



Drugs Acting On Uterus

By

Dr.Nashwa Abo-Rayah

associate prof. of clinical & experimental pharmacology

Mutah University- Faculty of Medicine- JORDAN

2024/2025



Objectives

- 1- Uterine contractions
- 2- Drugs affecting uterine contractions
- 3- Oxytocin: actions, mechanism of action, kinetics, indications, side effects , precautions and contraindications
- 4- Ergometrine: mechanism of action, uses, side effects and contraindications
- 5- Prostaglandins PGs: dinoprostone, gemeprost, misoprostol and dinoprost
- 6- Tocolytic drugs

Uterine contractions

Contracted at all time (mild)

- Uterine smooth muscle is characterized by high level of spontaneous contractile activity.

- It is innervated by autonomic nervous system
- Uterine contractions are muscle contractions of the uterine smooth muscle that occur during:

- Menstrual cycle

- Ovulation

- Pregnancy → mild intensity, irregular (braxton hicks)

- Labor → high intensity, regular

- Sexual stimulation & during lactation: due to oxytocin (love hormone)

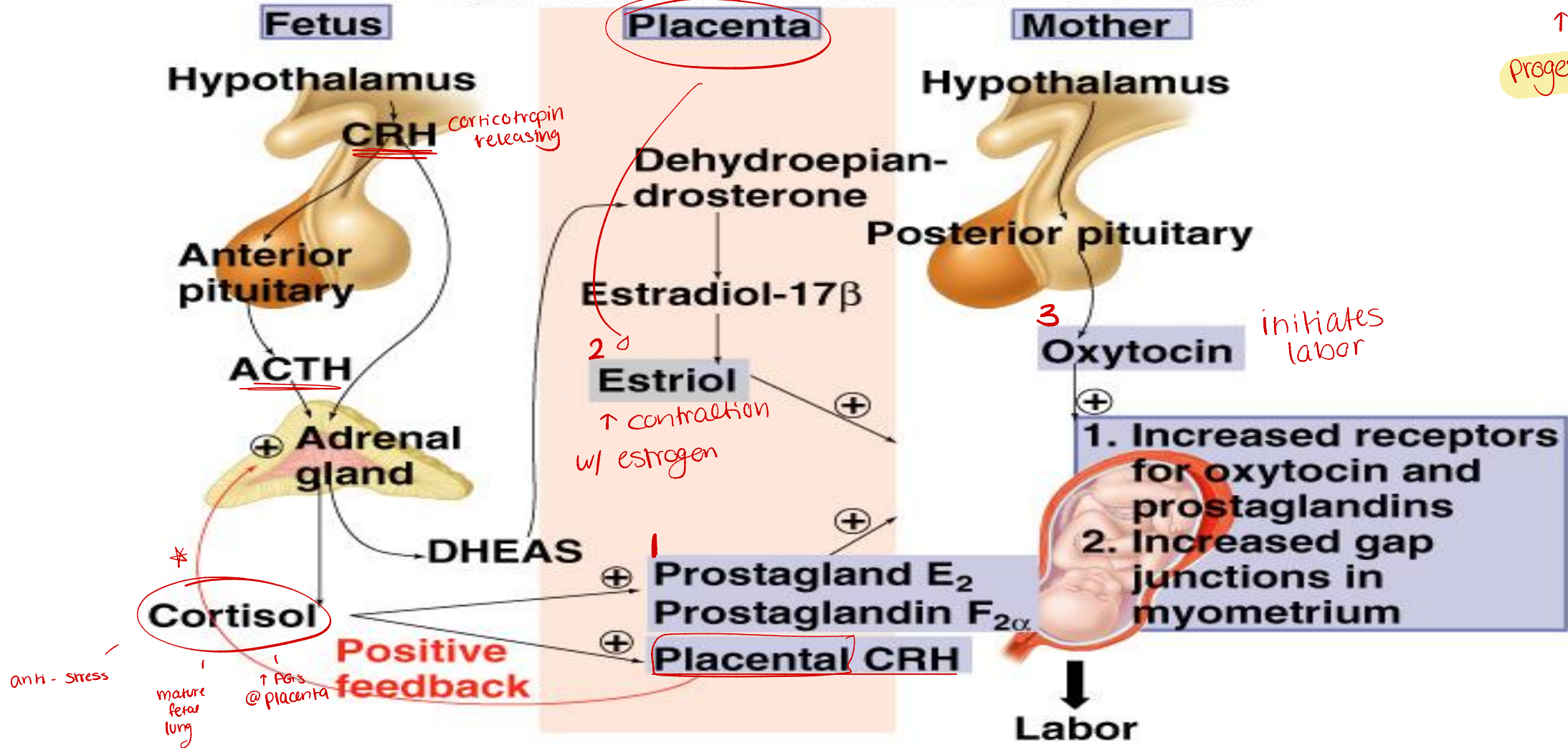
Parturition

Pregnancy hormones to
stop contractions

↑

Progesterone

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.



Drugs affecting uterus

Stimulatory

The main drugs used clinically to increase uterine contractility:

- Oxytocin
- Alph₁- Adrenoceptor agonists (ergot derivatives)
- PGE₂ or PGF_{2α}

- Inhibitory ^{↗ relaxant}
- B2- adrenoceptor agonists
- Calcium channel blockers
- Atosiban
- **Indications:**
- Prevention of preterm labor [⊗]

Indications of uterine stimulants

for all
↑ 3 groups

- 1- Induce or facilitate labor:

① • ^{<39w}Pre-term: diabetic mother- pre-eclampsia- Rh negative fetus ^{تسليم الحمل}

② • Incomplete abortion ^{→ post spontaneous abortion to clean up & avoid fibrosis}

③ • At-term: uterine ^{inert/ inactive}inertia Full, 39w

④ • ^{>39w}Post-term: delayed labor

- 2- Prevention of postpartum hemorrhage

- 3- Induction of abortion

1. Oxytocin

NOT
in exam
^

1. Oxytocin (Pitocin, Syntocinon) : Oxytocin and vasopressin are ^{qx} nonapeptide hormones, synthesized in hypothalamus, then transported to posterior pituitary where they are stored and released.

w/ vasopressin !
=
adverse effects

Pharmacological actions of oxytocin :

1. **Contraction of myoepithelial cells** surrounding secretory alveoli of breast leading to milk ejection in lactating females.

2. Induction of intermittent! uterine contractions and maintainance of labor:

- It contributes to initiation of parturition.
- Reaches peak during pushing phase of labor
- Oxytocin-induced contractions can be inhibited by:
- B₂-adrenoceptor agonists or by general anesthetics.

constant + durations
+
regular

example:
want to do C-section
instead of natural

3. Uterine involution

post-stretch to ↓
size back to normal
size, 60g in 6w

4. Oxytocin has weak antidiuretic or pressor activity.

] since similar to
vasopressin

Mechanism of action:

1. Stimulation of oxytocin receptors
2. Increasing PGs levels intrauterine

PKs of oxytocin

q-amino → protein
↑

- It is not given orally since it is destroyed by proteolytic enzymes of stomach and intestine (trypsin and chymotrypsin).
- It is **NOT** bound to plasma proteins
- Eliminated by liver and kidney (plasma $t^{1/2} \sim 5$ min): **IV** infusion.

Therapeutic uses of oxytocin

Induction of labor: given by IV infusion in:

- 1- Conditions requiring early vaginal delivery at 37-38 weeks: maternal diabetes, pre-eclampsia, Rh-immunization
- 2- Primary uterine inertia, and to enhance uterine contractions in **incomplete abortion** and **full-term labor**
- 3- **Delayed onset of labor at term**: post-maturity

Postpartum hemorrhage

Control of post-partum hemorrhage (PPH): (by IV infusion or IM injection with ergonovine)

alone
↑

To induce milk let-down after labor: by nasal spray.

N.B. Clinically oxytocin is given only when uterine cervix is soft and dilated !

Adverse effects:

⊗ intermittent anymore

Rare with proper supervision

1- With large IV infusion doses, **tetanic uterine contractions** can occur which obstructs intramural uterine blood flow causing:

90-100s, ⊗ intervals [continuous contraction] ↑ pressure artery

- fetal distress or death.
- Uterine rupture may occur esp. with obstructed labor.

2- with large doses , blood pressure increases due to vasoconstriction

3- Water intoxication can rarely occur due to large volume of IV infused fluid.
(may be fatal)

4- Increased incidence of neonatal jaundice: due to increased osmotic fragility of RBC + rupture of RBC release

Contra-indications:

1- Fetal distress too risky ~

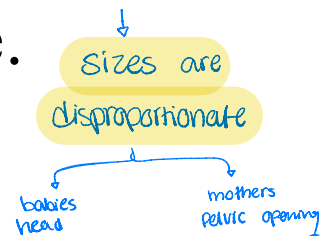
2- Prematurity of fetus

3- Fetal-malpresentation e.g. breech presentation & Cephalopelvic disproportion i.e. contracted pelvis: both predispose to uterine rupture.

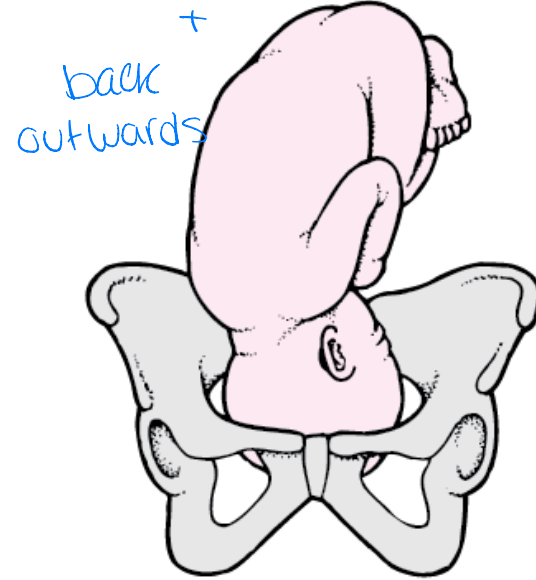
4- Prolapse of umbilical cord following rupture of fetal membranes.

↑ antidiuretic effect
=
water intoxication

→ dilutional hyponatremia
=
death

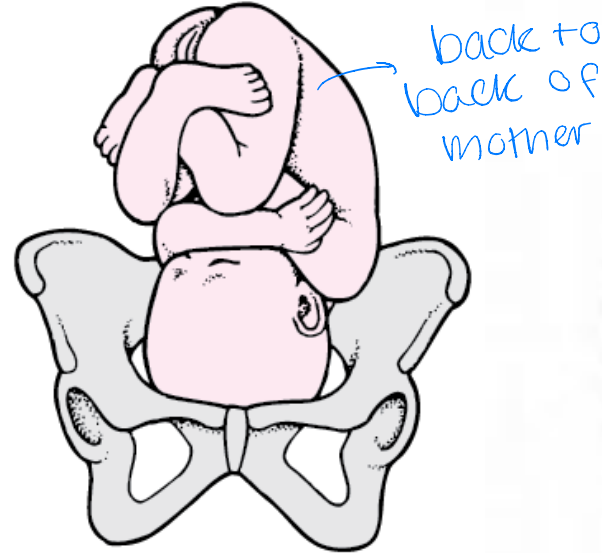


Facing Backward
Head First



Normal Position and
Presentation

Facing Forward



Abnormal
Position
Frank Breech



Complete Breech Footling Breech



Precautions to oxytocin use

not contraindicated
but
high risk

- Multiple pregnancy → 6th, 7th --- weak muscles, risk of rupture
- Previous c- section
- Hypertension [pressor]

2. Ergonovine (Ergometrine) and methylergonovine

- More selective than other ergot alkaloids in stimulating the uterus and is the ergot alkaloid of choice in obstetrics.

Mechanism of action:

1. ~~Powerful~~ **direct** action on uterine muscles
2. Possibly other actions (5-HT₂, alpha₁-adrenoceptor agonist actions).

indirect
action
↑

It helps to prevent postpartum hemorrhage by causing powerful, ~~⊗~~ **sustained** uterine contraction.

NOT appropriate for
labor

intermittent ⚡

Route of administration:

IV or IM at time of delivery of placenta or after delivery of fetus but **never** before.

↑
Sustained
Contractions
Prevent Birth

Side effects: Increased B.P

Contraindications VC risky for:

1. Hypertension as in pre-eclampsia
2. Peripheral vascular disease
3. Angina

3. Prostaglandins (PGs)

E2
F2
⊕ E1
medically

- **Mechanism of action as uterine stimulants:**
- 1- **Direct action:** via receptors
- 2- **Indirect action:** upregulation of oxytocin receptors

❑ **PGE2 (Dinoprostone)**: It is commonly used vaginal.

A. To stimulate uterine contractions for:

Induction of labor given as vaginal gel or insert

Note: If oxytocin is needed for induction of labor, it is given after 6 hours have passed after PG use to avoid excessive uterine contractions. *to avoid rupture*

Induction of abortion: vaginal suppository is used.

B. For softening the cervix at term: This shortens time to onset of labor and labor time.

PGE2 directly affects collagenase of cervix that breaks down the collagen network and softens it.

! oxytocin for dilation when
cervix is soft, what makes
it soft? PGE2

❑ **Gemeprost**: PGE1 analogue

- used as vaginal suppository to induce early medical abortion during **first trimester**.

❑ **Misoprostol**: PGE1 analogue

❑ oral or vaginal supp.:

used for induction of medical abortion in **second trimester** or when gemeprost is not available.

PGF_{2α} (Dinoprost) : less commonly used

May be given vaginally, intra-amniotically, or IV for induction of abortion in second trimester.

Intra-amniotic PGF_{2α} has up to 100% success rate with fewer and less severe adverse effects than IV.

Side effects:

PGF_{2α} causes more G.I. side effects (vomiting, diarrhea) than PGE₂.

Tocolytics

What are tocolytic drugs?

- Drugs that inhibit uterine contractions

عقب ڈاکی حینہ قبل سوئی

Indications:

- Delay, ~~inhibit~~ or ~~prevent~~ premature labor ($< \overset{\rightarrow 39w \text{ full}}{37}$ weeks of pregnancy).

- **Clinical hint:**

- Usually, regular uterine contractions can stabilize on bed rest & local warmth.
When this is insufficient, then a tocolytic drug is used. ← *هكذا ممكن تخفف إذا موكفاج*

N.B. ^{better} **Betamethasone** is given IV to mother or into cord blood to stimulate maturation of fetal lung (by enhancing surfactant formation); it is preferred to **dexamethasone** because it is bound less to plasma proteins.

1. B₂ - adrenoceptor agonists

bronchial asthma
dilatation + relaxation

- Ritodrine, Terbutaline, salbutamol
- **Mechanism of action:** increasing cAMP in myometrium
- **Route of administration:**
- oral or IV infusion in 5% dextrose
- **Side effects:**
 - 1. Tachycardia ↑ dose / long term = loss of B₂ selectivity → affect B₁
 - 2. Sometimes acute left ventricular failure in mother occurs due to overload of infusion fluid and marked tachycardia.
 - 3. Hypokalemia
 - 4. Hyperglycemia

2. Calcium channel blocker *anti-HTN*

- The short acting Nifedipine or Nicardipine
- **Mechanism of action**: inhibition of Ca influx in myometrium
- **Route of administration**: oral
- **Side effects**: hypotension

3- Atosiban

- Analogue of oxytocin that act as competitive antagonist, inhibiting oxytocin binding to its receptor.
- Given IV

4- Miscellaneous drugs → not used clinically

- Progesterone
- Halothane
- Nitroglycerine: NO donor: increasing cGMP in myometrium = relaxation
- Indomethacin: COX inhibitor ↓ PG₁

Contraindications of tocolytics

له مكنوع يمنع الولادة

- 1- Chorioamnionitis: infection of fetal membranes
- 2- Congenital anomalies → abortion probability
- 3- Late pregnancy: more than 34 weeks
- 4- Placenta abruption → hemorrhage + bleeding
- 6- Pre-eclampsia

