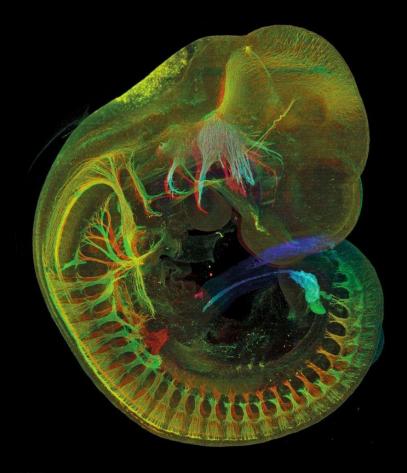


## Introduction to embryology III

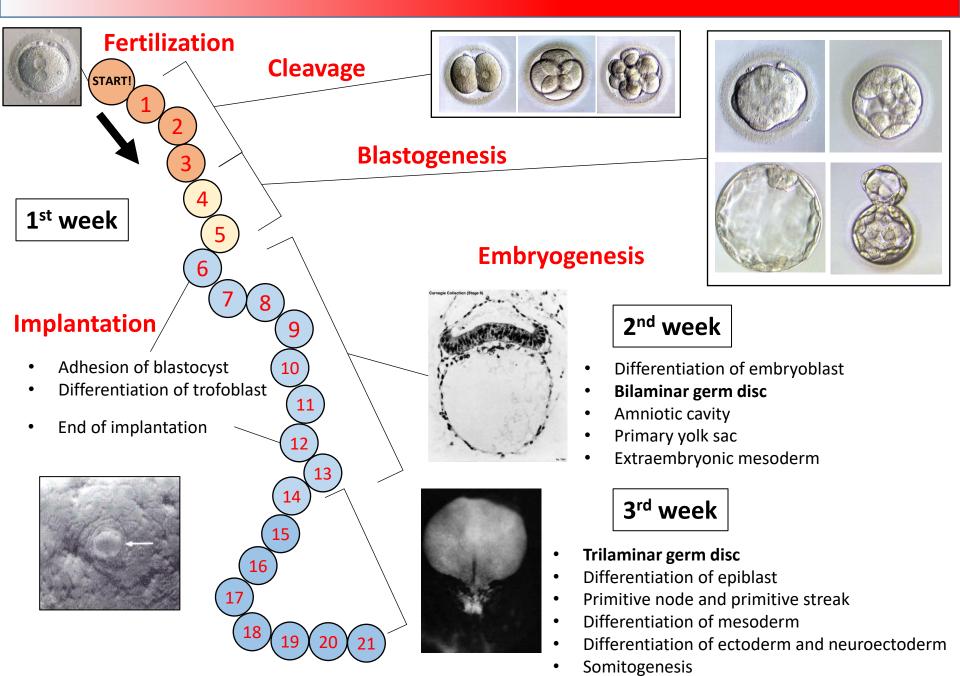
Petr Vaňhara



- 1. Revision of early embryogenesis
- 2. Trilaminar germ disc embryo
- 3. Folding of embryo
- 4. Pregnancy and fetal development
- 5. Prenatal diagnostics

@ RPS/BNPS

#### FIRST EVENTS IN HUMAN LIFE

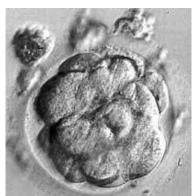


# 1<sup>st</sup> week **Mechanism of diferentiation** "cell polarity" "inside-outside"







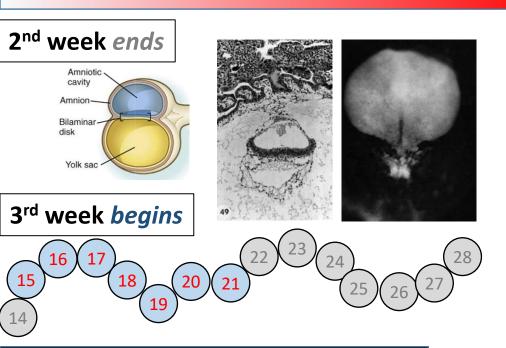


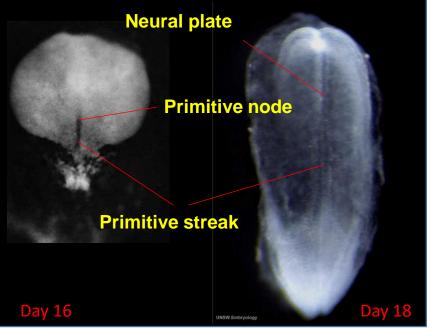


16-cell embryo is still totipotent – later (32-cell), it loses the full developmental potential
 → cell are determined.

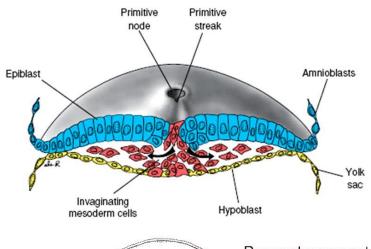
mesoderm

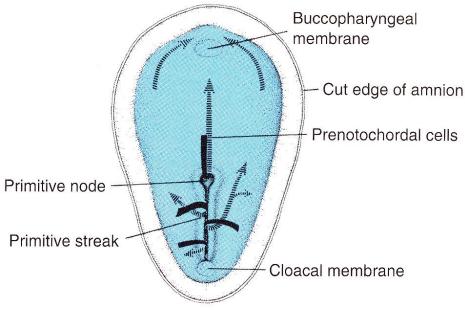
#### 1st-2nd week **BILAMINAR GERM DISC** Amniotic cavity Blastocoele Day 5 Day 7 1/2 Amnion Imphoblast amniotic cavity Open roof of Day 6 Cytotrophoblast amniotic cavity Bilaminar disk Hypoblast Extraembryonic mesoderm В Yolk sac Parietal Day 8 endoderm Amniotic **Epiblast** Secondary membrane Splanchnic mesoderm Hypoblast of yolk sac Extraembryonic coelom Extraembryonic mesoderm Day 9 amnion Primary villus epiblast hypoblast Primary Secondary yolk sac Remains of yolk sac lacunae primary yolk sac syncytiotrophoblast end of 2<sup>nd</sup> week cytotrophoblast primary yolk sac extraembryonic coelom extraembryonic



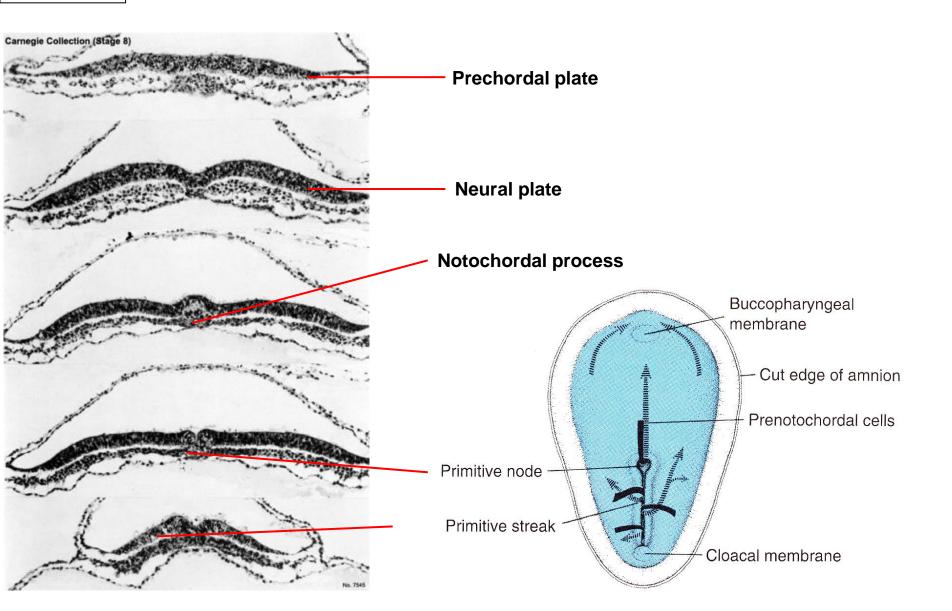


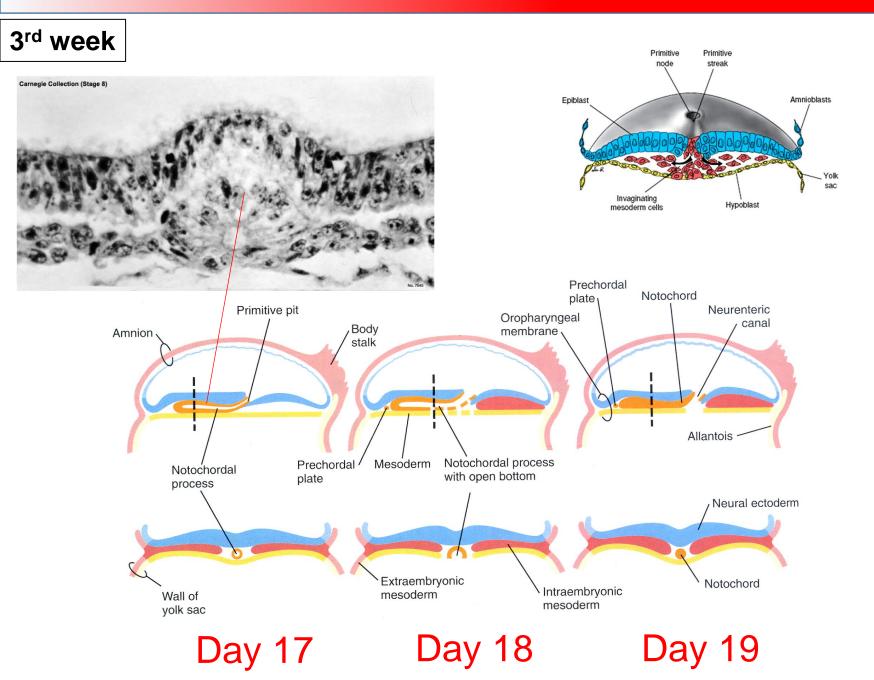
#### **NEW STRUCTURES**



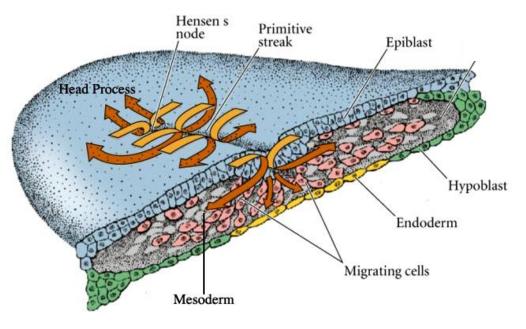


#### 3<sup>rd</sup> week



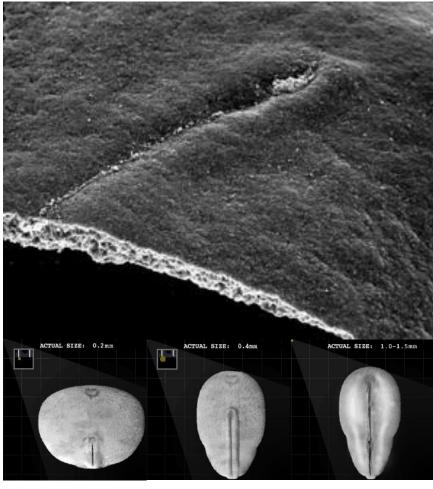


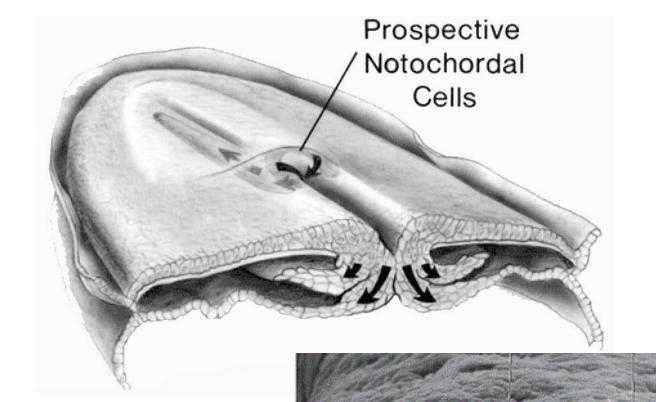
#### 3<sup>rd</sup> week



@ 2000 Sinauer Associates, Inc.

### A new cell population appears - MESODERM

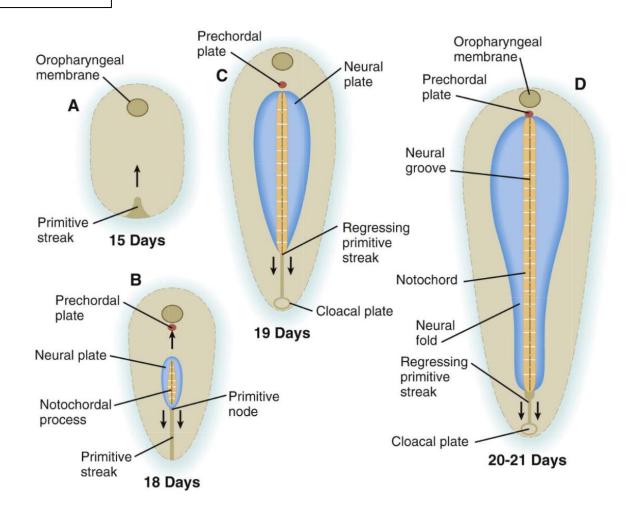




End of 2<sup>nd</sup> week

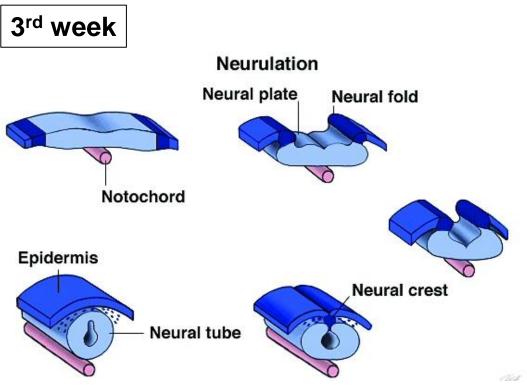
End of 3<sup>rd</sup> week

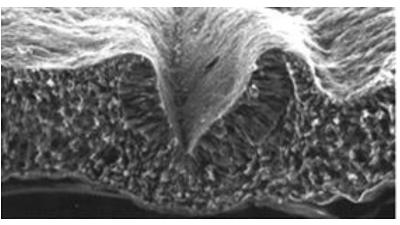
#### 3<sup>rd</sup> week

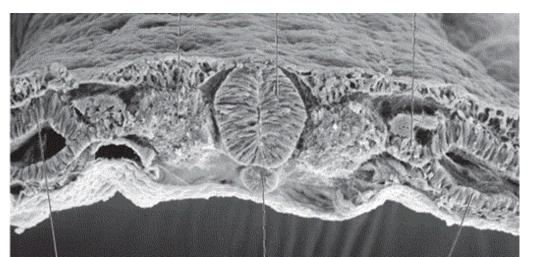


Notochord induces
differentiation of ectoderm
– cellular basis of nerve
system is established –
NEUROECTODERM

NEURULATION NEURAL TUBE

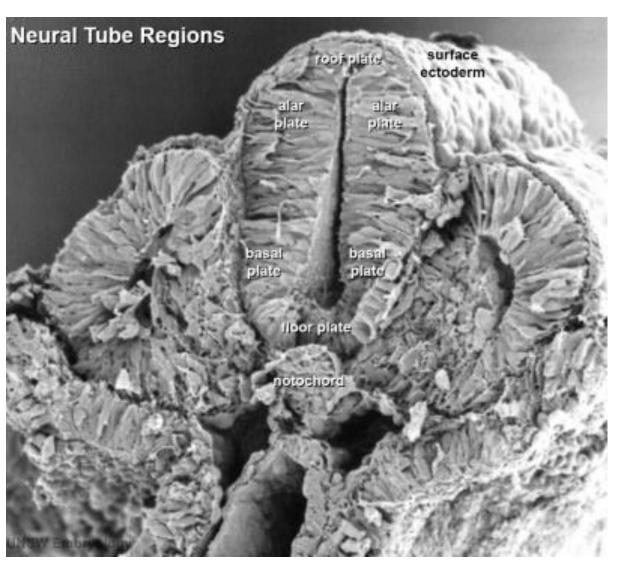


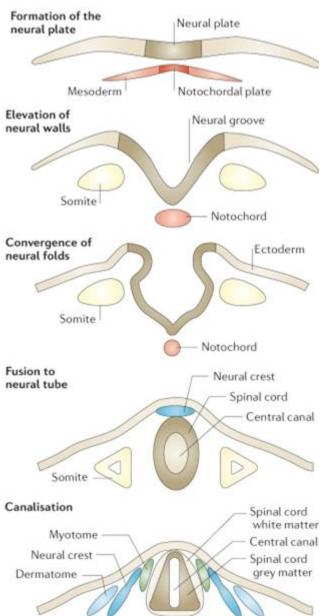


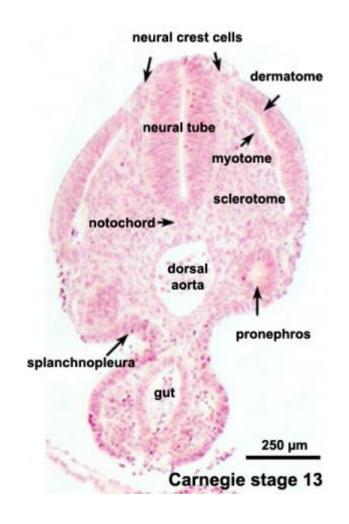


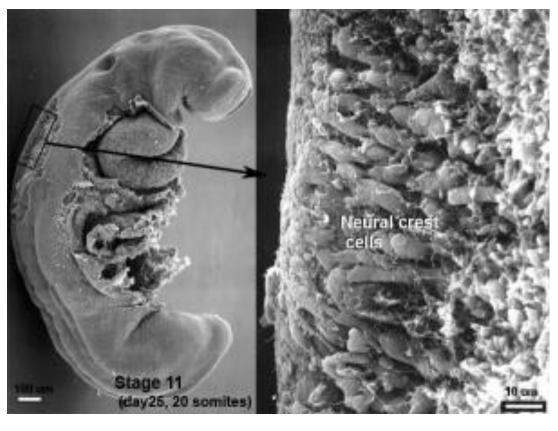
- Neural plate
- Neural folds
- Neural tube
- Neural crest

3<sup>rd</sup> – 4<sup>th</sup> week

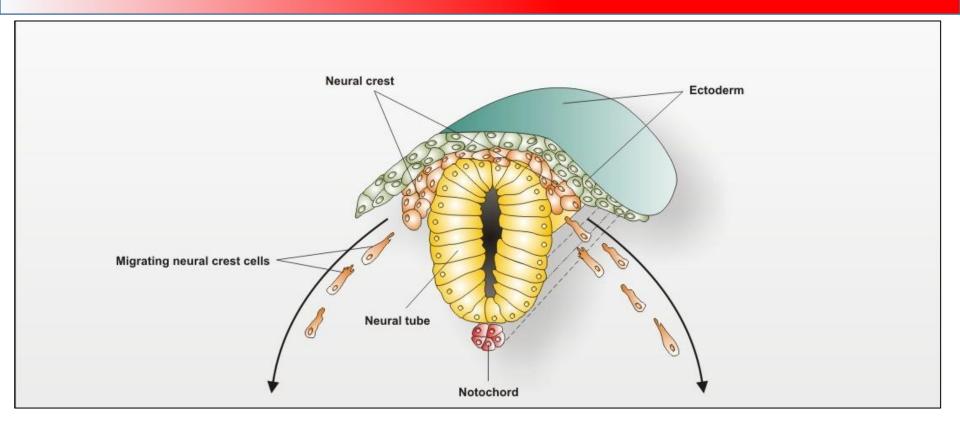






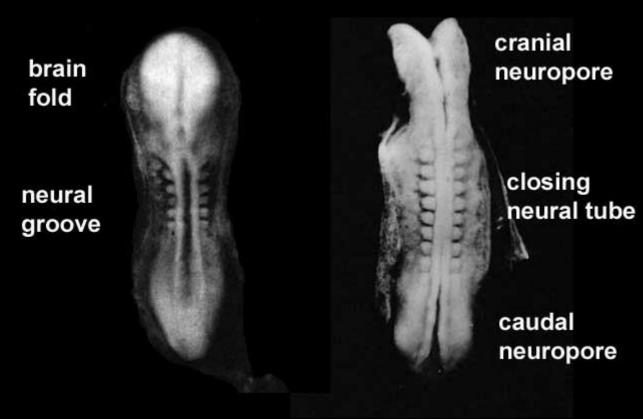


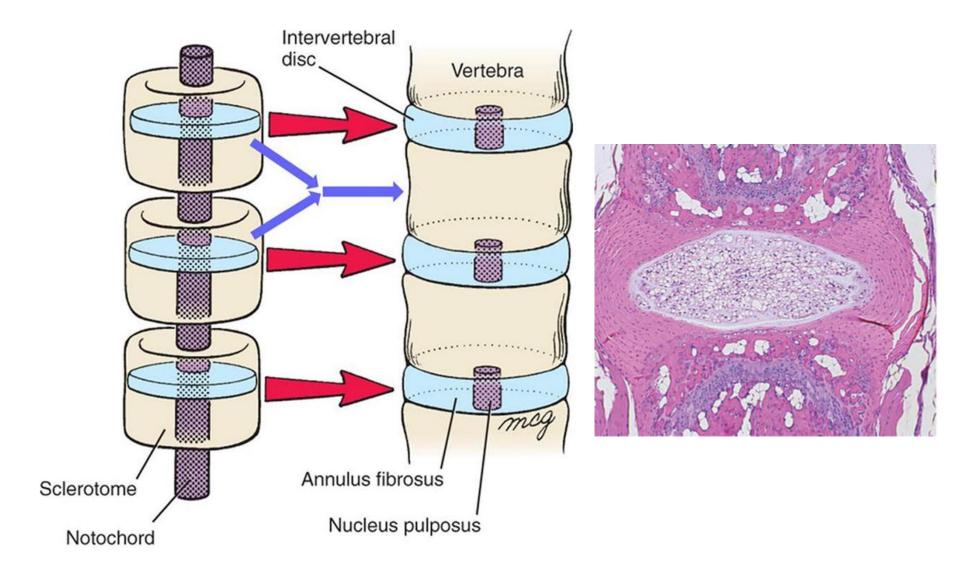
#### **NEURAL CREST AND ITS DERIVATIVES**



- Peripheral nerve system including glia
- Enterochromaffin cells
- Melanocytes
- Craniofacial connective tissue
- Odontoblasts
- Adrenal medulla
- ...

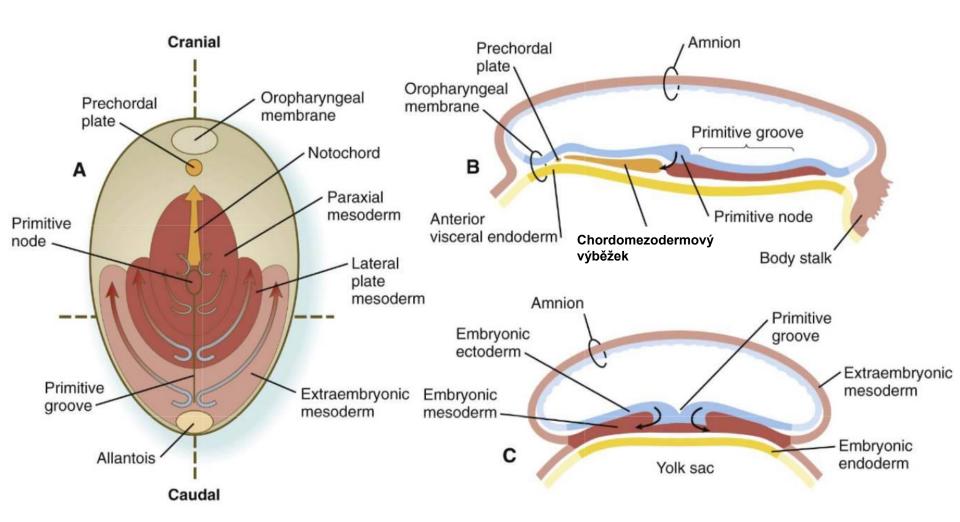




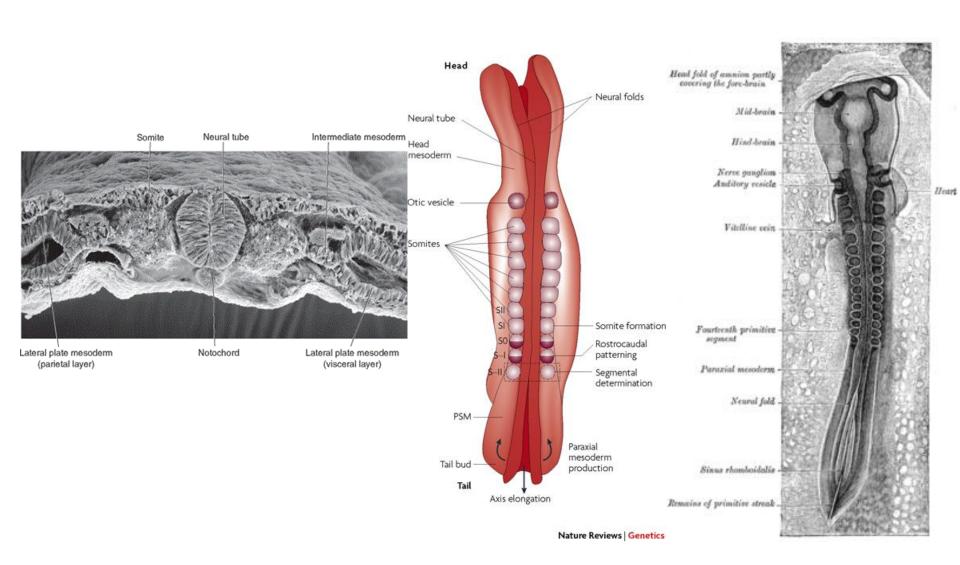


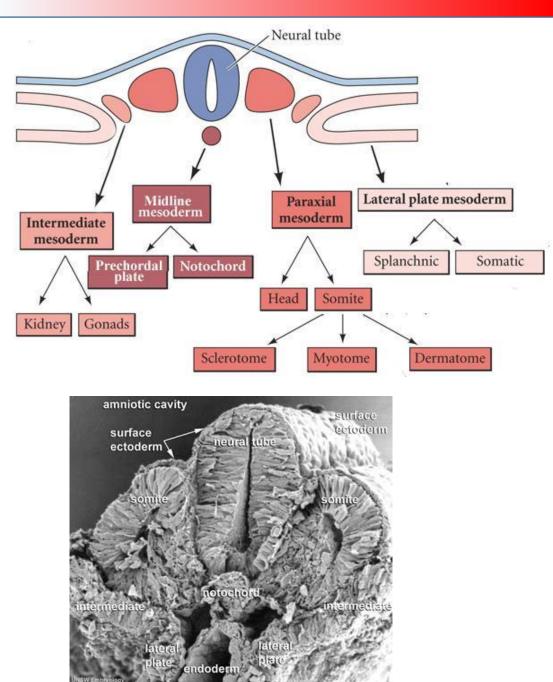
GASTRULATION MESODERM

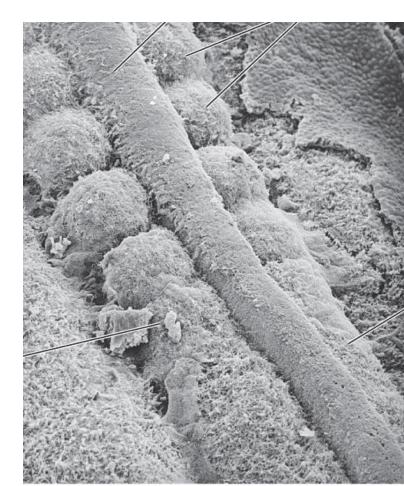
3<sup>rd</sup> week



3<sup>rd</sup> – 4<sup>th</sup> week

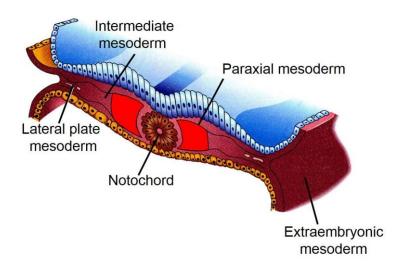




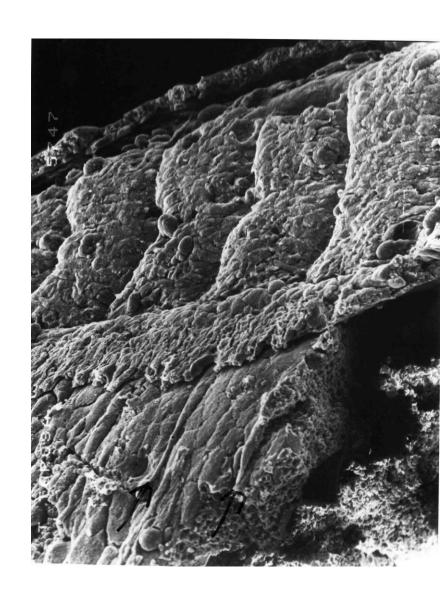


#### OTHER DERIVATIVES OF MESODERM

#### $3^{rd} - 4^{th}$ week



- heart, cardiovascular system
- urogenital system
- muscle and skeletal system
- hematopoietic and lymphatic systems
- connective tissue, dermis
- mesothelium



#### DEVELOPMENTAL DISORDERS DURING GASTRULATION

 Primitive streak is a temporary embryonic structure. Persistent primitive streak causes sacrococcgyeal teratoma.





Failure of primitive streak leads to absence of mesoderm in affected region -

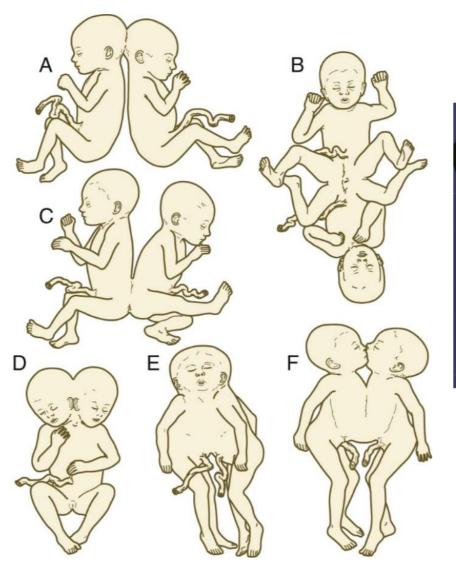
sirenomelia

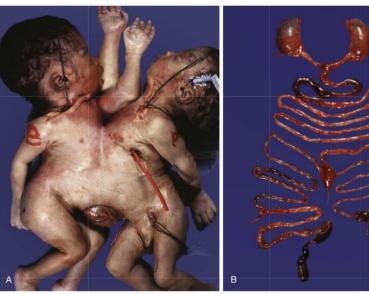
- limbs
- urogenital system
- GIT



#### **DEVELOPMENTAL DISORDERS DURING GASTRULATION**

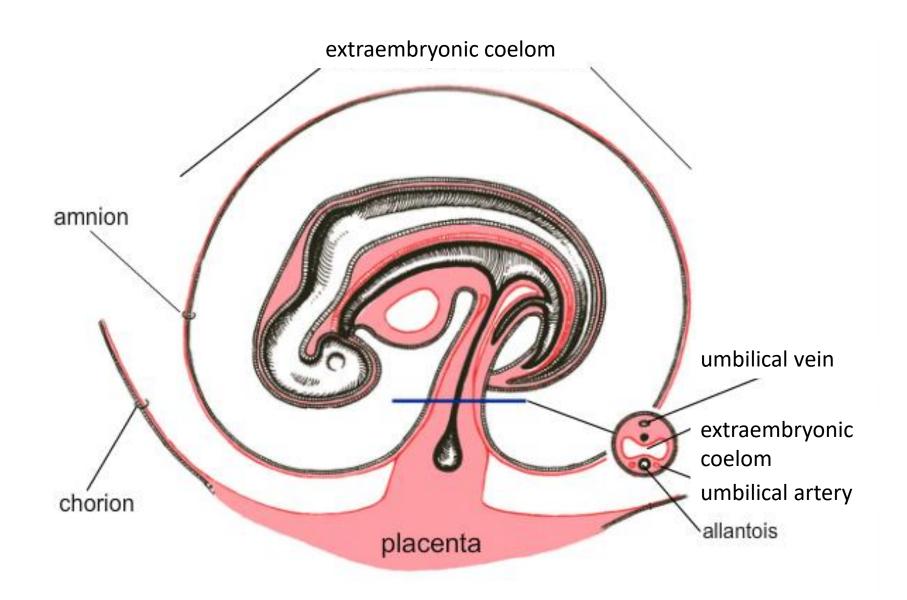
• If two primitive streaks form, conjoined twins may develop

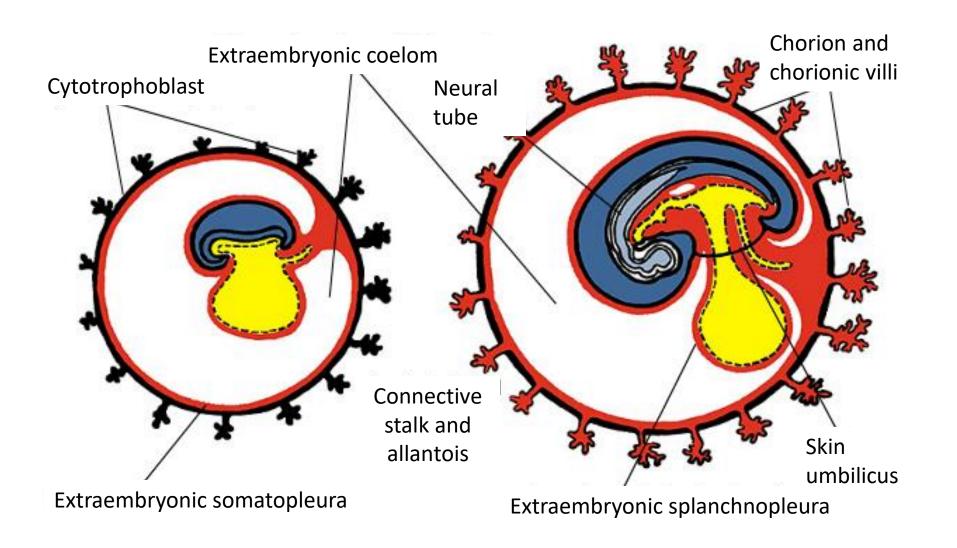




Hinagut 4<sup>th</sup> week Foregut Endoderm Amniotic cavity Heart Connecting Ectoderm tube. stalk Angiogenic cell cluster Allantois Pericardial. cavity uccopharyngeal membrane Cloacal membrane A B Buccopharyngeal membrane Cloacal membrane Lung bud Liver bud Midgut Heart als-R tube Remnant of the buccopharyngeal membrane Allantois Vitelline duct Yolk sac

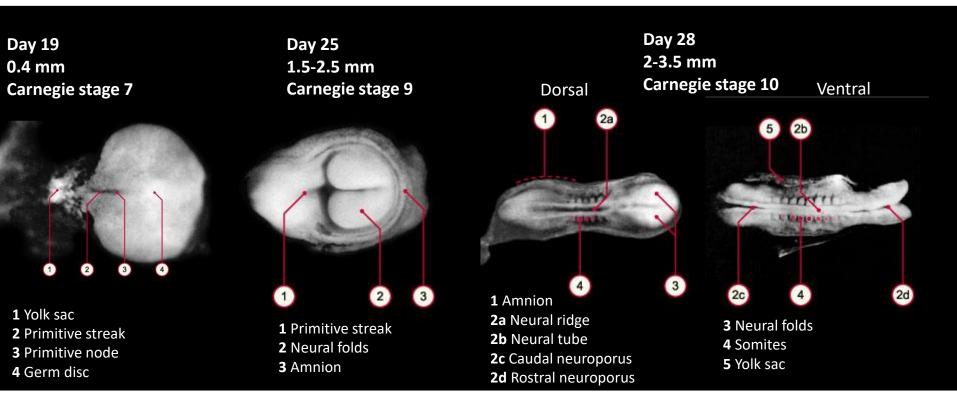
D





- bilaminar → trilaminar germ disc
- cephalocaudal and lateral flexion of embryo

http://www.embryology.ch/anglais/iperiodembry/carnegie03.html#st710









- Mesoderm segmentation
- Primitive gut
- Esophagotracheal diverticulum
- Heart (starts beating day 22-23)
- · Limb buds
- Primary brain vesicles, closing of neuropores
- Differentiation of neural crest
- Origin of thyroid and anterior pituitary
- Ectodermal placodes, optic vesicle
- Liver diverticulum
- Septum transversum

- · Segmentation of mesoderm continues
- Posterior pituitary
- · Heart septation begins
- Lung buds branch pseudoglandular stage of lung development
- Cochlea grows
- · Lens vesicle, nasal placodes
- Fourth brain ventricle forms
- · Pharyngeal arches, ridges and pouches
- · Limb buds grow
- Hematopoiesis in liver
- Retinal pigment

- Derivatives of endodermal pharyngeal pouches (parathyroid, thymus)
- · Adrenal gland
- · Heart and lungs descended to thorax
- Innervation of limbs, differentiation of myoblasts
- Face development maxillary and mandibulary processes, palatine, choans
- Telencephalon stratifies archicortex, paleocortex and neocortex. Choroid plexus
- · Rotation of stomach
- · Pancreatic diverticula fuse



- Secretion from endocrine pancreas
- Growth of liver, growth and luminization of bile ducts
- · Ossification of limbs begin
- · Development of brain nuclei

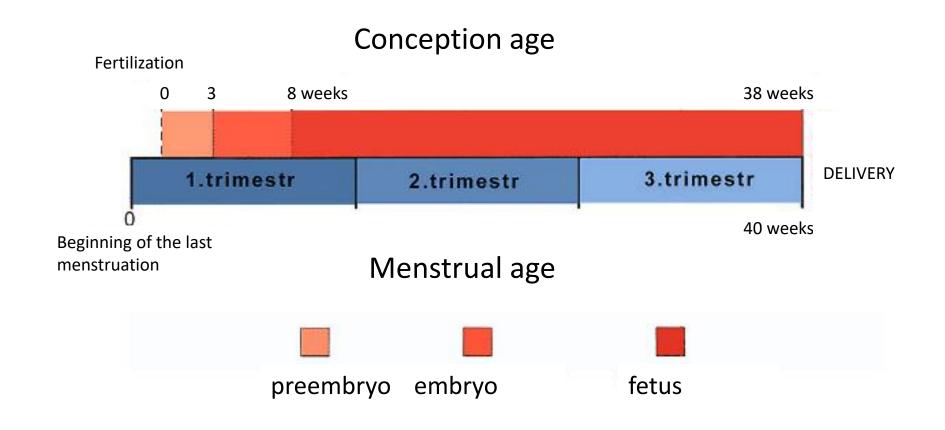


- Joints of upper and later lower limbs allow rotation
- Fingers grow
- Stratification of cerebellar cortex
- Perforation of anal membrane
- Herniation of intestinal loops
- Testes produce testosterone
- Nose, meatus, eyelids, developer, external ears start to grow
- Backbone 33-34 cartilaginous vertebrae
- · Embryonic tail diminished

#### **LENGTH OF PREGNANCY**

280 days (= 40 weeks = 10 lunar months) from the first day of the last **menstruation**266 days (= 38 weeks) **from ovulation** (gestation age)

Calculation of term: First day of the last menstruation + 1 year - 3 months + 7 days



FETAL DEVELOPMENT month 4-5



- Fetus swallows amniotic fluid necessary for GIT development
- Rapid growth of head (non-proportional to rest of body)
- Eyelids fuse
- · Ossification centers visible by ultrasound examination
- Development of external genitalia
- Kidneys produce urine, other organs start to work
- · Skeletal muscles innervated
- · Physiological umbilical hernia, in 12th weeks reposition of intestinal loops



- Rapid growth of fetus
- Ossification of skeleton
- · Face growths, mandible visible
- Apparent external genitalia

FETAL DEVELOPMENT month 5-9



- Limbs growth
- · Mother feels fetal movements
- Vernix caseosa, lanugo
- Short hairs and eyelashes
- · Fetus reacts to sound and later to light
- Lungs start to produce surfactant
- · Limit of viability



- · Eyelids open
- · Wrinkled skin with visible capillaries
- Subcutaneous fat
- Hairs grow
- · Maturing of organ systems



- Subcutaneous fat accumulates in limbs
- · Smooth, red skin
- · Hallmarks of full term fetus



Full term – related to <u>length of pregnancy</u> (menstrual age)

- preterm (<37 weeks)</li>
- full term (38 40 weeks)
- after term (>42 weeks) (meconium in amniotic fluid)

**Fetal maturity** – <u>development</u> of fetus: **mature** X **immature** 

#### HALLMARKS OF FETAL MATURITY

#### Major:

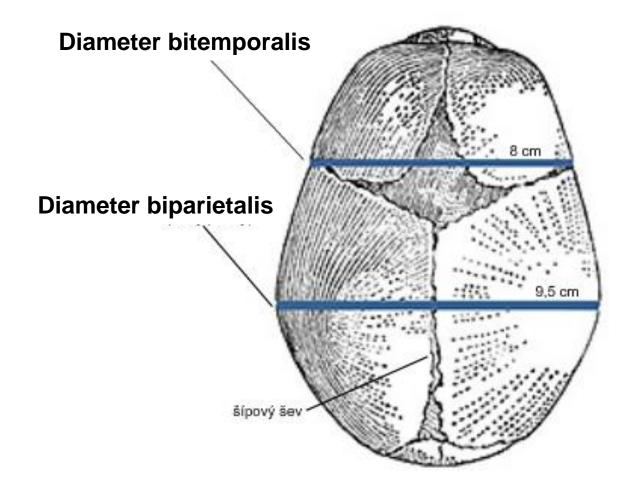
- length (50 51 cm),
- weight (around 3500 g, physiological range 2500 4000g),
- head sizes
- boys testes in scrotum, girls labia majora over labia minora

#### Minor:

- eutrophic fetus, subcutaneous fat
- skin is not blue (no cyanosis), lanugo remains on shoulders and back,
- eyelashes, hairs several cm long, nails over fingertips
- cranial bones hard, anterior and posterior fontanelle are palpable, and separated
- newborn cries and moves (Apgar score)

FULLTERM NEWBORN HEAD SIZE

Diameter bitemporalis – 8,00 cm (join of the most distant points on sutura coronaria) Diameter biparietalis – 9,5 cm (join of midpoints of tubera parietalia)

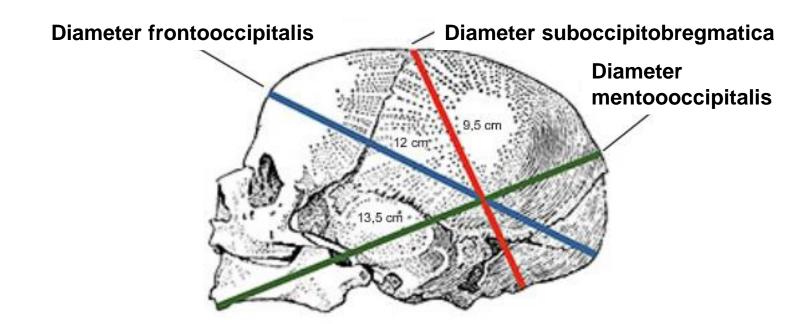


FULLTERM NEWBORN HEAD SIZE

#### **Oblique sizes:**

Diameter frontooccipitalis – 12.0 cm (join of forehead midpoint and most distant point of occiput)

- Circumferentia frontooccipitalis 34.0 cm
- Diameter suboccipitobregmatica 9.5 cm (join of protuberantia occipitalis externa and midpoint of large fontanelle)
- Circumferentia suboccipitobregmatica 32.0 cm
- Diameter mentooccipitalis 13.5 cm (join of chin midpoint and most distant point of occiput)
- circumferentia mentoocipitalis 35 36 cm
- Diameter biacromialis 12.0 cm, circumferentia biacromialis 35 cm
- (join of acromion acromion)



#### **RULE OF HASSE**

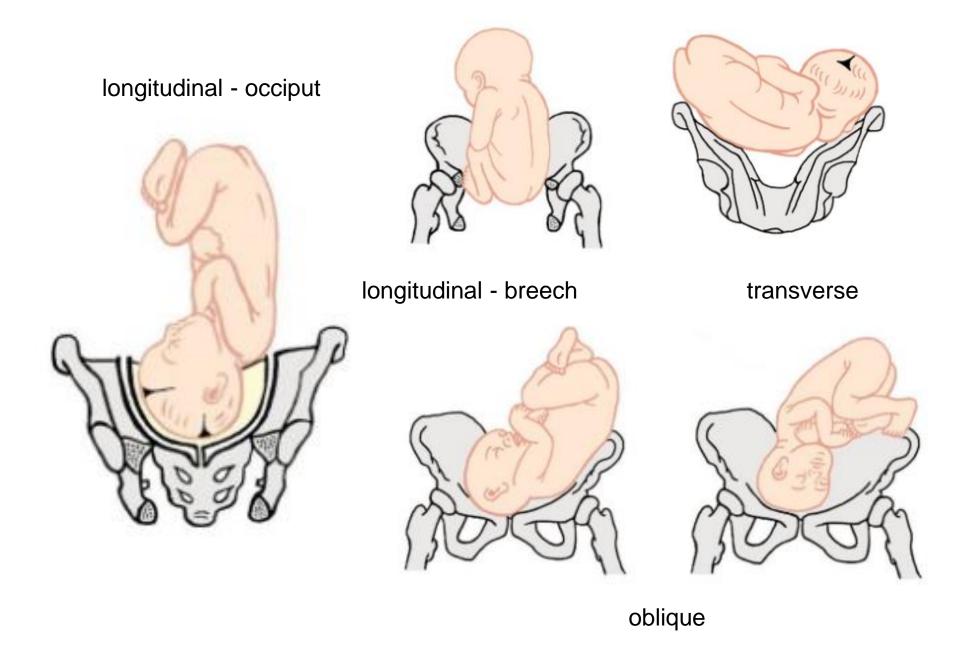
- forensic medicne
- 3. 5. lunar month: length in cm = square of month
- 6. 10. lunar month: length in cm = months multiplied by 5

<b>Lunar month</b>	Length of fetus[cm]
3	9
4	16
5	25
6	30
7	35
8	40
9	45
10	50

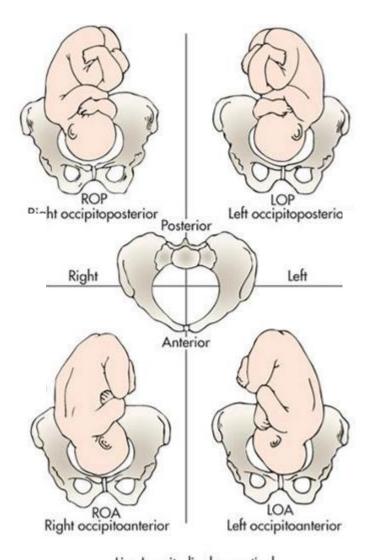
#### **FETUS IN UTERUS**

- 1. **LIE** (SITUS) = relationship of the long axis of the fetus to that of the mother
- longitudinal: (99 %)
- transverse: (1 %) perpendicular axes
- oblique: unstable → longitudinal or transverse position
- 2. **POSITION** (*POSITIO*) = fetal backbone relative to uterus ridge
- first= left (back to the left)
- second= right (back to the right)
- first/second common/less common
- 3. **FETAL HABITUS** (*HABITUS*) = relationship of one fetal part to another
- regular = head and limbs in flexion
- irregular = everything else
- 4. **PRESENTATION** (*PRAESENTATIO*) = that part of the fetus lying over the pelvic inlet; the presenting body part of the fetus.
- occiput (most common)
- vertex, forehead, face (1%)
- breech
- trunk, shoulder

# SITUS



# 2. COMMON "RIGHT"



1. COMMON "LEFT"

2. LESS COMMON

Lie: Longitudinal or vertical Presentation: Vertex Reference point: Occiput Attitude: General flexion 1. LESS COMMON

# **HABITUS AND PRESENTATION**

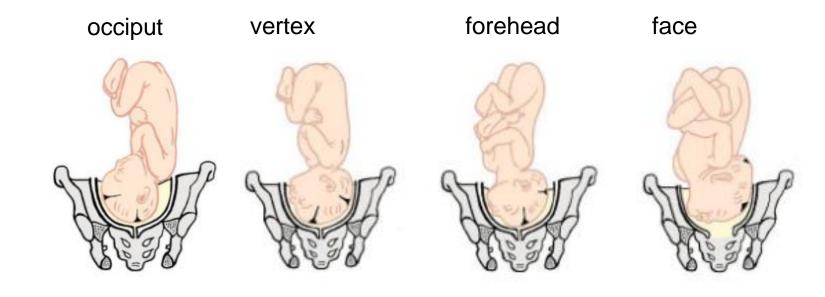


# **HABITUS**

irregular (any other)

regular

# **PRESENTATION**



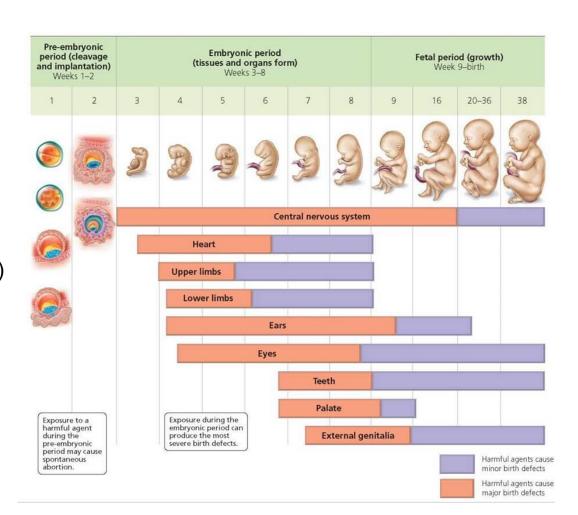
#### PHYSIOLOGICAL IMPOSITION OF FETUS IN UTERUS

- LIE LONGITUDINAL HEAD FIRST
- POSITION FIRST COMMON
- HABITUS REGULAR
- PRESENTATION OCCIPUT



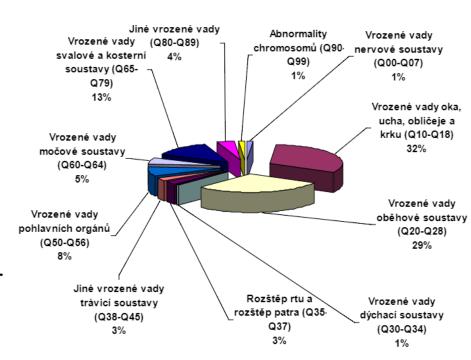
#### INTRODUCTION TO TERATOLOGY

- Congenital disorders due to abnormal developmental events
- Genetic (inherited) or nongenetic (external) causes
- Teratogens
- Critical developmental periods
- Life style (alcohol, smoking, drugs)
- Infections (rubeola, HIV, toxoplasmosis)
- Lack or abundance of key substances (folic acid × retinoids)
- Chronic diseases (medical treatment)



#### INTRODUCTION TO PRENATAL DIAGNOSTICS

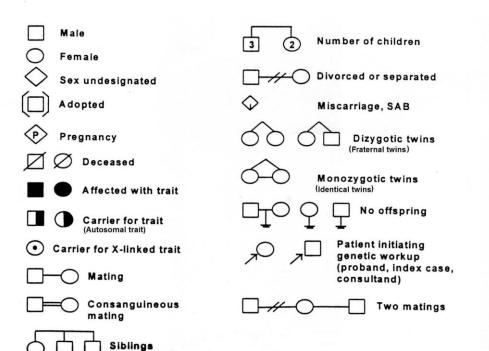
- Interdisciplinary care biochemistry, genetics, gynecology and obstetrics, neonatology – parts of fetal medicine
- Revealing high risk pregnancies, access to preventive and therapeutic care
- Preventing delivery of fetuses with severe congenital malformations
- Support of delivery of genetically high-risk babies
- Planning and providing clinical care
- Genetic counselling
- Biochemical and ultrasound screening
- Karyotyping and DNA diagnostics
- Clinical diagnostics
- Indication:
- congenital disorders in family
- positive screening in 1<sup>st</sup> or 2<sup>nd</sup> trimester
- abnormal finding by ultrasound
- maternal age (over 35 years)

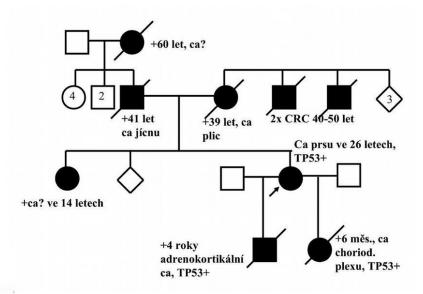


ČR 1994-2008

#### **GENETIC COUNSELING**

- Anamnesis (case history)
- Preconception counselling
- Explaining of examination results, causes, clinical symptoms, therapeutic options
- Minimization of risk of repeated disease
- Providing diagnosis and information for free choice
- Providing precise diagnosis and risk estimation
- Providing care during pregnancy and later

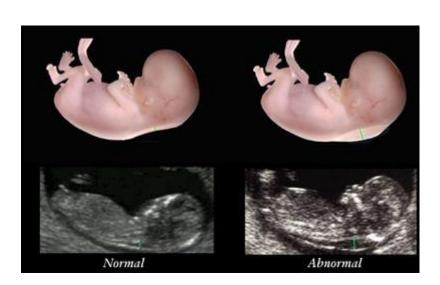




NONDIRECTIVE ALL EXAMINATIONS AND PROCEDURES ARE VOLUNTARY

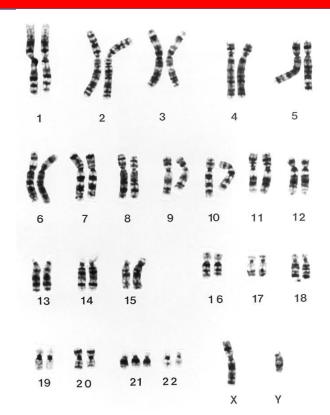
#### **BIOCHEMICAL SCREENING**

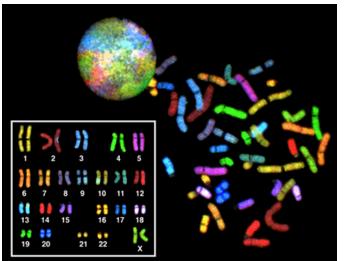
- Non invasive
- Revealing high risk pregnancies chromosomal aberrations and clefts
- Screening is not a diagnostics → further examinations (amniocentesis, karyotype, US)
- Tests between weeks 14-16 ("TRIPLE test")
- low sensitivity and specificity (50-60%), high false positivity (70%)
- AFP, E3, hCG
- chromosomal aberrations, abnormal closing of neural tube, defects of body walls
- Combined screening in week 11-13
- chromosomal aberrations Down: 47,XY,+21, Edwards: 47,XY,+18, Patau 47,XY,+18
- US -nuchal translucence, NT
- PAPP-A, hCG (multiplies of median, MoM)
- age included in algorithm
- output: screening positive vs. negative (limit 1:100)



#### **INVASIVE DIAGNOSTICS**

- Amniocentesis
- 16<sup>th</sup>-20<sup>th</sup> week
- US controlled amniotic fluid aspiration
- Cell culture, karyotype
- Risk of miscarriage 0.5-1%
- Chorion villus biopsy
- 10<sup>th</sup>-13<sup>th</sup> week
- Karyotype, molecular genetic examination
- Risk 0.5-1%
- Cordocentesis
- 22<sup>nd</sup> week
- Sampling of venous umbilical blood
- Now diagnostics and therapy of blood diseases (anemia, infections), or diagnostics in multiple pregnancies
- Risk 1%
- Fetoscopy
- Transabdominally (earlier transcervically)
- Visualization and fetal biopsy
- Risk 3-10%, done rarely

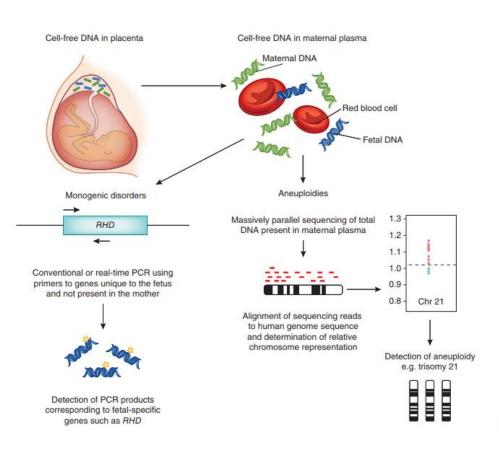




#### **ULTRASOUND DIAGNOSTICS**

- 6-8<sup>th</sup> week
- confirmation of pregnancy, heart action
- number of fetuses
- 13-14<sup>th</sup> week
- nuchal translucence (risk > 3 mm)
- nasal bone (present × absent), **minor markers** (omphalocele, tricuspidal regurgitation, abnormality in ductus venosus flow, enlargement of urinary bladder-megavesica)
- fetal size
- 20-22<sup>nd</sup> week
- detailed screening
- fetal biometry (biparietal diameter, head circumference, length of femur)
- head and CNS (shape, cavity in septum pellucidum, ventricles, cerebellum, cisterna magna), face (lip, jaws, nose, orbits, profile), heart (action, size, axis, 4-chamber projection, outflow tracts, ...), thorax (pathological structures), abdominal cavity (stomach, intestine, kidneys, urinary bladder, umbilicus and umbilical vessels), backbone, limbs, palms, feet
- placenta, volume of amniotic fluid
- 30<sup>th</sup> week
- fetal size
- volume of amniotic fluid
- placenta (exclusion of *placenta praevia*)

#### ADVANCEMENTS IN MOLECULAR GENETICS



Analysis of cell-free fetal DNA in maternal blood Since 12<sup>th</sup> week Massive paralell sequencing (Next-Gen Sequencing) Common aneuploidies (trisomy 21,13,18) Monogeneous disorders

SOP-M8 NEINVAZIVNÍ DETEKCE ANEUPLOIDIÍ CHROMOZOMŮ 13, 18 A 21 POMOCÍ MULTIPLEX PCR A MASIVNÍHO PARALELNÍHO SEKVENOVÁNÍ (MPS)

Test Clarigo se značkou "CE", která je nezbytná pro provedení tohoto vyšetření v zemích EU, splňuje základní požadavky Směrnice Rady IVD 98/79/EC pro in vitro diagnostiku.

#### VÝSLEDEK VYŠETŘENÍ:

Chromozom	Stav	Fetální frakce	Předpokládané pohlaví plodu
13	normální	7,1 %	ženské
18	normální		
21	normální		

#### ZÁVĚR:

Analýzou volné fetální DNA cirkulující v krvi těhotné nebylo zjištěno zvýšené riziko aneuploidie chromozomů 13, 18 a 21.

Komentář: doporučujeme genetickou konzultaci.

Pozn: Při patologickém nálezu je výsledek nutné ověřit některým z invazivních postupů (např. odběr plodové vody, choriových klků, kordocentéza s následnou QF-PCR analýzou nebo stanovením klasického karyotypu apod.).



# THANK YOU FOR ATTENTION

pvanhara@med.muni.cz
http://www.med.muni.cz/histology