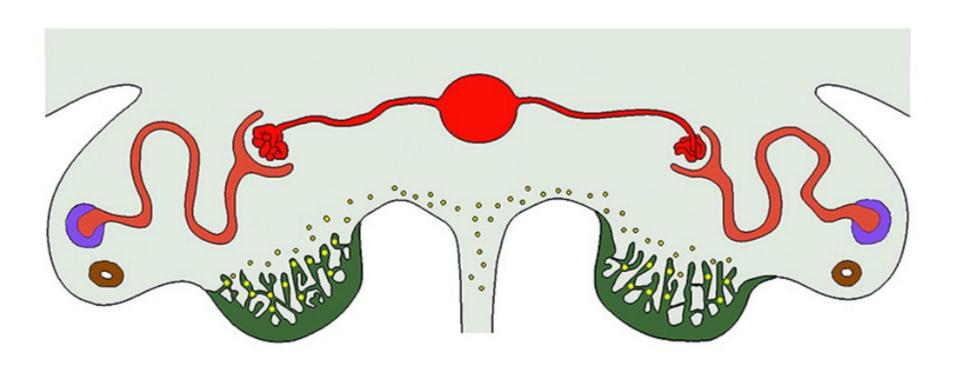
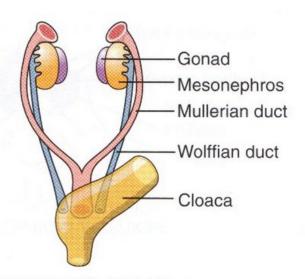
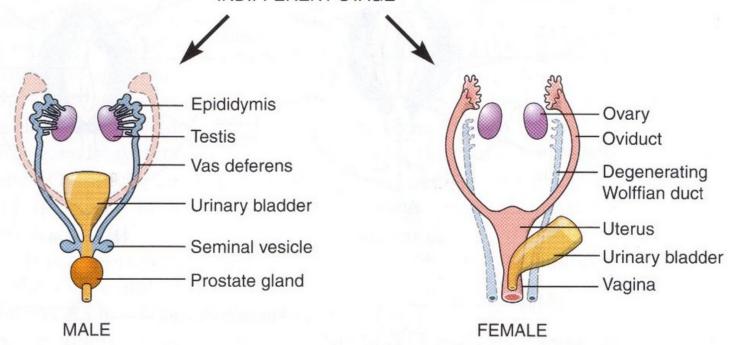
Development and teratology of reproductive system.

Anna Mac Gillavry 05.04.2021



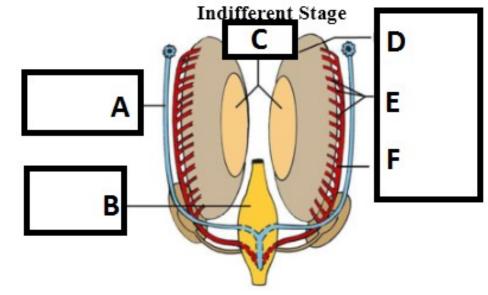


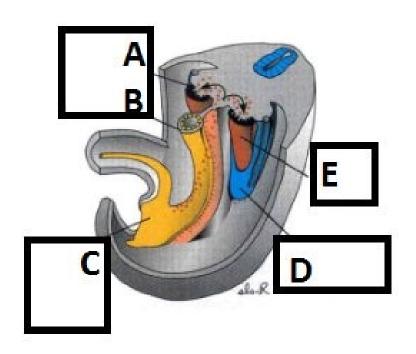
INDIFFERENT STAGE



Development of gonad

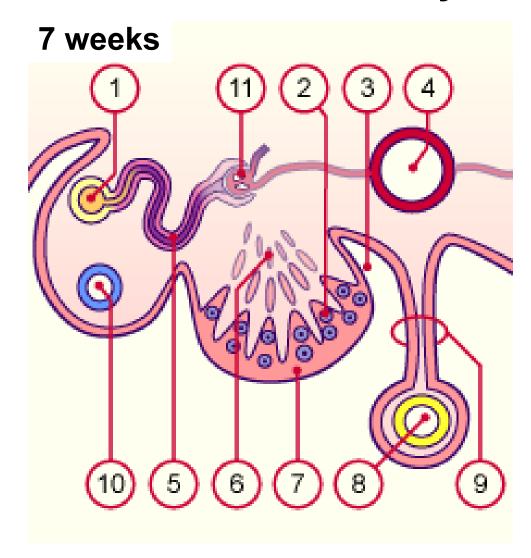
- indifferent gonad to 6th-7th week of development
 - germ cell (wall of yolk sac)
 - <u>genital (gonadal)</u><u>ridge</u> mesoderm
 - coelomic epitheliummesodermprimitive sex cords





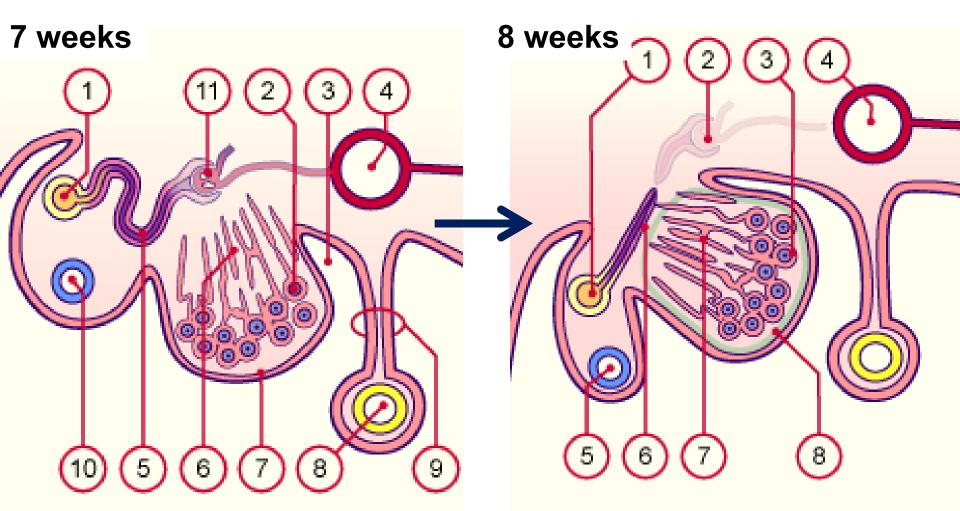
Development of gonad - differentiation to ovary

- 2 generations of cords
 - medullary disappers
 - 7th week cortical cords arise →form follicles, remain close to the surface
- germ cells– oogonia

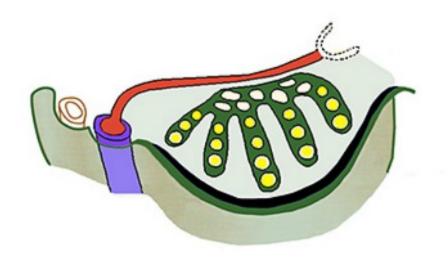


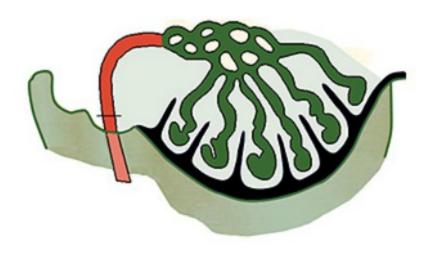
Development of gonad - differentiation to testis

- 1 generation of genital cords, separating from the coelomic epithelium



http://www.embryology.ch/anglais/ugenital/diffmorpho02.html

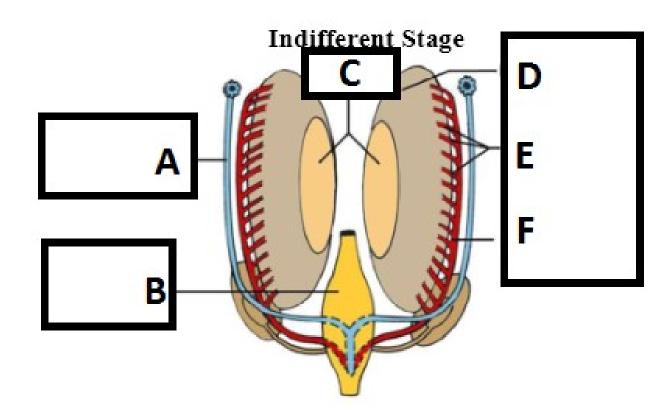




Development of genital ducts

indifferent stage

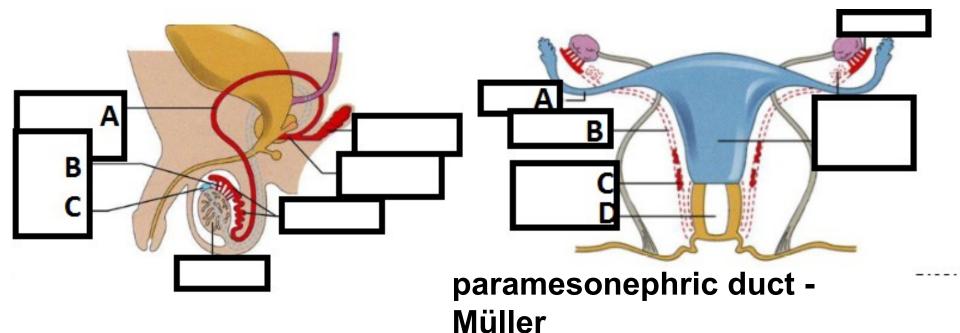
- mesonephric duct (Wolffian)
- paramesonephric duct (Müllerian) longitudinal invagination of coelomic epithelium at the urogenital cord



Differentiation

towards male sex

towards female sex



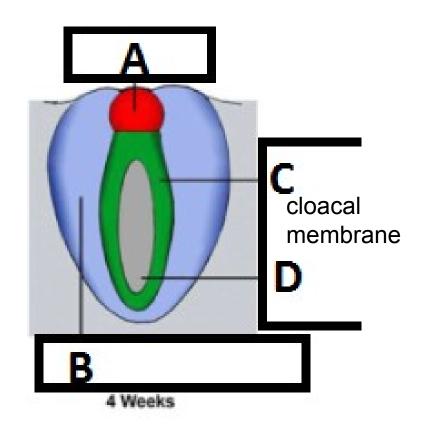
mesonephric duct – Wolff

- vertical
- horizontal do not fuse →uterine tube
- vertical both ducts fuse → uterovaginal primordium uterus and cranial part of vagina

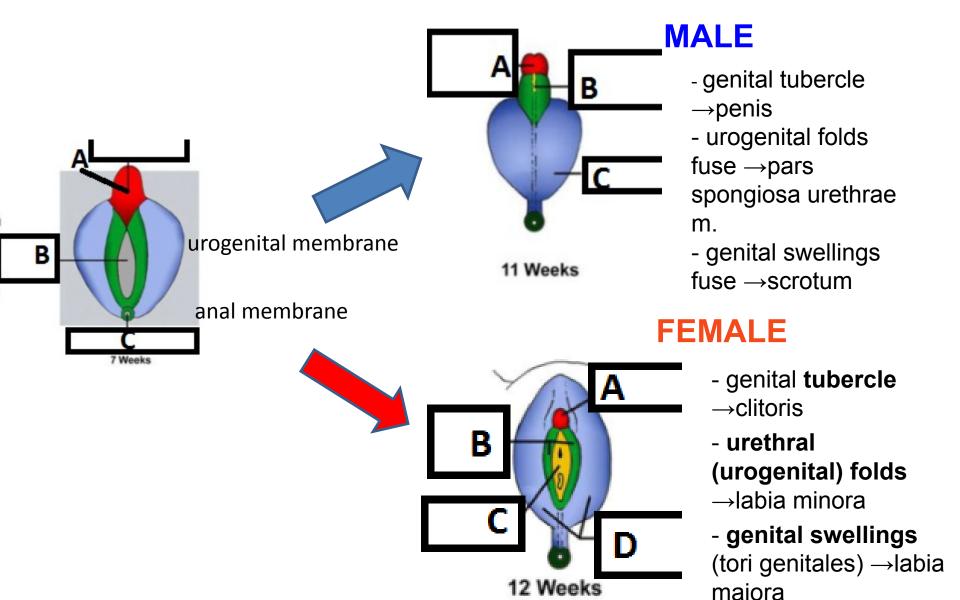
Development of external genitalia

indifferent stage

- cloacal **folds** around the cloacal membrane
 ⇒anteriorly **urethral** (urogenital) folds
 - →posteriorly anal folds
- genital tubercle (phallus)
- genital swellings



Development of external genitalia - differentiation



Disorders of sexual development "hermaphrodites"

- Ambigous genitalia: clitoral hypertrophy/small penis with hypospadia
- Ovotestes less than 1 in 20000 in 70 % of cases, the caryotype is 46, XX
- Genotype does not match the phenotype
- congenital adrenal hyperplasia, most common cause;
- androgen insensitivity syndrom (AIS): complete (CAIS), mild (MAIS) or partial (PAIS)
- 5-a-reductase deficiency affects males

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

- Duplication of the uterus
- uterus didelphys
- uterus bicornus
- uterus arcuatus
- Uterus bicornus unicollis
- Cervical atresia
- Vaginal atresia

Defects in male genetalia

- 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