

FEMALE REPRODUCTION SYSTEM

OÖGENESIS

DEVELOPMENT:

6-8 weeks

GERMINAL EPITH.

hormonally
independent

OÖGONIA
mitotic division

FOLLICLE
PRIMORDIAL

24 weeks

OÖCYTES I.

7×10^6

1. meiosis
prophase

2×10^6

birth

hormonally
dependent
(cyclic)

puberty

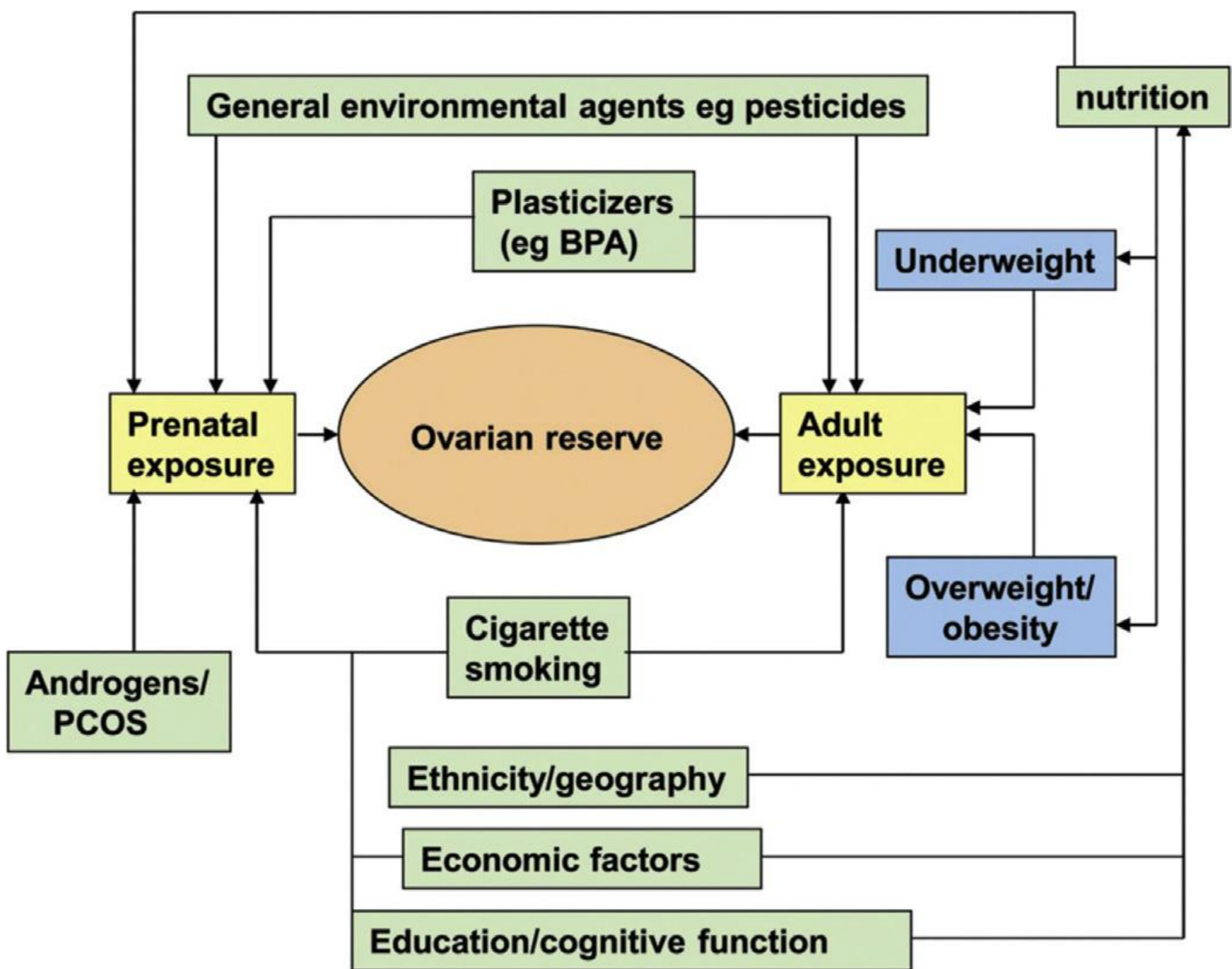
OÖCYTES II.
haploid
2. meiosis
metaphase
OVUM

3×10^5
DOMINANT
ATRETIC
GRAAF
OVULATION

2. meiosis – end

climacterical

0



UTERINE CYCLE

ovarian

uterine

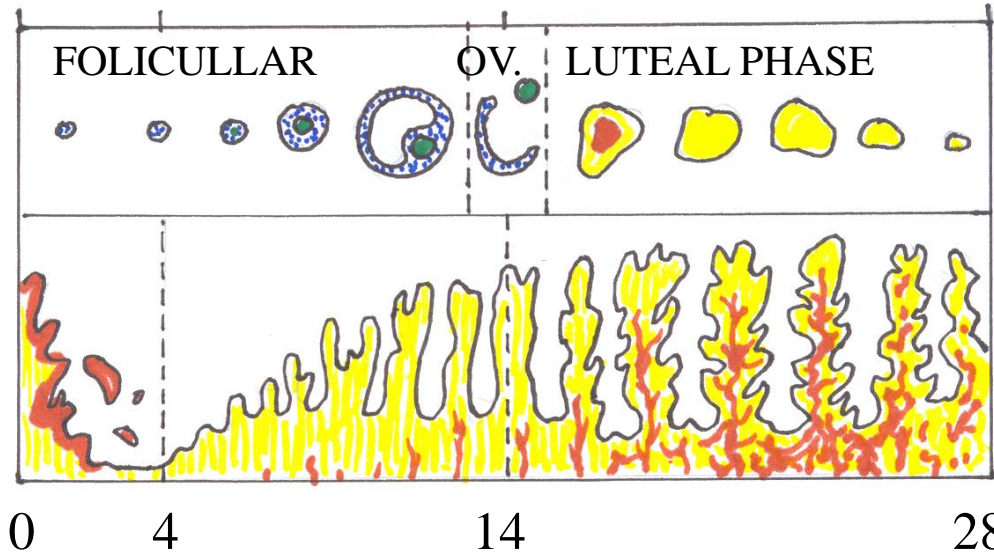
gonadoliberin
(GnRH)

FSH, LH

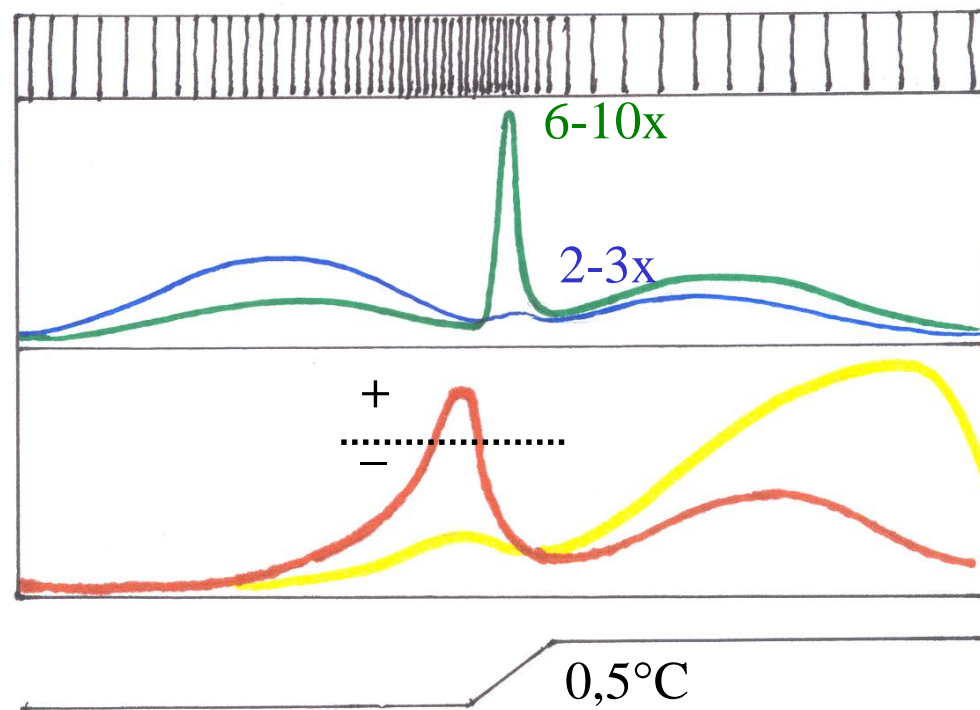
estradiol

progesteron

basal temper.



0 4 14 28
MENS. PROLIPHER. SECRETORY PHASE



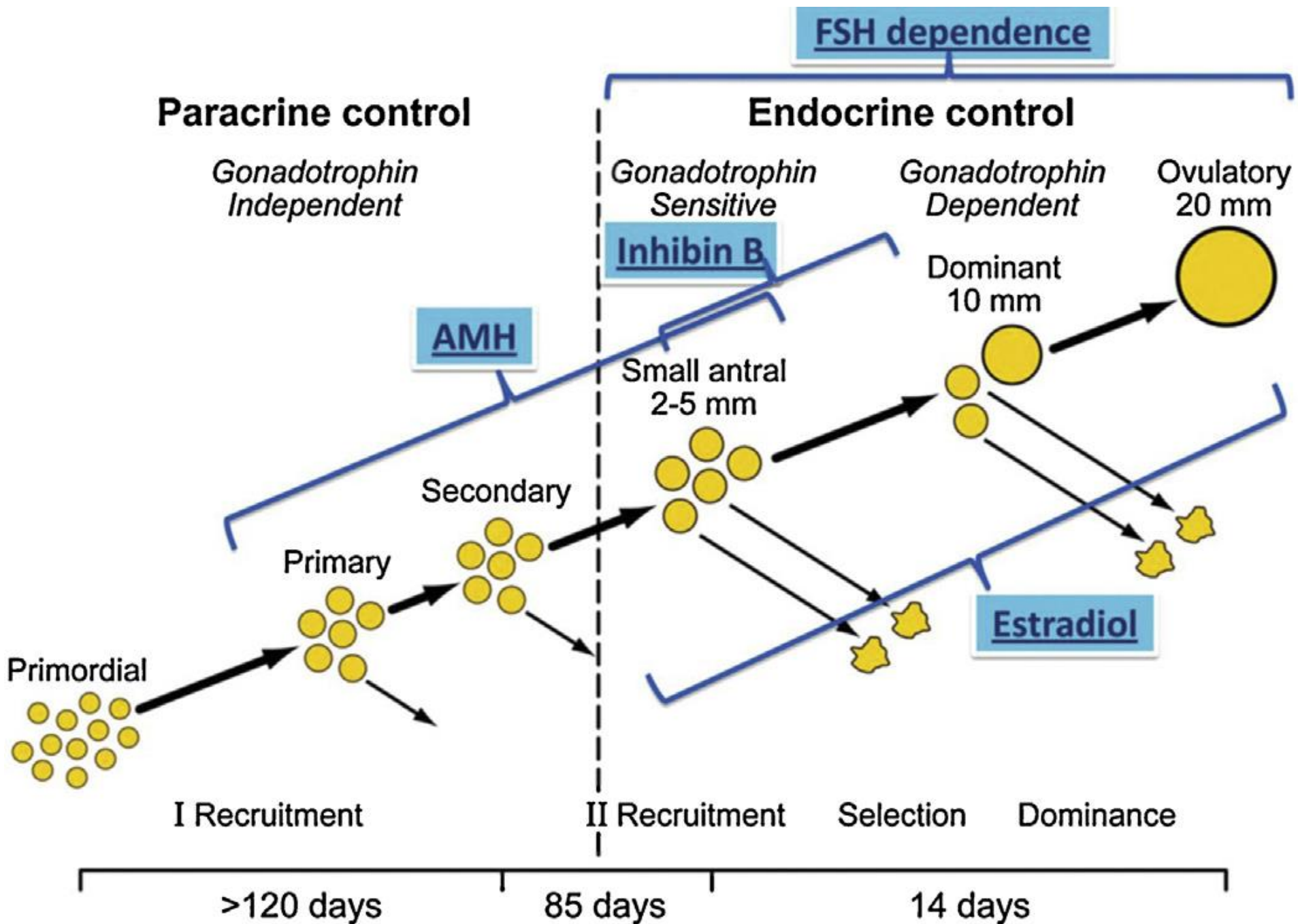
6-10x

2-3x

+

-

0,5°C

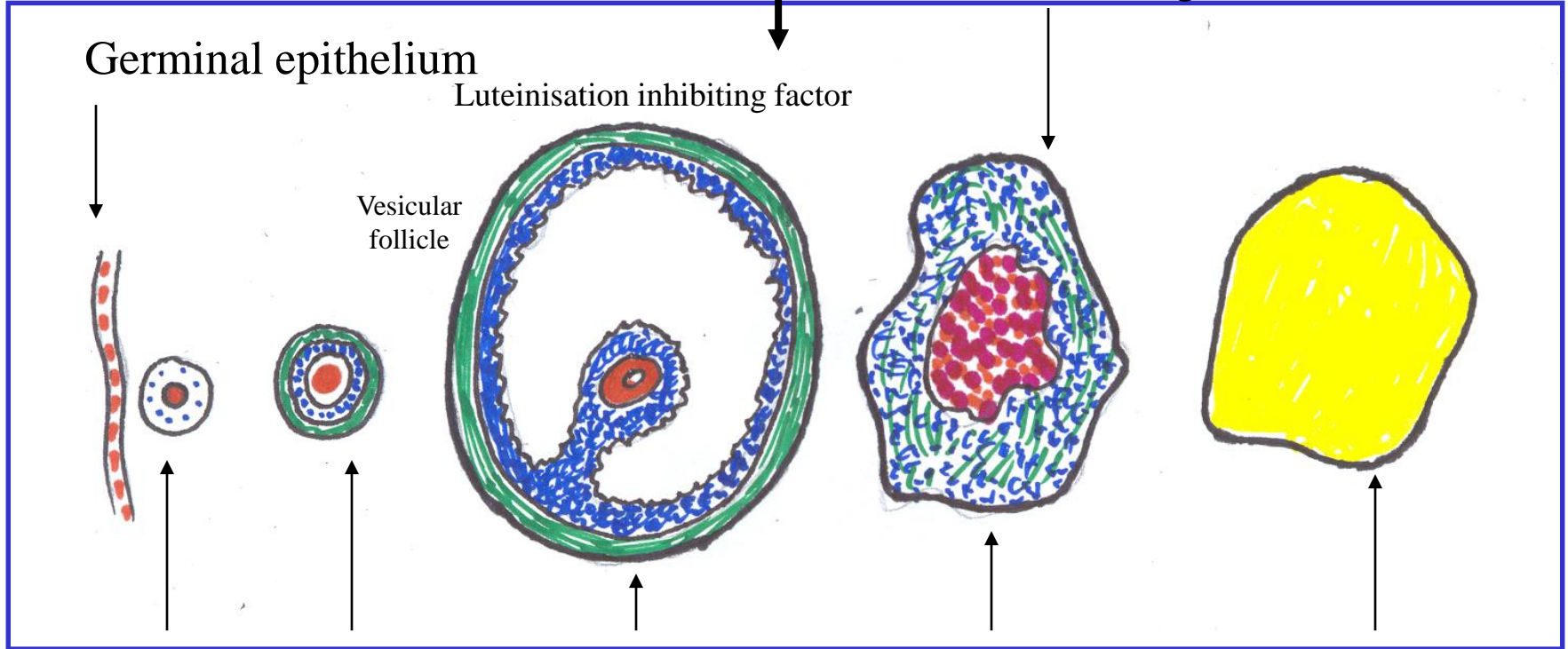


OVARIAN CYCLE

Oocyte-maturation inhibiting factor

OVULATION

methrorrhagia



Primordial

Primary
follicle

Graaf

Corpus haemorrhagicum C. luteum

25 μ

150 μ

up to 2 cm

estradiol (estrogens)

progesteron
(progestins)

VESICULAR FOLLICLE

PRIMARY FOLLICLE - FSH

Growth acceleration of primary follicle – change into vesicular follicle:

1) estrogens released into follicle stimulate granul. cells



increased number of receptors for FSH – POSITIVE FEEDBACK

(higher sensitivity for FSH!!!)

2) Increased number of receptors for LH (estrogens and FSH) – another acceleration of growth due to „higher sensitivity“ to LH

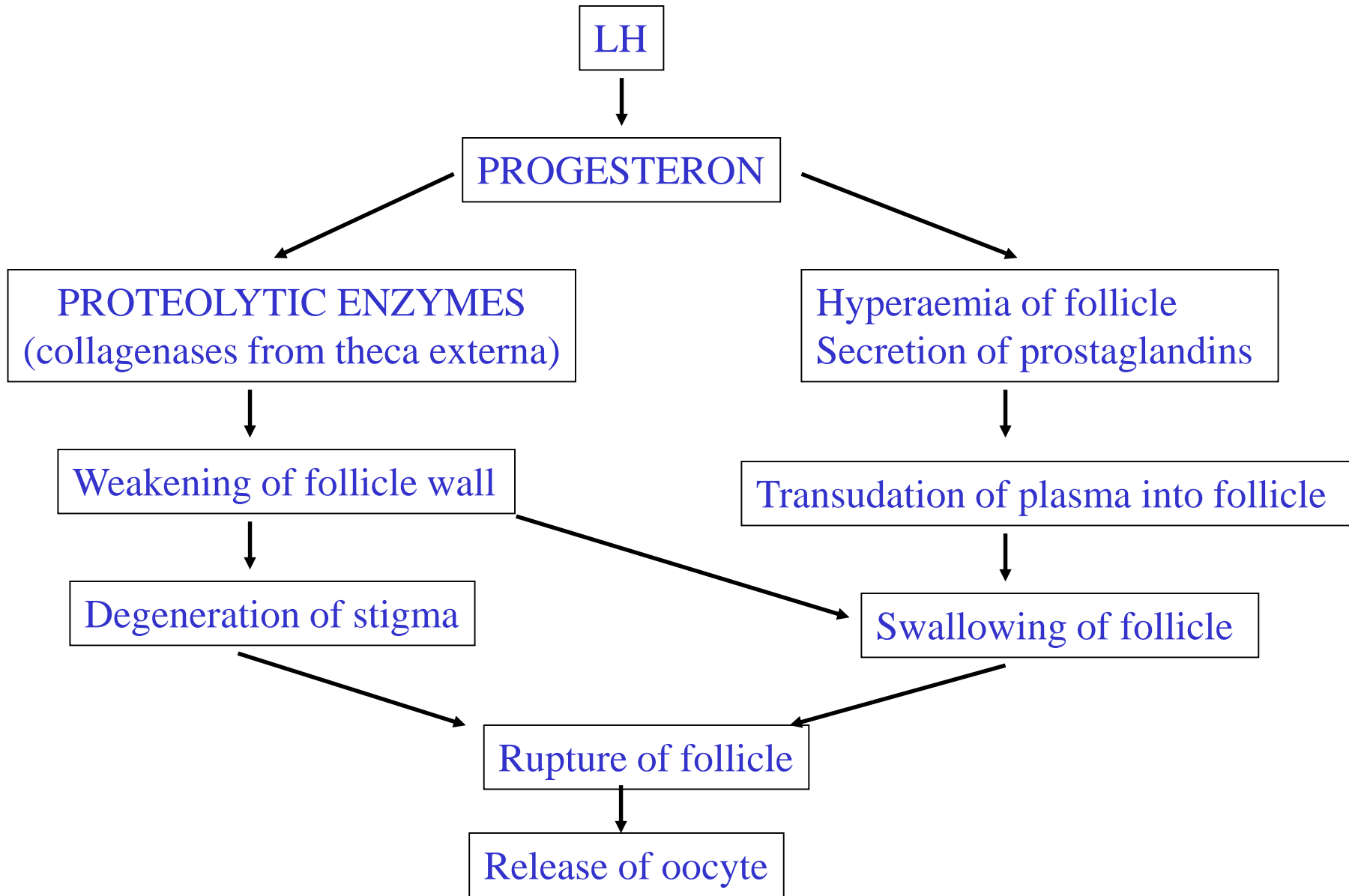
3) Increased estrogens and LH secretion accelerates growth of theca cells, secretion is increased

→ **explosive growth of follicle**

DOMINANT FOLLICLE

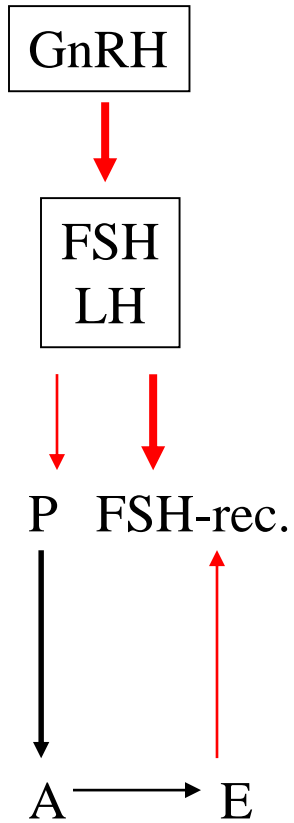
1. High level of **estrogens** from the fastest-growing follicle
2. **Negative** feedback on FSH production from adenohypophysis
3. Drop in **FSH** secretion
4. „**Dominant** follicle“ continues in growing due to **intrinsic positive** feedback
5. Other follicles grow slowly and subsequently become **atretic**

MECHANISM OF OVULATION

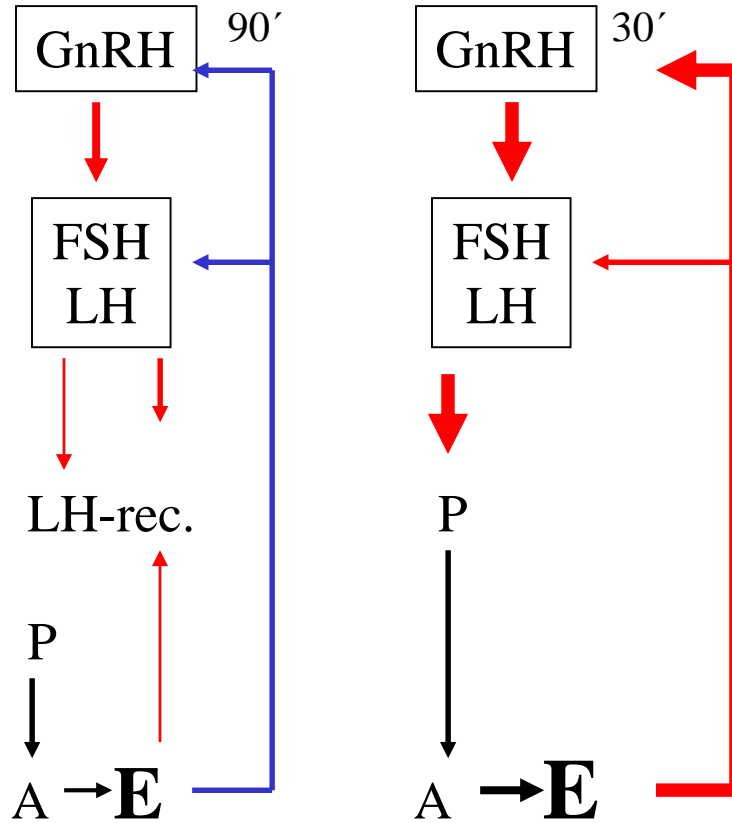


HUMOURAL REGULATION OF CYCLE

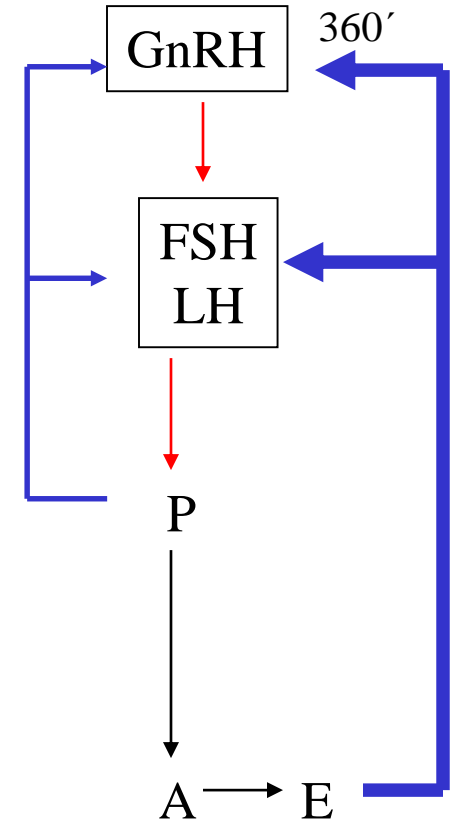
Follicular phase



Ovulation



Luteal phase



Artesia of follicle (except of one)

Feedback -/+

Involution of corpus luteum

EFFECTS OF OVARIAL HORMONES

E

P

Ovaries:	maturation of follicles	
Hysterosalpinx:	motility	motility
Uterus:	proteosynthesis	proteosynthesis
	vascularisation and proliferation of endom.	secretion of endom. glands
	motility	glycogen
		motility
Cervix:	colliquation of „plug“	creation of „plug“
Vagina:	cornification of epithelium	proliferation of epithelium
Mamma:	growth of terminals	growth of acines

Secondary sexual signs	+	-
Adipose tissue:	store (predilection), (critical amount)	-
Bone tissue:	absorption	-
	closure of fissures	-
	development of pelvis	-
Total water retention:	+	+
Sexual behaviour:	+	-

ASSISTED REPRODUCTION TECHNIQUES

1. STIMULATION OF OOGENESIS (maturation of more follicles)
2. STIMULATION OF SPERMIOGENESIS (vit. E)
3. INSEMINATION (treated sperm, applied deeply into uterus)
4. IVF (in vitro fertilisation)

IVF PROCEDURES

1. STIMULATION OF OVARIES
2. TIMING OF TAKING THE OOCYTES
3. EXTRACORPOREAL FERTILISATION OF OOCYTES
4. EMBRYOTRANSFER AND MAINTAINANCE THERAPY

Ad 1) **PROTOCOLS OF OVARIAL STIMULATION** (short of long stimulation protocols)

Stimulation of ovaries – **FSH** and **LH**, 3. - 12. day of cycle, SOMETIMES combined with **GnRH** agonists or antagonists

Ad 2) **TIMING OF TAKING THE OOCYTES**

Between 12. and 17. days of cycle, US controlled, after stimulation of oocyte maturation by hCG, aspiration from follicular liquid in analgesia or anaesthesia

Ad 3) **EXTRACORPOREAL FERTILISATION OF OOCYTES** (cultivation of sperm and oocytes in vitro for 48 hrs; test of sperm surviving – min.40%; micromanipulation techniques – ICSI a AH = gentle rupture of zona pellucida; prolonged cultivation – up to 120 hrs)

Ad) **EMBRYOTRANSFER** (transfer of max. 3 embryos in stage of morula or blastula; genetic examinations) and **MAINTENANCE THERAPY** (progesterone)

CONTRACEPTION (BIRTH CONTROL)

- RHYTHM METHOD
- SPERMICIDE SUBSTANCES
- COITUS INTERRUPTUS
- CONDOM, PESSARY
- IUD
- HORMONAL CONTRACEPTIVES – risk of failure less than 1%
- VASECTOMY AND LIGATION OF HYSTEOSALPINX

Hormonal curettage (excochleation). Substitution therapy in climacterium.

HORMONAL CONTRACEPTION

- block of ovulation by suppression of hypothalamic releasing hormones
(block of preovulatory surge of LH)
- changes of character of cervical plug (progestin thickens mucus)
- changes of endometrium (suppression of its growth)
- changes of hysterosalpinx motility

Combined hormonal contraceptives:

- **monophasic** (amount of oestrogen and gestagen is stable)
- **biphasic** and **triphasic**
- **combiphasic** contraceptives (after 7 days gestagen content increases and oestrogen content decreases)

15 μ g estrogenu
60 μ g progestinu