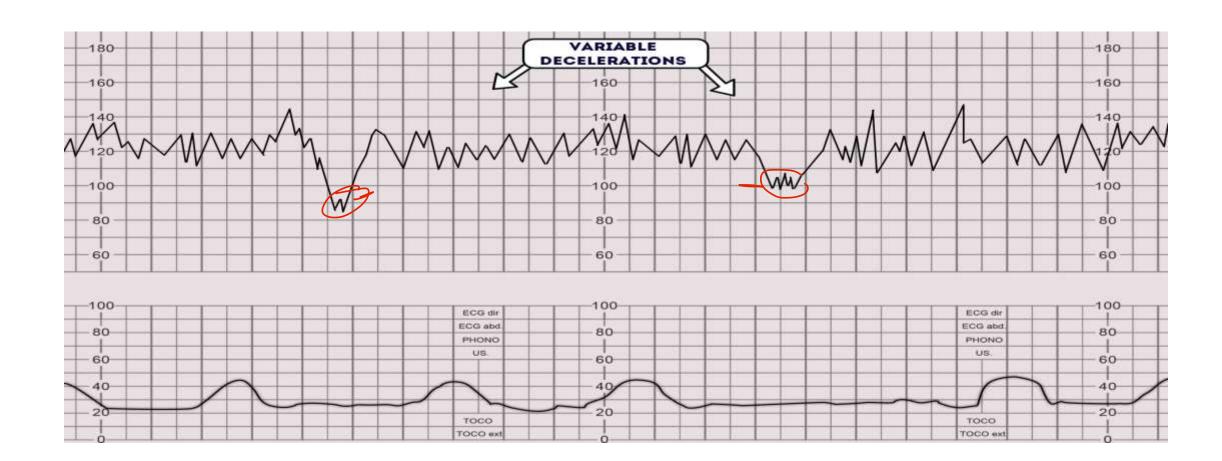


Variable decelerations caused by umbilical

- depth

cord compression

-location
-shape



Variable Decelerations

• Variable decelerations are V-shaped. Variable in onset, duration and depth. They may occur with contractions or between contractions.

• Typically, they have an abrupt onset and rapid recovery (in contrast to other types of decelerations which gradually slow and gradually recover).

Variable Decelerations

cold complexsion

- The accelerations before and after a variable deceleration are known as the shoulders of deceleration.
- Their presence indicates the fetus is not yet hypoxic and is adapting to the reduced blood flow.
- The presence of persistent variable decelerations indicates the need for close monitoring.
- Variable decelerations without the shoulders are more worrying, as it suggests the fetus is becoming hypoxic

Management of Variable Decelerations

- Changing maternal position to improve uterine blood flow
- IV hydration to increase maternal blood volume, presumably leading to increased uterine blood flow
- Administering oxygen to the mother to try to get some additional oxygen through to the fetus. (The least useful).

 • Decreasing or discontinuing oxytocin infusion to slow down or stop
- contractions that are provoking the decelerations.

 We may consider Amnioinfusion to improve oligohydramnios
- We may consider Tocolytic drugs to slow down or stop contractions that are provoking the decelerations.
- Digital elevation of the fetal head out of the maternal pelvis to ease pressure on the umbilical cord left lateral position a material position

Management of Variable Decelerations

- Occasional mild or moderate variable decelerations are common and not considered threatening.
- Severe variable decelerations dip below 60 BPM for at least 60 seconds ("60 x 60").
- If persistent and not correctable by simple means, they can be threatening to fetal well-being.
- Like persistent, non-remediable late decelerations, fetuses demonstrating persistent, non-remediable severe variable decelerations should be delivered promptly.

Late decelerations

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Uteroplacental insufficiency is a failure of the placenta to deliver enough oxygen and nutrients to the fetus.

Cause: Uteroplacental Insufficiency

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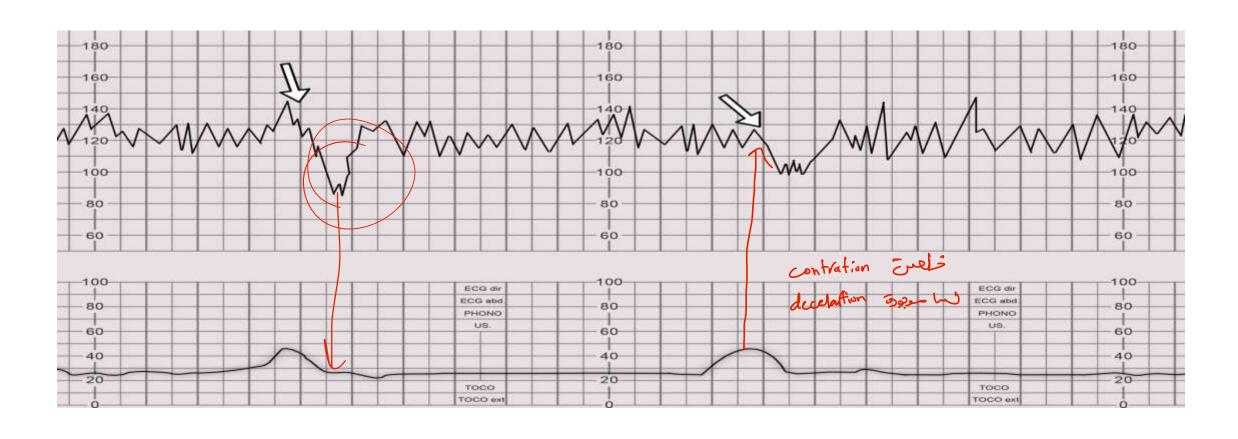
• Insufficiency is caused by uterine tachysystole (uterine hyperstimulation), maternal hypotension, epidural or spinal anesthesia, IUGR, hypertension & preclampsia, intraamniotic infection, or placental abruption.

- Late decelerations begin at the peak of the uterine contraction and recover after the contraction ends
- Severe, repetitive late decelerations usually indicate fetal metabolic acidosis.

Compensate

inth whome

Late Deceleration



Management of Late decelerations

- supplessure of Isc -> A serious detain -> for to mom and baby
- Maternal left lateral position to improve uterine blood flow
- Correct maternal hypotension with IV fluids
- ✓ Administer O2 by mask
- Stop oxytocin infusion
- Consider tocolytic drugs to slow down or stop contractions that are provoking the decelerations
- Vaginal examination (To evaluate cervical condition & descent)

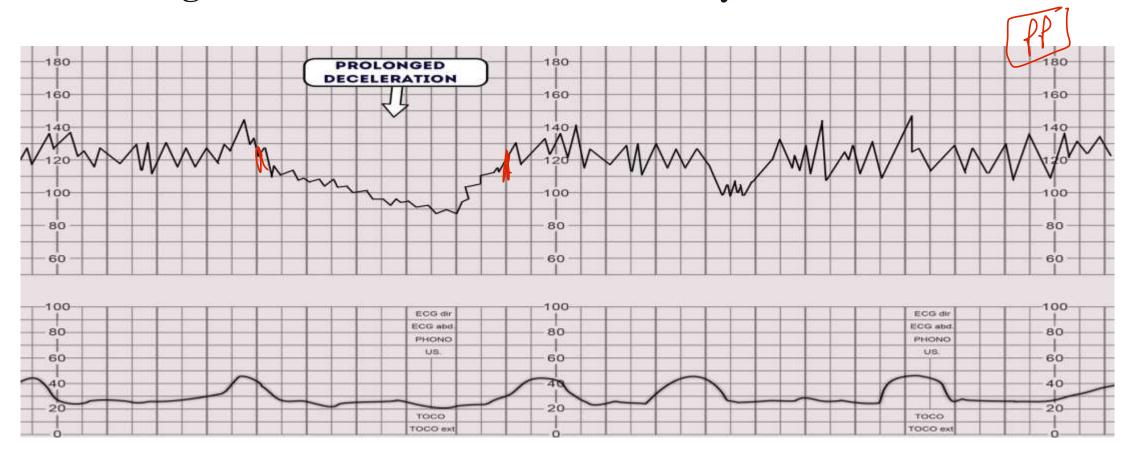
dilation

- If persistent perform fetal scalp PH
- If persistent consider immediate delivery

Prolonged deceleration:

If it lasts between 2-3 minutes it is classed as non-reassuring.

If it lasts longer than 3 minutes it is immediately classed as abnormal.



Causes of prolonged decelerations

- Severe bradvcardia: FHR less than 80 beats per minute. lasting longer than 3 minutes is an ominous finding and may be associated with fetal acidosis.
- Could be transient due to:
- 1. Supine hypotension -> ateris compless Isc
- 2. Paracervical block
- 3. **Epidural** and spinal anesthesia
- Other important possible causes:
- ► Prolonged cord compression ✓ Msofty





Rapid fetal descent may cause strong vagal response



• If associated with vaginal bleeding we have to think of: Major abruptio placenta, rupture uterus and vasa previa.







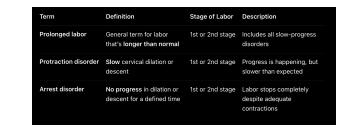
A sinusoidal pattern usually indicates one or more of the following:

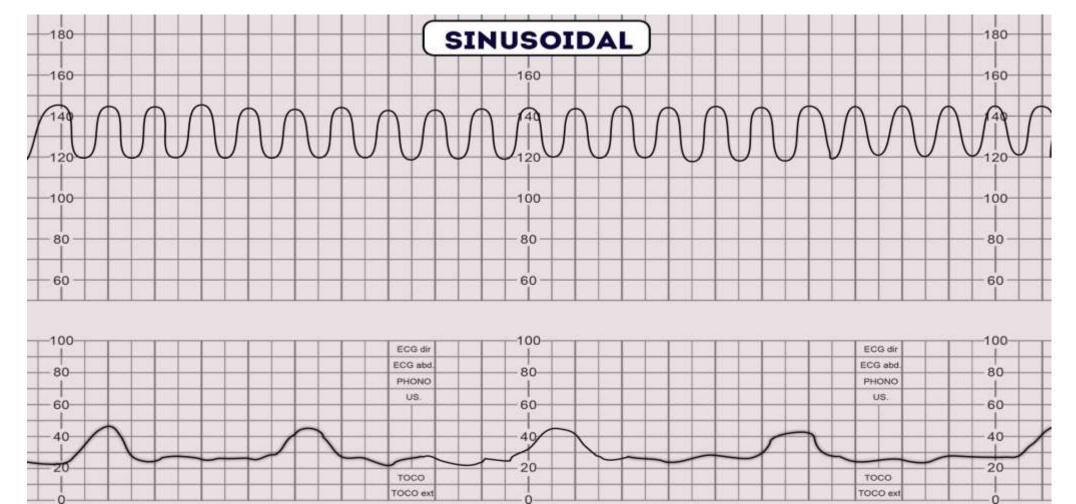
Severe fetal hypoxia

Severe fetal anaemia (Rh disease)

Fetal/maternal haemorrhage







Causes of Sinusoidal Pattern

• Sinusoidal pattern is **very concerning** as it is associated with high rates of **fetal morbidity** and **mortality**.

- It occurs in association with:
- Severe fetal hypoxia
- Severe fetal anemia
- Fetal-maternal hemorrhage
- Ruptured vasa previa

Remember

- Uterine hyper-contractility is the most frequent cause of a pathological CTG
- Baseline and variability are the most important features on a CTG they are indicative of hypoxia.
- Remember a fetus will protect its heart muscle as a priority.....the other organs and the brain will suffer the hypoxia first.
- Interpret the CTG in the full clinical context and understanding of the fetal reserve.

Category1: Strongly predictive of normal fetal acid-base status at the time of observation. Requires no specific management

hormal CTG

Category I (Normal)

Category I FHR tracings include all of the following:

- Baseline rate: 110-160 beats/min
- Baseline FHR variability: moderate
- Late or variable decelerations: absent
- Early decelerations: present or absent
- Accelerations: present or absent

Category II FHR (Intermediate)

- The fetal heart rate tracing shows **ANY** of the following:
- Tachycardia, bradycardia without absent variability, minimal variability, absent variability without recurrent decelerations, marked variability, absence of accelerations after stimulation, recurrent variable decelerations with minimal or moderate variability, prolonged deceleration > 2minute but less than 10 minutes, recurrent late decelerations with moderate variability, variable decelerations with other characteristics such as slow return to baseline, and "overshoot".
- Not predictive of abnormal fetal acid-base status, but requires continued surveillance and reevaluation.

Category II FHR (Intermediate)

- X
 - Not predictive of abnormal fetal acid-base status, so management depends on clinical circumstances.
 - Search for the underlying cause (ex: maternal hypotension) and correct it.
 - Some interventions:
 - Change the mother's position to the left lateral recumbent.
 - Reduce the infusion rate of oxytocin
 - Increase intravenous fluids.

<u>Category 3:</u> Predictive of abnormal fetal acid-base status. In addition to measures in category II treatment, may consider:

Fetal scalp blood sampling Ultrasound doppler velocimetry

Delivery
almo(m) pathological

Category III (Abnormal)

Category III FHR tracings include either of the following:

- Absent baseline FHR variability and any of the following:
 - Recurrent late decelerations
 - Recurrent variable decelerations
 - Bradycardia
- Sinusoidal pattern

Secondary Tests of Fetal Wellbeing

- ·Fetal Scalp Sampling we don't do it
- Scalp stimulation
- Acoustic stimulation

the use of sound or auditory stimuli to elicit specific responses or effects in individuals

- •Fetal pulse oximetry
- •Fetal Electrocardiogram Analysis

Fetal Blood Sampling (Fetal Scalp PH)

- FBS may be used in cases of abnormal CTG.
- A vaginal examination needs to be performed prior to the procedure to assess the nature and position of the presenting part.
- <u>Contraindications</u>: <u>maternal infection</u>, women seropositive to hepatitis B, C, or to HIV, suspected fetal blood disorders, uncertainty about the presenting part, preterm fetus.
- CTG + FBS results in a reduction in cesarean deliveries when compared with CTG alone

Fetal Scalp PH abridian To

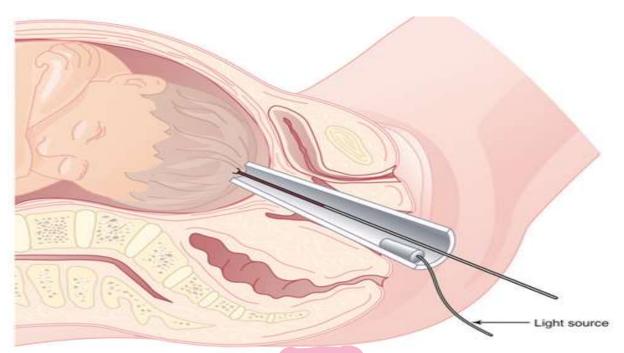


FIGURE 9-6 This technique of fetal scalp blood sampling via an amnioscope is still used in many centers. After making a small stab incision in the fetal scalp, the blood is drawn off through a long, heparinized capillary tube.

Fetal Blood Sampling (Fetal Scalp PH)

	Interpretation	рН	Lactate
			(mmol/L)
	Normal	≥ 7.25-7.35	< 4.2
	Repeat in 30 mins	7.21 – 7.24	4.2 - 4.8
	Birth expedited	≤ 7.20	> 4.8
	Urgent delivery	< 7.15	> 5.0

abnormal e

Dr Amal Barakat

Thank you The End of Intrapartum Fetal Monitoring