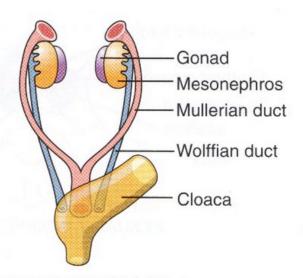
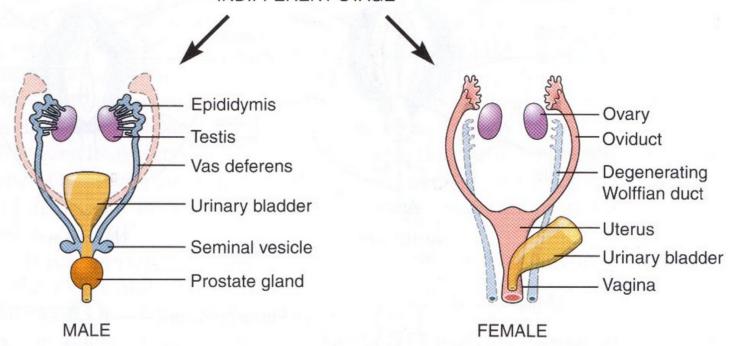
# Development and teratology of reproductive system

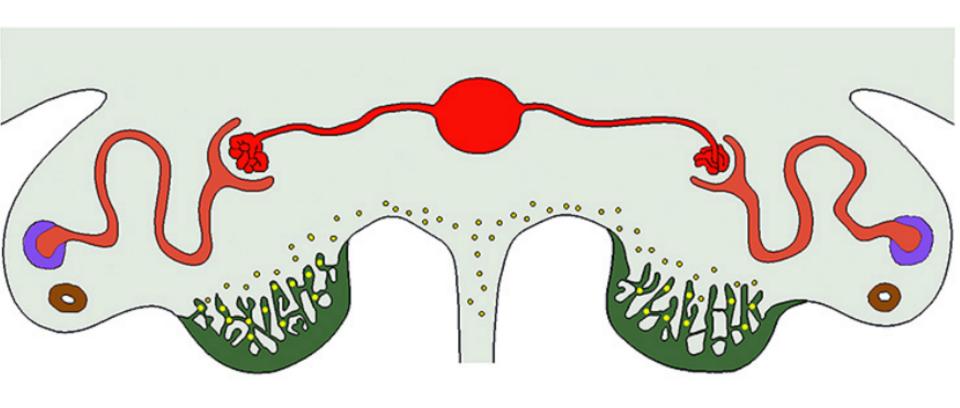
Anna Mac Gillavry 24.04.2023



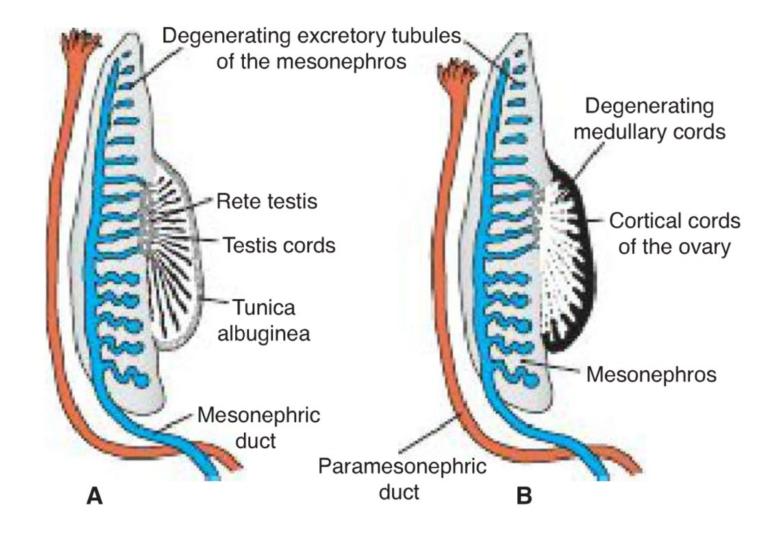
#### INDIFFERENT STAGE



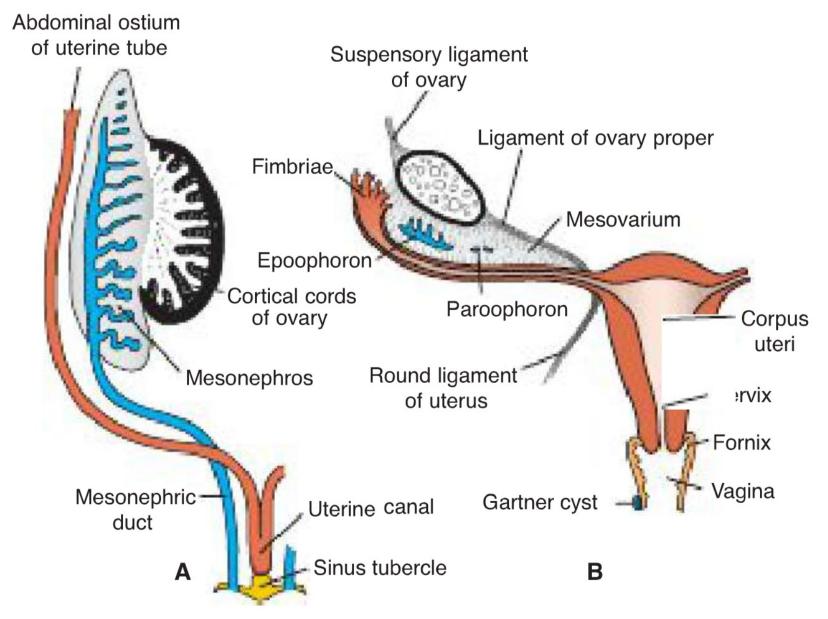
# Indifferent stage of the gonads



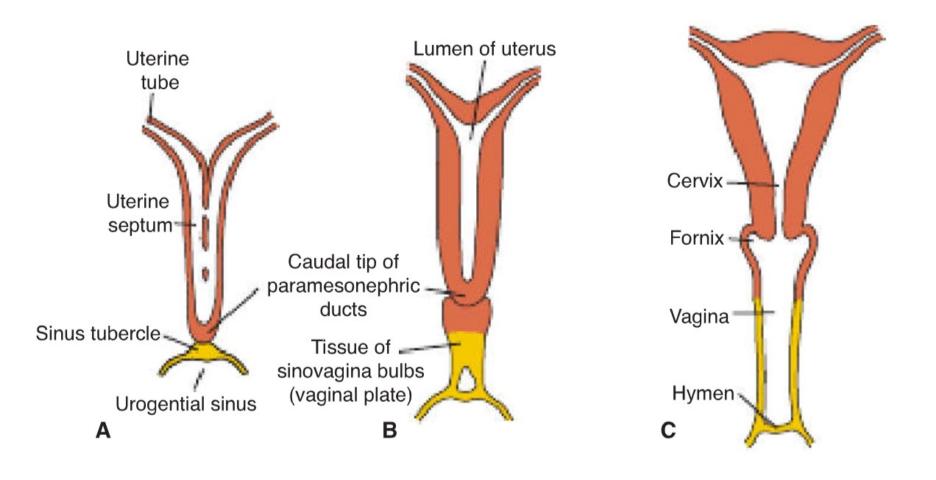
# Indifferent stage of the genital ducts



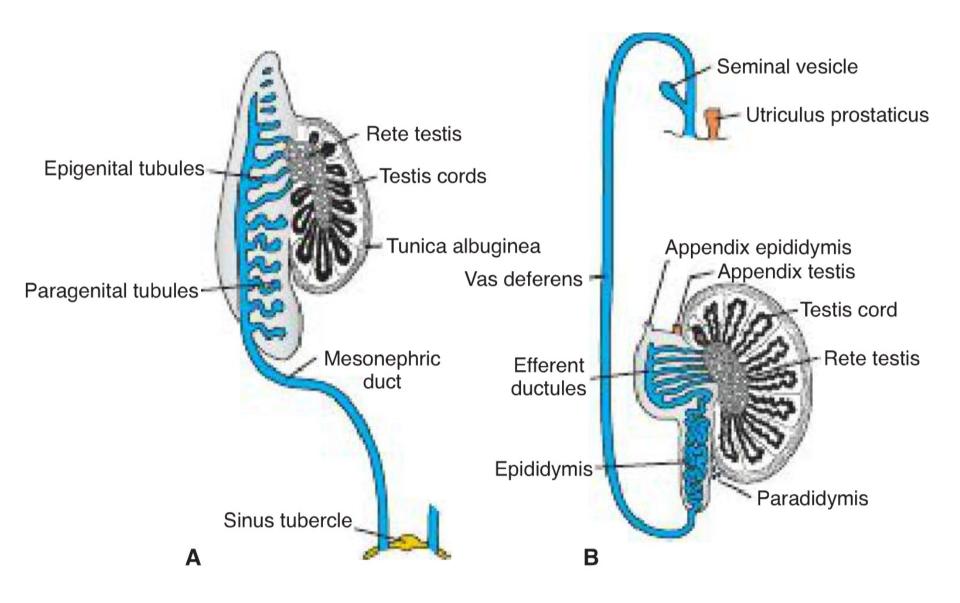
### Differentiation of the ducts - females



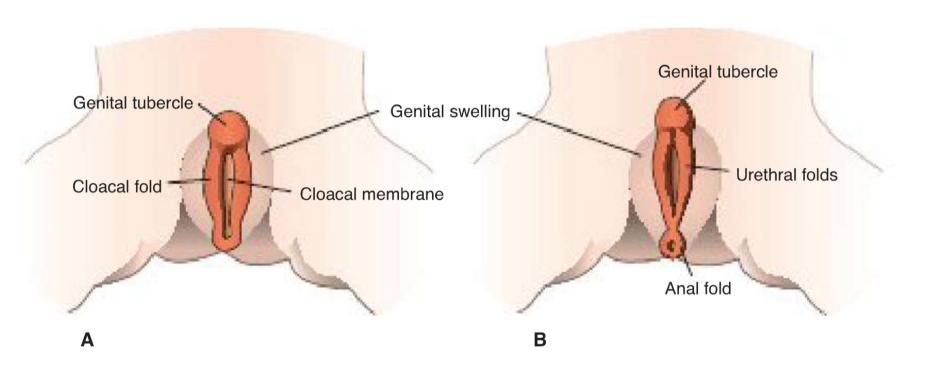
# Development of the vagina



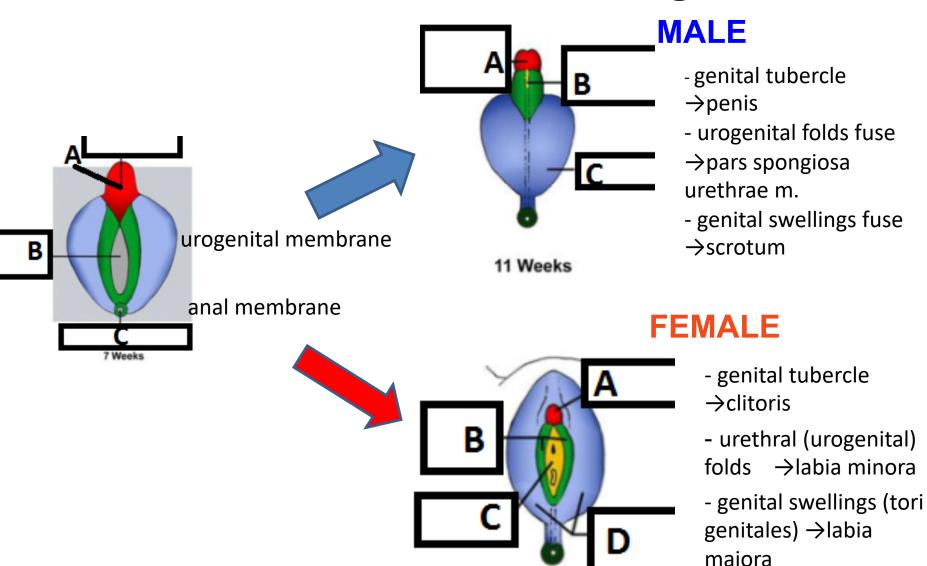
### Differentiation of the ducts - males



# Indifferent stage of the external genitalia



# Differentiation of external genitalia



12 Weeks

# Disorders of sexual development (DSDs)

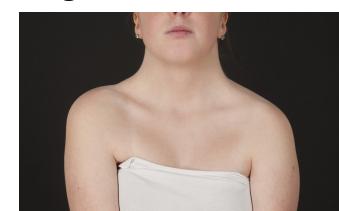
- Ambigous genitalia: clitoral hypertrophy/small penis with hypospadia
- 46, XX DSDs:
- congenital adrenal hyperplasia, most common cause – 60 % of all DSDs
- 46, XY DSDs:
- androgen insensitivity syndrom (AIS): complete (CAIS), mild (MAIS) or partial (PAIS)
- 5-a-reductase deficiency: testosteron convertion into dihydrotestosteron is impaired

Ovotesticular DSDs - the caryotype is 46, XX in 70 % of cases

#### Chromosomal abberations

Klinefelter syndrom – 47, XXY (XXXY...) – 1 in 1000 males: decreased fertility, small testes, decreased testosteron levels, gynecomastia in app. 33%

- Gonadal dysgenesis oocytes are absent:
- Swyer syndrome XY female gonadal dysgenesis –
  point mutation or deletions of the SRY gene
- Turner syndrome 45, X

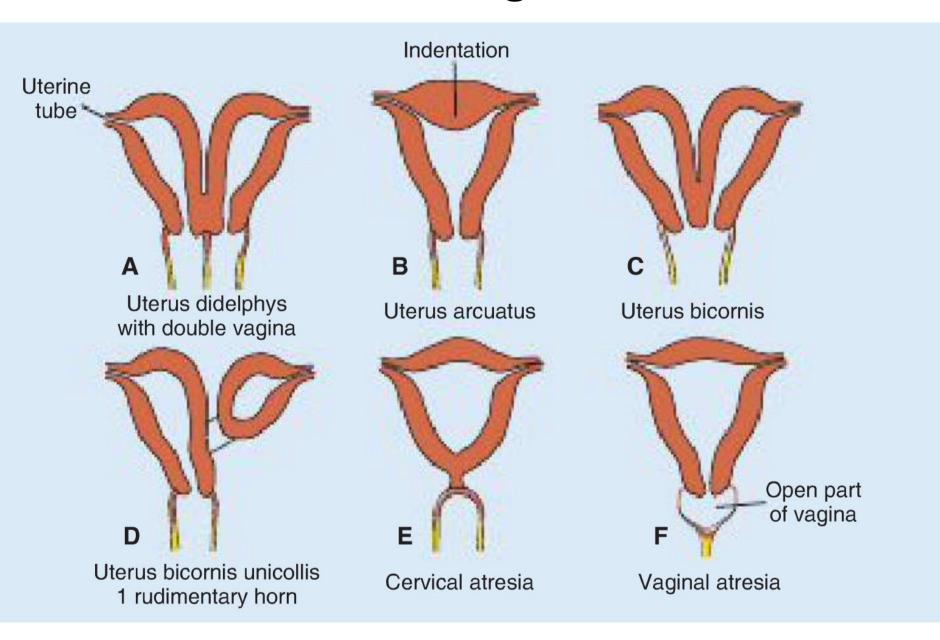


# Hernias and cryptorchidism

- Congenital inderect inguinal hernia
- Hydrocele of the testis and/or spermatic cord

Cryptorchidism

# Uterine and vaginal defects



# Defects in male genitalia

- Hypospadia 3 to 5 in 1000 births
- Epispadia 1 in 30000 most often associated with extrophy of the bladder and abnormal closure of the ventral body wall
- Micropenis insufficient androgen stimulation
  primary hypogonadism, hypothalamic or
  - pituitary disfunction
- Bifid penis