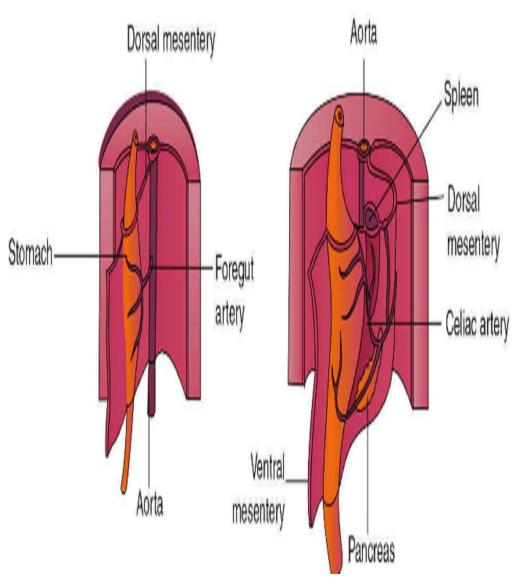
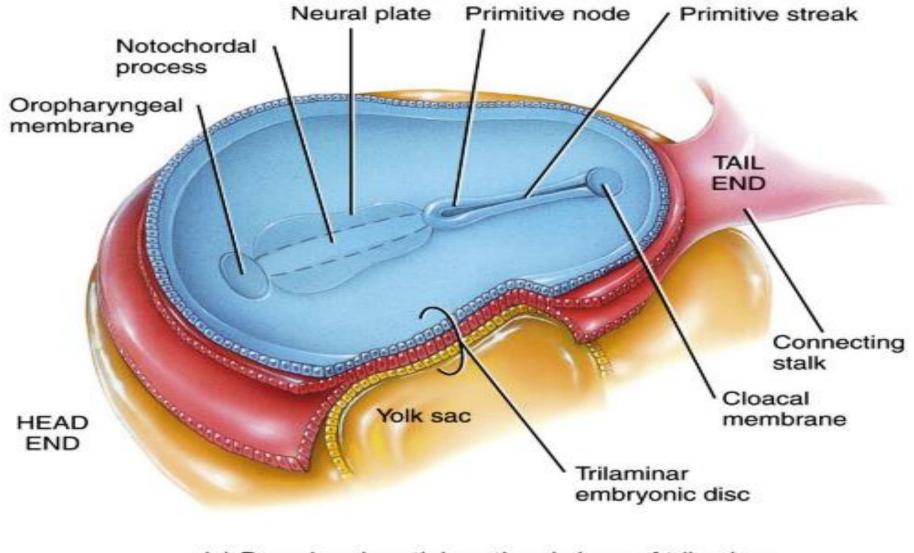
# DEV. OF ESOPH., STOM&CH & DUO.



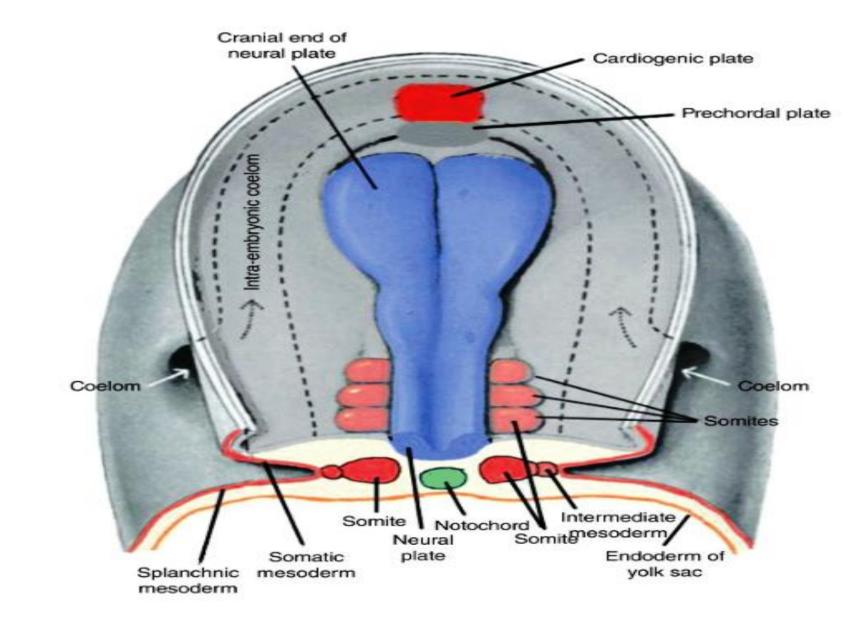
#### BÝ DR ABULMAATÝ MOHAMED ASSISTANT PROFESSOR ANATOMÝ & EMBRÝOLOGÝ MUTAH UNIVERSITÝ





 (a) Dorsal and partial sectional views of trilaminar embryonic disc, about 16 days after fertilization

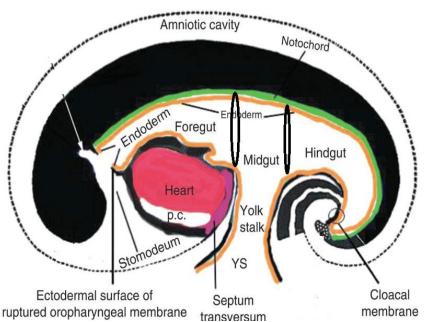
# REV.



#### REV. Amniotic cavity Amniotic Notochord Neural tube cavity Neural plate(tube) Notochord Ectoderm -Ectoderm Oropharyngeal membrane (prechordal plate) Endoderm YS Heart Prechordal Endoderm plate YS Cardiogenic plate Amniotic cavity Notochord Amniotic cavity Oropharyngeal membrane Notochord Endoderm Endoderm Foregut oregut Hindgut Midgut Hindgut Midgut Heart Endoderm Heart p.c. Yolk p.c. Yolk stalk stalk Stomodeum Cloacal YS Stomodeum membrane Septum Cloacal Ectodermal surface of transversum Septum ruptured oropharyngeal membrane membrane transversum

### DIVISIONS OF GUT

-folding of embryonic disc leading to incorporation of large part of the yolk sac (lined by endoderm) inside embryo leading to formation of the primitive gut primitive gut is divided into 3 parts:



- 1- foregut: included in head fold & ends blindly by BPM
- 2- hindgut: included in tail fold & ends blindly by CM
- 3-midgut: middle part & is connected to the
- yolk sac by vitellointestinal (vitelline) duct
- ant. intestinal portal: junction () foregut and midgut
- post. intestinal portal: junction () midgut and hindgut.
- -The wall of gut is formed of endodermal lining
- & covering of visceral (splanchnic) mesoderm.

## FOREGUT

#### **Extents:**

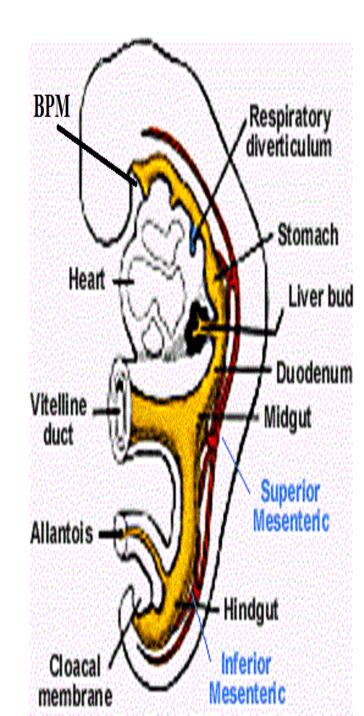
from buccopharyngeal membrane BPM to ant. intestinal portal (origin of liver bud). **Parts & derivatives :** 

divided by laryngeotracheal(respiratory) diverticulum into:

- 1- cranial (pharyngeal) part:
- from the BPM to
- the laryngeotracheal diverticulum
- will form: post. Part mouth cavity and pharynx

2- caudal part:

- -from the laryngeotracheal diverticulum to origin of liver bud
- -will form esophagus, stomach,
- upper half of duodenum
- liver, pancreas and biliary system

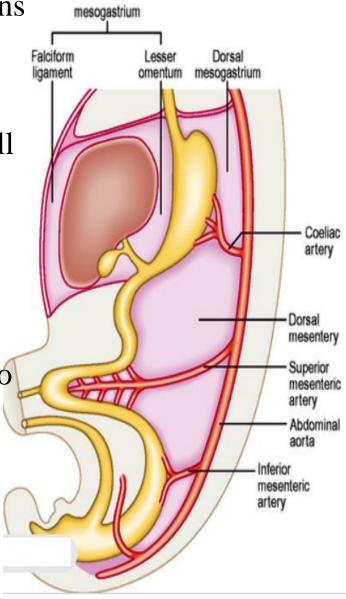


# MESENTERY

**Ventral mesentery:-**

**Def. :-** 2 layers of peritoneum connect the organs to posterior & anterior abdominal wall

Site:-Connect the gut to anterior abdominal wall extent:- from lower end of the esophagus to 1<sup>st</sup> inch of duodenum (the part opposite the stomach is called ventral mesogastrium) Fate:- liver will develop inside it dividing it into Lesser omentum :- between the liver & gut falciform ligament:- between the liver & anterior abdominal wall



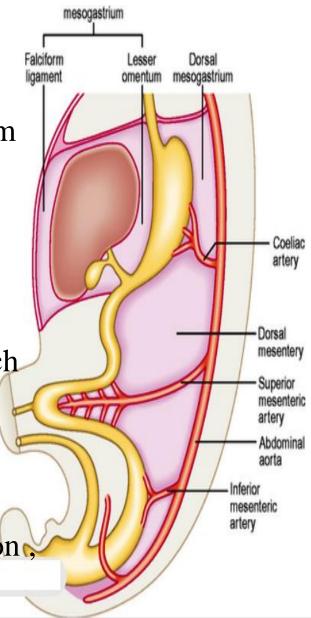


#### **Dorsal mesentery:-**

Site:-Connect the gut to posterior abdominal wall extent:- from lower end of the esophagus to rectum Parts &Fate:-

Dorsal mesogastrium :- opposite the stomach will form greater omentum Dorsal mesoduodenum :- opposite the duo. Will disappear except 1<sup>st</sup> inch Mesentery proper:-opposite the jejunum &ileum will form the mesentery mesocolon :- opposite the colon will disappear except transverse mesocolon

sigmoid mesocolon, mesoappendix

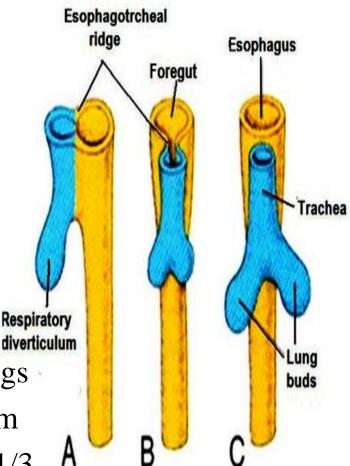


### ESOPHAGUS

- **A- developmental source:**
- caudal part of foregut,
- from pharynx
- (level of laryngeotracheal diverticulum)
- to stomach

#### **B- development:**

- $\Box$  at 4th week esophagus is short but
- elongates with the descent of the heart and lungs
- $\hfill\square$  muscles develop from surrounding mesoderm
- & are striated in upper 2/3 & smooth in lower 1/3
- □ proliferation of the lining endoderm then recanalization to
- change epithelium from columnar to stratified squamous
- $\hfill\square$  closure of edges of laryngeotracheal groove to separated esophagus from trachea



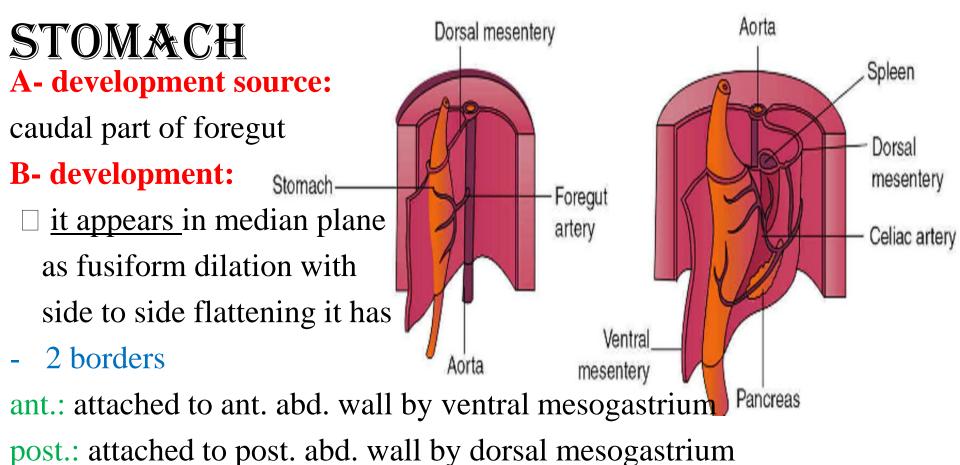
### ESOPHAGUS

#### **C- congenital anomalies:**

- 1- esophageal atresia: obliterated esophagus due to failure of recanalization
- 2- esophageal stenosis: local narrowing of esophagus due to incomplete recanalization
- 3- short esophagus: accompanied by thoracic stomach
- 4- trachea- esophageal fistula with esophageal atresia

Herniated Stomach

- $\Box$  due to defective closure of laryngeotracheal groove
- $\Box$  commonest anomaly
- □ forms
- a- fistula above atresia
- b- fistula below atresia: commonest
- c- fistula above & below atresia
- d- H shaped fistula without atresia



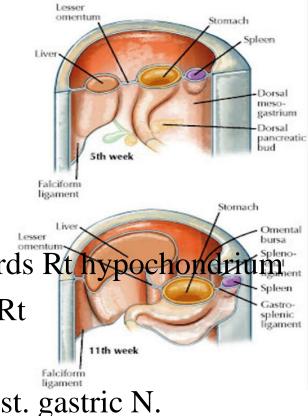
- -2 surfaces
- Rt: supplied by Rt vagus Lt: supplied by Lt vagus
- □ <u>change in shape:</u> by differential growth
- -post. border grows rapidly forming the greater curvature.
- -ant. border grows slowly forming the lesser curvature

#### STOM&CH B- development:

□ <u>change in position:</u> by rotation

#### 1st rotation:

- 90 to Rt (clockwise) around its longitudinal axis
- by growth of liver in ventral mesogastrium towards Rt hypochondrin
- results in 1- lesser curvature become directed to Rt ,greater curvature become directed to Lt
- 2- Rt surface become post. & Rt vagus become post. gastric N.
- Lt surface become ant. & Lt vagus become ant. gastric N.
- 3- The Rt part of peritoneal cavity becomes the lesser sac behind stomach
- 4- Ventral mesogastrium forms lesser omentum & peritoneal lig. of liver Dorsal mesogastrium forms gastrosplenic lig., gastrophrenic lig., lienorenal lig. and greater omentum



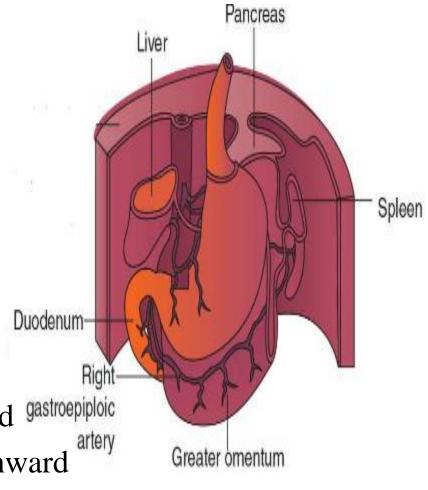
## STOM&CH

### **B- development:**

#### change in position:

2nd rotation:

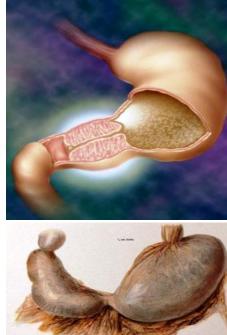
- 900 to Rt(clockwise) around antero-posterior axis
- Due to presence of diaphragm
- results in
- lesser curvature become directed upward gastroepiploic artery greater curvature become directed downward
- □ <u>muscles develop</u> from surrounding splanchnic mesoderm
- proliferation of circular layer at pyloric end
- To form pyloric sphincter

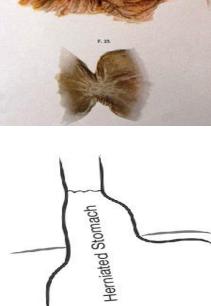


### STOM&CH

#### **C- congenital anomalies:**

- 1- hypertrophic pyloric stenosis
- one of the commonest anomalies of stomach.
- due to abnormal thickening of pyloric sphincter.
- 2- Hour-glass stomach
- stomach with local constriction
- dividing it into 2 parts
- 3- thoracic stomach
- stomach in thoracic cavity due to short esophagus
- 4- transposition of stomach
- stomach in Rt hypochondrium due to abnormal growth of liver to Lt hypochondrium

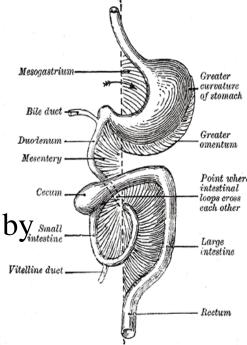




# DUODENUM

#### developmental sources:

- 1- upper part from terminal part of foregut
- 2-lower part from proximal part of midgut
- N.B: junction () foregut & midgut is marked in adult by small opening of bile duct in second part of duodenum Vitelline duct **development:**
- □ terminal part of foregut & proximal part of midgut grow rapidly formation of u shaped duodenal loop that is convex ant. &covered by peritoneum & attached to post. abd. wall by mesoduodenum
- $\Box$  loop rotate 90 o to Rt due to rotation of stomach convexity become to Rt
- □ Fixation: Mesoduodenum degenerate
- □ duodenum become retroperitoneal except 1st inch



- absorped

# DUODENUM

- **Congenital anomalies:**
- 1- atresia &stenosis
- 2-diverticulae
- 3-persistence of mesoduodenum

