

Lecture 2
Drugs inducing ovulation



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introduction:

Causes of Failure to Ovulate (Female infertility)

- 1- Hormonal Problems these are the most common causes of anovulation. The process of ovulation depends upon a complex balance of hormones and their interactions to be successful, and any disruption in this process can hinder ovulation. There are three main sources causing this problem
- 2- Failure to produce mature eggs In approximately 50% of the cases of anovulation, the ovaries do not produce normal follicles in which the eggs can mature. Ovulation is rare if the eggs are immature and the chance of fertilization becomes almost nonexistent.

Note: Polycystic ovary syndrome, the most common disorder responsible for failure to produce mature eggs, includes symptoms such as amenorrhoea, hirsutism, anovulation and infertility. This syndrome is characterized by a reduced production of FSH, and normal or increased levels of LH, estrogen and testosterone.

3- Malfunction of the hypothalamus (Low GnRH)

<u>Hypogonadotropic</u> The hypothalamus is the portion of the brain responsible for sending signals to the pituitary gland, which, in turn, sends hormonal stimuli to the ovaries in the form of FSH and LH to initiate egg maturation. If the hypothalamus fails to trigger and control this process, immature eggs will result.

- 4- Malfunction of the pituitary gland (Low LH & FSH) (Hypogonadotropic) The pituitary's responsibility lies in producing and secreting FSH and LH. The ovaries will be unable to ovulate properly if either too much or too little of these substances is produced. This can occur due to physical injury, a tumor or if there is a chemical imbalance in the pituitary.
- **5- Hyperprolactinaemia:** it is usually caused by Prolactin secreting pituitary tumors. High prolactin levels would inhibit the action of **Gonadotrophs (FSH & LH)**

6-hypergonadotrophic : in this case all we do is conception control , no place for therapy .. usually only surgical option.

Drugs used to induce ovulation

1-ANTIESTROGENS (SERMs)

E.g. Clomiphene & Tamoxifen

MOA:

Selective Estrogen Receptor Modulators [SERMs] → compete with estrogen on estrogen receptors & antagonise the action of estrogen on the hypothalamus & anterior pituitary gland.

A. <u>Clomiphene</u>: compete with the endogenous estrogen receptors in:

hypothalamus; it inhibits the negative feedback of endogenous estrogen on hypothalamus → This will lead to the increased pulsatile release of GnRH → ↑ gonadotrophin production [FSH & LH]

- cause growth maturation & rupture of follicles = OVULATION
- ➤ Ant.pituitary; Aresponse of gonadotrophins to GnRH

Indications

- **➤** Female infertility that is not due to ovarian or pituitary failure **→** Normogonadotrophic
- >The success rate for ovulation →80% & pregnancy →40%.

The discrepancy between 2 rates is due to the antiestrogenic effects of clomiphene on uterus, cervix & vagina

Method of administration:

Clomiphene given → 50 mg/d for 5 days from 5th day of the cycle to the 10th day. If no response give 100 mg for 5 days again from 5th to10th day (double the dose) The drug can be repeated not more than 6 cycles.

ADR:

- 1.Hot Flushes & breast tenderness (Anti-estrogenic effect)
- 2. Fatigue
- 3. Weight gain
- 4. Hair loss (reversible)
- 5. Hyperstimulation of the ovaries & high incidence of multiple birth. (10%)
- 6. Gastric upset (nausea and vomiting)
- 7. Visual disturbances (reversible)
- 8. Nervous tension & depression
- 9. Skin rashes

Note: variety of other symptoms such as nausea and vomiting increased nervous tension, depression, fatigue, breast soreness. However, these appear to result from the hormonal changes associated with an ovulatory menstrual cycle rather than from the medication.

B-TAMOXIFEN

Is similar & alternative to clomiphene ,But <u>differ in being Non-Steroidal</u>
Used in palliative treatment of hormone-dependent / estrogen receptor- positive advanced breast cancer

But why not clomiphene used in such cases of breast cancer?
Tamoxifen has an antiestrogenic effects in the breast while clomiphene doesn't; it is

Tamoxifen has an antiestrogenic effects in the breast while clomiphene doesn't; it is actually has a weak estrogen agonist effect in the breast.

Note: Tamoxifen has a potent antiestrogenic effects in the breasts (But maybe estrogen agonist in the uterus). That is why it must be used with caution while treating a female with breast cancer and has a history of endometrial hyperplasia.

2. GONADOTROPIN RELEASING HORMONE (GnRH)

Analgoues with agonist activity:

Leuprolin, Goserelin

Mechanism:

Native GnRH is naturally produced by hypothalamus in a pulsatile manner. It is triggered when the negative feedback inhibition of ovarian hormones is lost by the end of the cycle. This activates FSH release from pituitary that stimulates growth and maturation of ova early during the follicular phase of the cycle. It also mediates estrogen induced LH surge that triggers ovulation.

GnRH-Agonists → bind to the receptors & mimic the native hormones provided it is given pulsatile.

GnRH and agonists, given S.C. in a pulsatile(drip) to stimulate gonadotropin release $(1-10 \ \mu g \ / \ 60-120 \ min)$ Start from day 2-3 of cycle up to day 10. Given continuously, when gonadal suppression is desirable e.g. precocious puberty and advanced breast cancer in women and prostatic cancer in men.

Uses:

- 1-Induction of ovulation in patients with hypothalmic amenorrhea (GnRH deficient)
- 2-In ASSISTED REPRODUCTION is part of a protocol for OVUM RETRIEVAL (Induction of ovulation)

Prof. Alhumyed said: there're some studies about using (continuous GnRH agonist) as a contraceptive, because it is going to inhibit the ovulation when it's given continuously.

ADR:

GIT disturbances, abdominal pain, nausea....etc

- > Headache
- ➤ Hypoestrogenism on long term use →
- -Hot flushes
- -Decreased Libido
- -Osteoporosis
- ➤ Rarely ovarian hyperstimulation → (ovaries swell & enlarge)

3.GONADOTROPHINS (FSH & LH)

Are naturally produced by the pituitary gland

For therapeutic use, extracted forms are available as;

- 1. Human Menopausal Gonadotrophin(hMG) → extracted from postmenopausal urine → contains LH & FSH → MENOTROPIN
- 2. Human Chorionic Gonadotrophin(hCG) extracted from urine of pregnant women → contains mainly LH) → PREGNYL
- N.B. Now, new available preparations by recombinant technology.

Mechanism:

Given sequentially

- ➤ Preparations of FSH → act on ovary directly, stimulating growth & maturation of Graafian Follicle(s)
- ➤ Preparations of LH → act just to induce ovulation

Uses:

> Stimulation & induction of ovulation in infertility 2ndry to gonadotropin deficiency (pituitary insufficiency)

Success rate for inducing ovulation is usually >75 %.

Method of administration:

hMG is given i.m or subcutaneous every day starting at day 2-3 of cycle for 10 days (to induce the development of the follicle) followed by

hCG on (10th - 12th day) for OVUM RETRIEVAL (Ovulation or rupture of the follicle) within 36 hours when we indicate: intrauterine insemination or intercourse.

ADR:

FSH containing preparations (MENOTROPIN);

- ✓ Fever
- ✓ Ovarian enlargement (hyper stimulation)
- ✓ Multiple Pregnancy (approx. 20%)

LH containing preparations; Headache & edema

4-Dopamine (2) Receptor Agonists (BROMOCREPTINE)

Is an ergot derivative (not a hormone)

MOA:

D₂ R Agonists binds to dopamine receptors in the anterior pituitary gland & inhibits prolactin secretion .

Uses: Female infertility 2_{ndry} to hyperprolactinaemia (hypogonadotrophic)

ADR:

- > GIT disturbances; nausea, vomiting, constipation
- > Headache dizziness & orthostatic hypotension
- > Dry mouth & nasal congestion
- Insomnia

Note: Orthostatic hypotension is due to stimulating D2 receptors, which are found in peripheral mesenteric and renal vascular beds, that causes vasodilatation.

Summary

✓ The main Cause of Failure to Ovulate (Female infertility) is hormonal imbalance e.g high estrogen, low gonadotrophs(FSH & LH) due to pituitary or hypothalamic disorders, or high prolactin levels usually due to prolactinoma.

✓ 1-Selective Estrogen Receptor Modulators [SERMs] E.g. Clomiphene & Tamoxifen

- ✓ Clomiphene is a partial agonist at estrogen receptors.
- ✓ **Clomiphene:** compete with the **endogenous estrogen receptors** in hypothalamus & ant.pituitary ,thus removing the negative feedback effect of estrogens from the hypothalamus & Ant.pituitary leading to an increase in gonadotrophs' (FSH & LH) release.
- ✓ Clmoiphene is used in **Female infertility that is not due to ovarian or pituitary** failure → Normogonadotrophic

- ✓ Clomiphene given \Rightarrow 50 mg/d for 5 days from 5th day of the cycle to the 10th day. (we repeat this method of dosing for 3 cycles)
- \checkmark If no response for 3 cycles: the dose is doubled in the next 3 cycles with the same method of administration.
- \checkmark The drug can be repeated not more than 6 cycles.
- \checkmark Its main ADRs : Visual disturbances, hair loss which can be reversible , .Hot Flushes & breast tenderness that are due to the Antiestrogenic effect of Clomiphene
- \checkmark Tomoxifin Is similar & alternative to clomiphene ,But differ in being Non-Steroidal
- ✓ Tamoxifin has an **Antiestrogenic effects on the breast** (Used in cases of breast cancer)
- ✓ 2-Leuprolin, Goserelin are synthetic analogues of GnRH with agonistic activity
- ✓ given S.C. in a pulsatile(drip) to stimulate gonadotropin release
- ✓ Starts from day 2-3 of cycle up to day 10.
- ✓ Given continuously, when gonadal suppression is desirable e.g. precocious puberty and advanced breast cancer in women and prostatic cancer in men
- ✓ It's main ADR is Hypoestrogenism on long term use → Hot flashes, decreased Libido ,Osteoporosis

√ 3-Gonadotrophs

- ✓ 1. Human Menopausal Gonadotrophin (hMG) → extracted from postmenopausal urine → contains LH & FSH → MENOTROPIN
- ✓ 2. Human Chorionic Gonadotrophin (hCG) extracted from urine of pregnant women → contains mainly LH) → PREGNYL
- ✓ They are used in Stimulation & induction of ovulation in infertility 2_{ndry} to gonadotropin deficiency (pituitary insufficiency)
- √ hMG (MENOTROPIN)is given i.m every day starting at day 2-3 of cycle for 10 days
 (to induce the development of the follicle) followed by
- \checkmark hCG (PREGNYL) is given on (10th 12th day) for OVUM RETRIEVAL (Ovulation or rupture of the follicle) .
- ✓ ADRs:

FSH containing preparations (**MENOTROPIN**);

- ✓ Fever
- ✓ Ovarian enlargement (hyper stimulation)
- ✓ Multiple Pregnancy (approx. 20%)

LH containing preparations (MENOTROPIN or Pregnyl); Headache & edema

- ✓ **MENOTROPIN**, Leuprolin, and Goserelin may lead to ovarian swelling or enlagement
- ✓ 4-BROMOCREPTINE: D₂ R Agonists binds to dopamine receptors in the anterior pituitary gland & inhibits prolactin secretion.
- ✓ Used in Female infertility 2ndry to hyperprolactinaemia
- **✓ 5-METFORMIN** is used in IN POLYCYSTIC OVARIAN SYNDROME to
- **♦** body weight &↑ response to ovulation induction drugs

Questions:

- 1- Drugs useful in the treatment of infertility include all of the following EXCEPT:
- a. Human chorionic gonadotropin.
- b. Bromocreptine.
- c. Gonadotropin-releasing hormone.
- d. Prolactin.
- 2- Bromocreptine causes the following:
- a. Prolactin release.
- b. Vomiting.
- c. Uterine contraction.
- d. Impotence.
- 3- The most prominent action of bromocriptine is:
- a. Dopamine D2 agonist.
- b. Dopamine D2 antagonist.
- c. Dopamine D1 antagonist.
- d. Alpha adrenergic antagonist.

Answers:

- 1- D
- 2- B
- 3- A