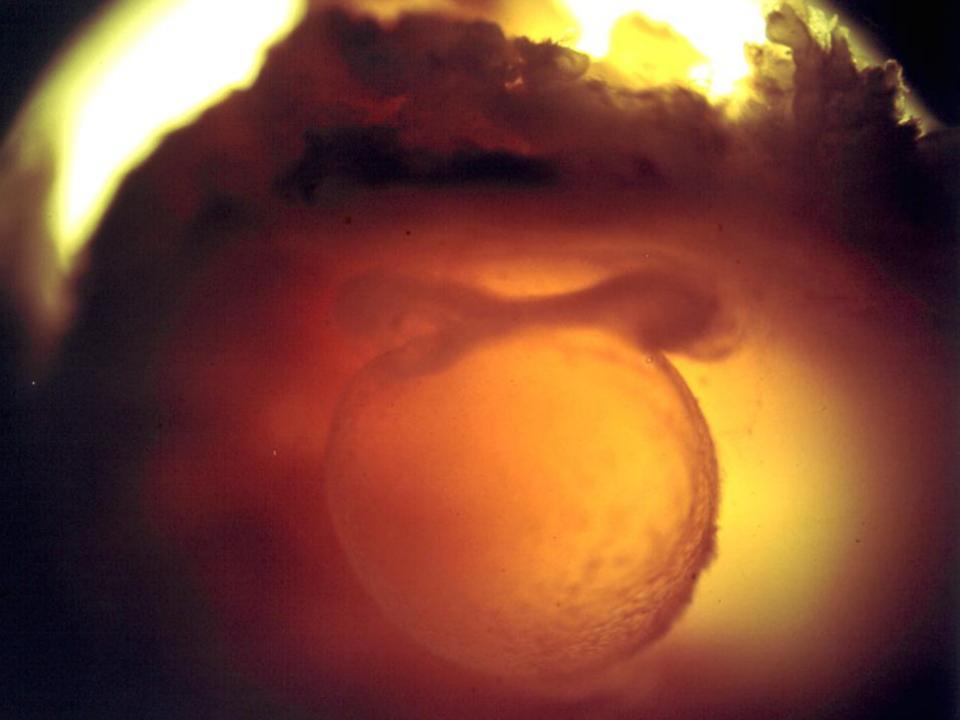
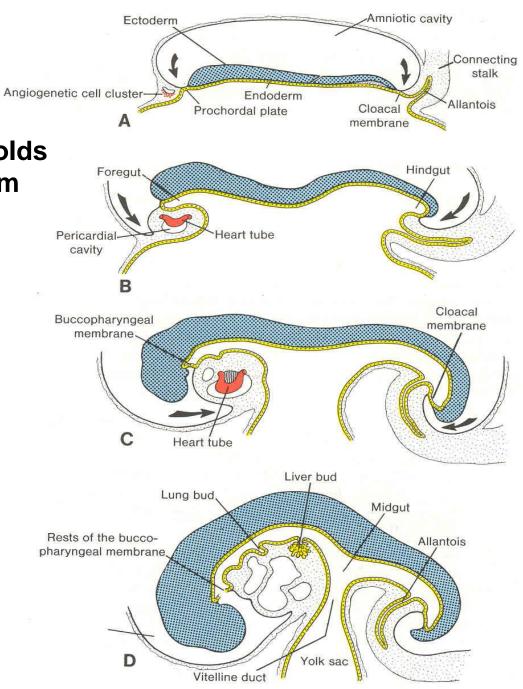
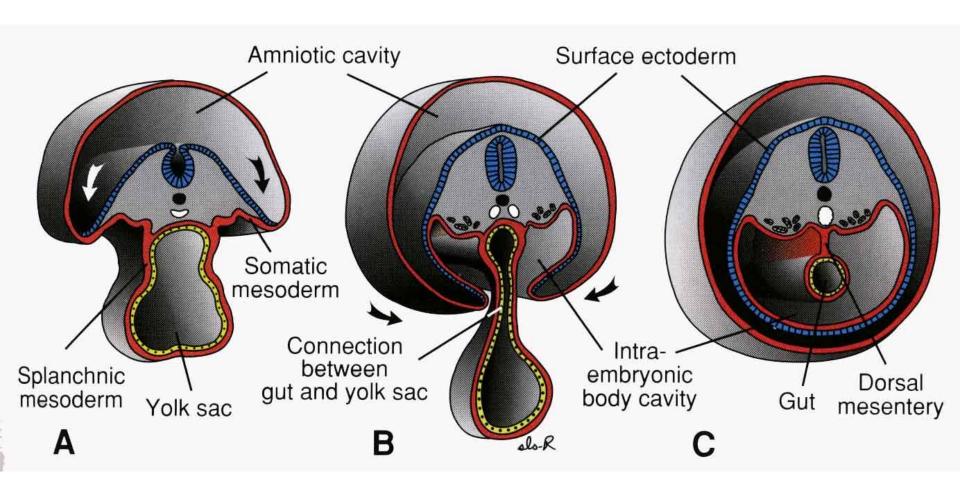
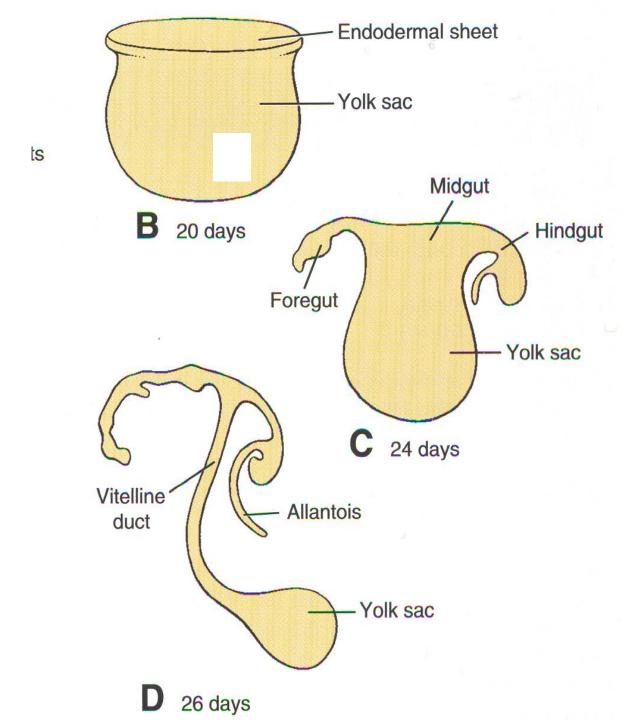
DEVELOPMENT OF THE DIGESTIVE SYSTEM

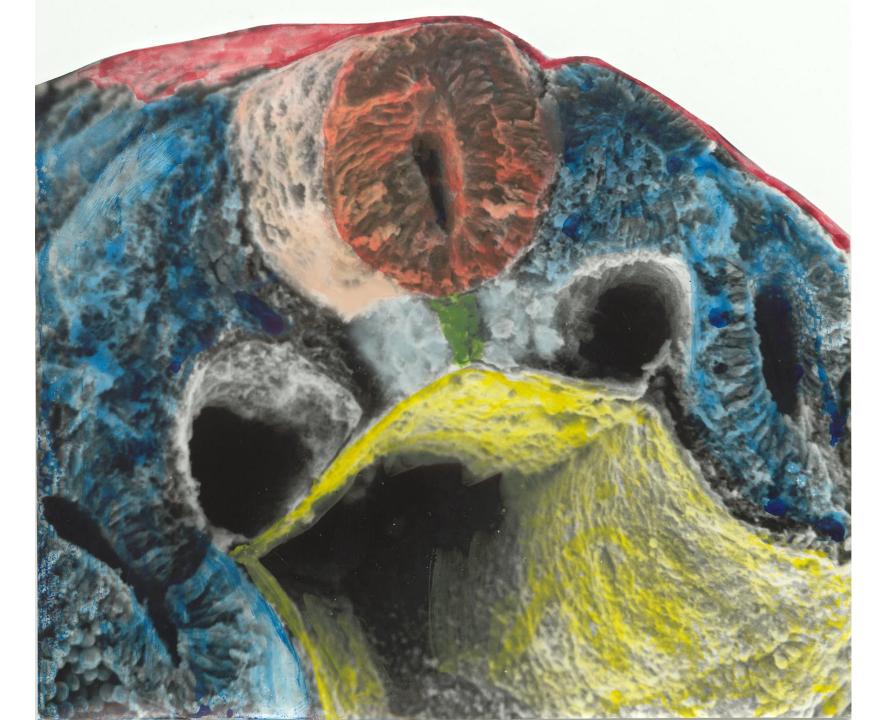


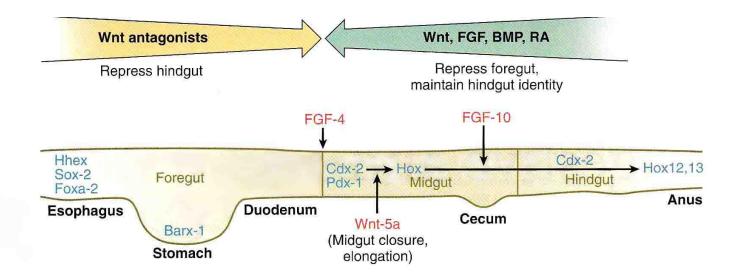
folding of the embryo and development of amniotic folds changes the trilaminar germ disc into the three-dimensional tube





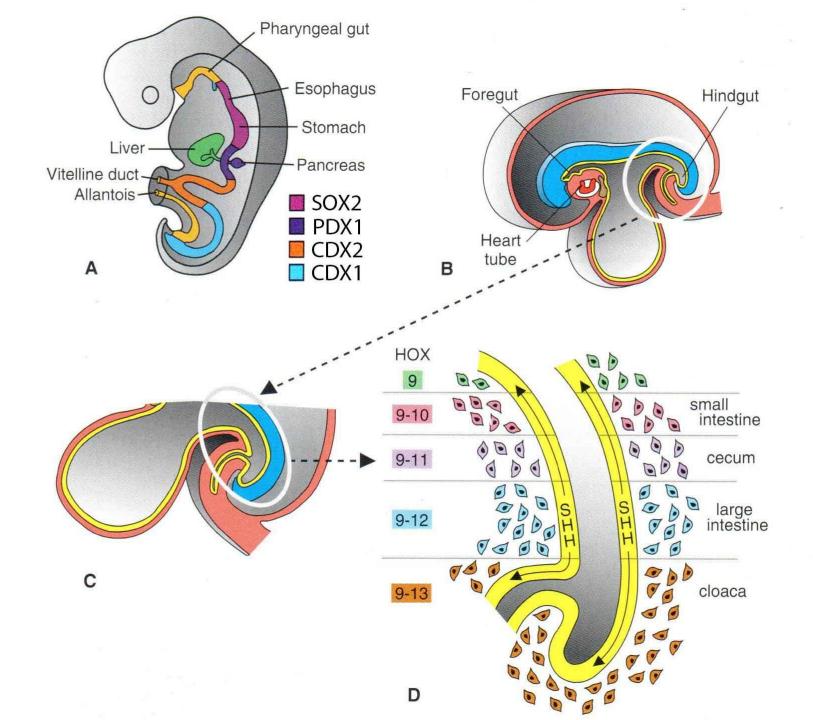


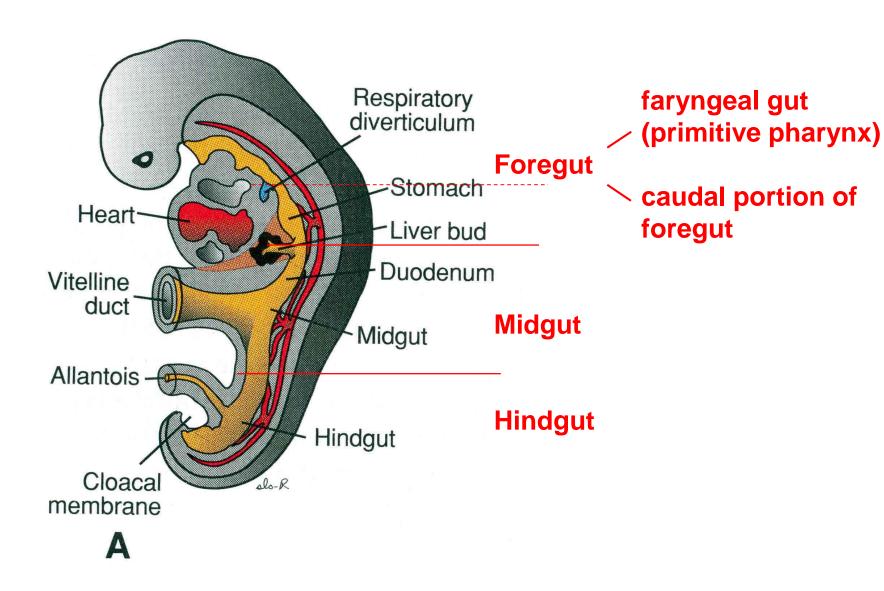


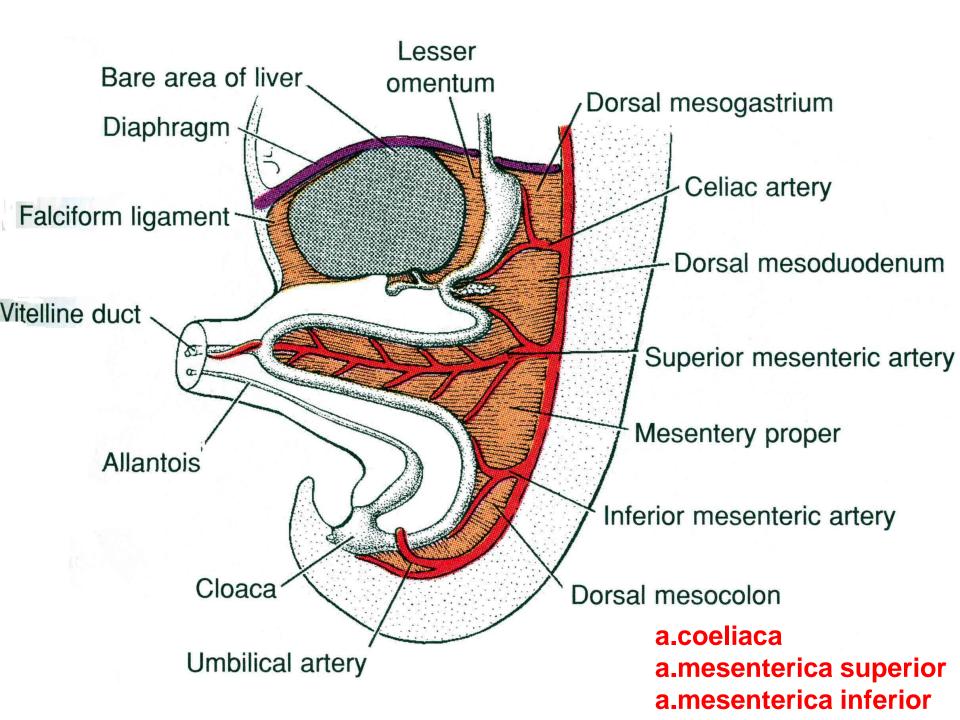


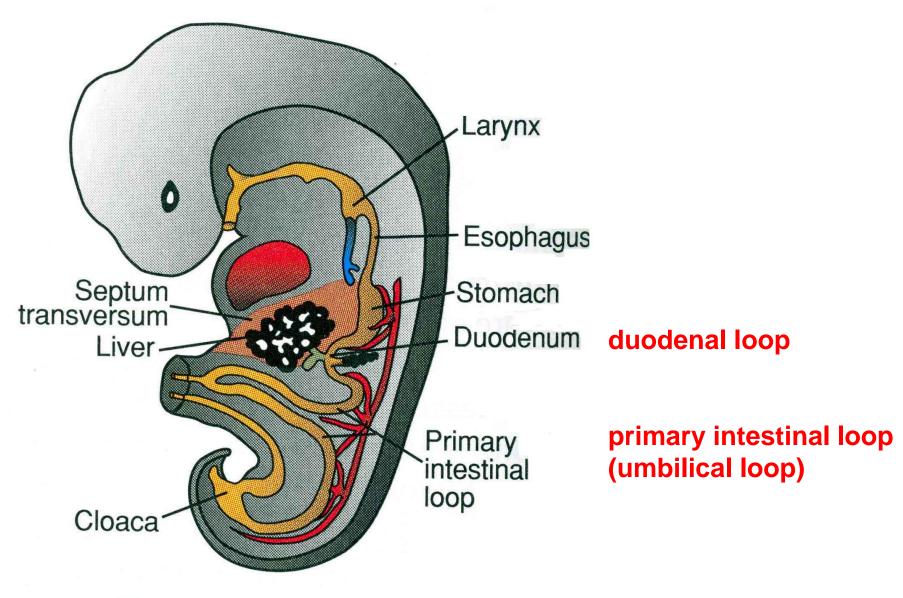
Red letters signaling molecules

blue letters transcription factors



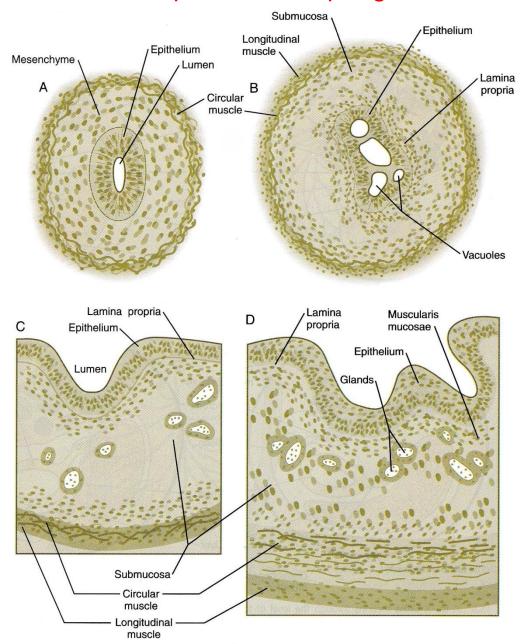




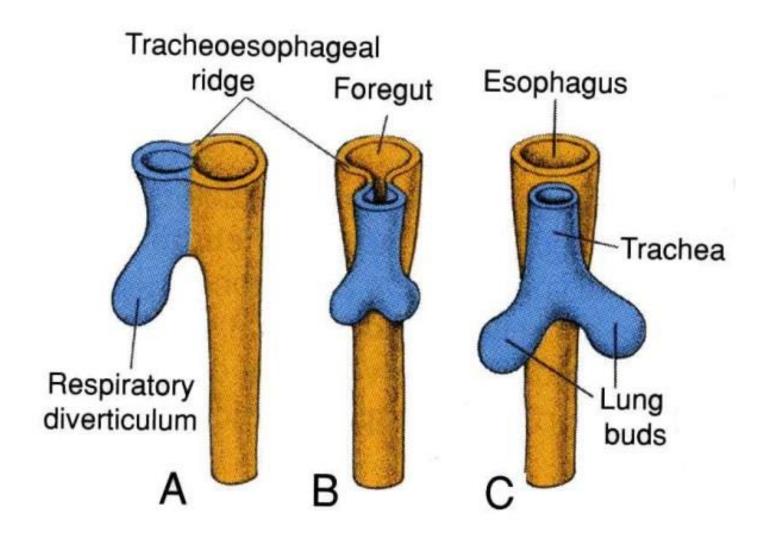


B

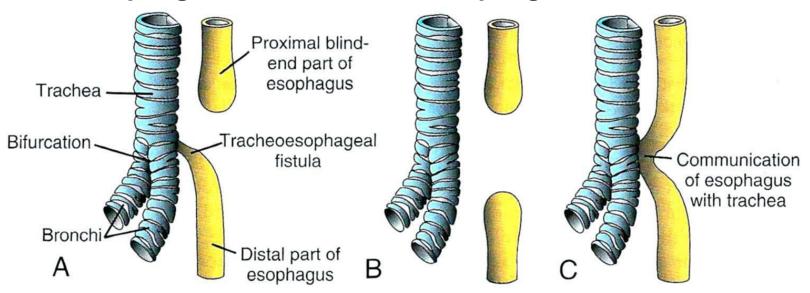
Development of esophagus

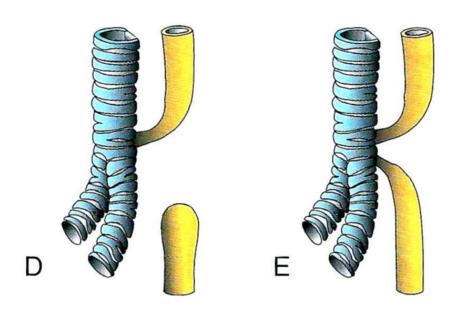


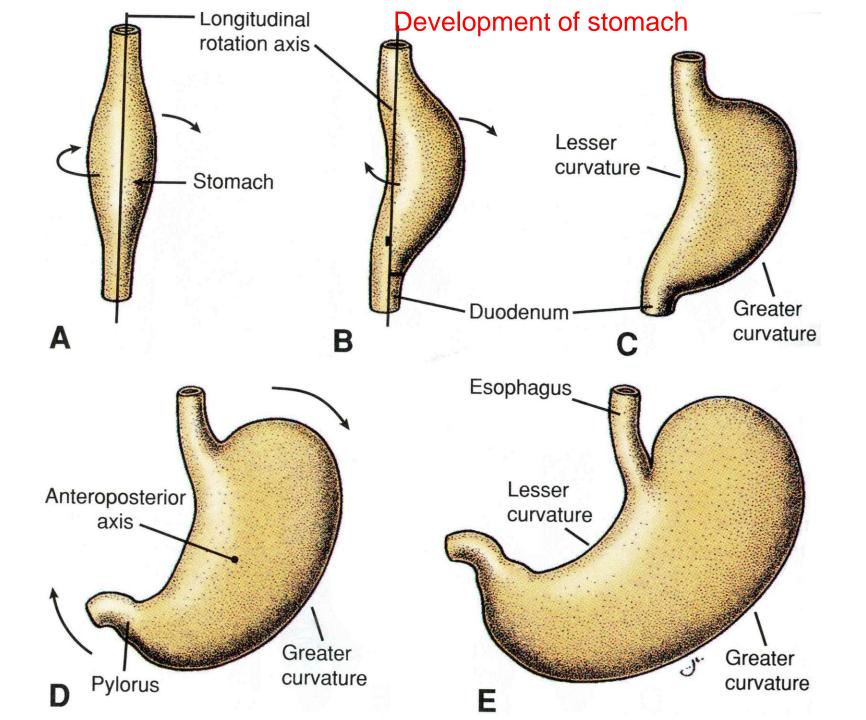
Tracheoesophageal septum



Esophageal atresia, tracheoesophageal fistula







Histogenesis

mucosal secretory cells develop step by step

gastric mucosa forms at the end of the 2nd month (rugae, first gastric pits)

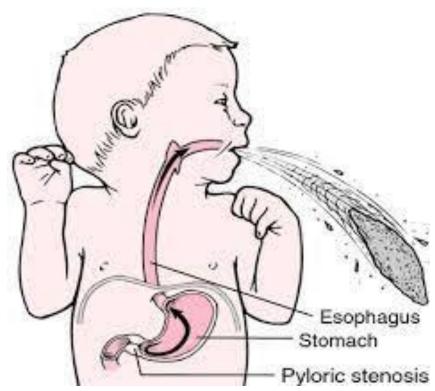
early fetal period – individual cell types featuring gastric mucosa

specific cell types complete their differentiation at late fetal period

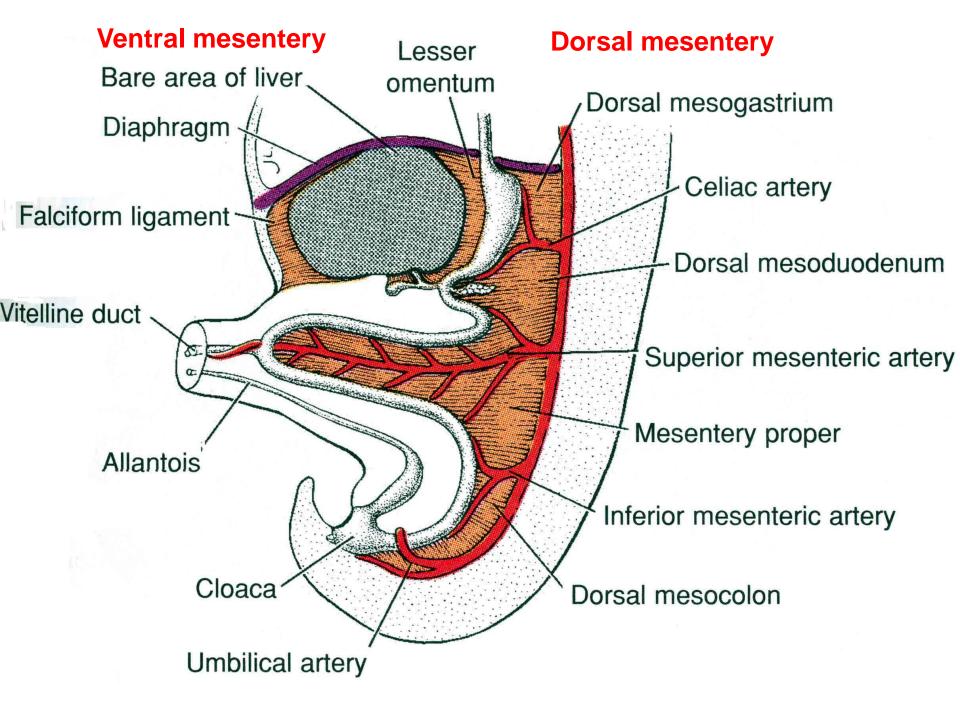
production of HCl early after birth

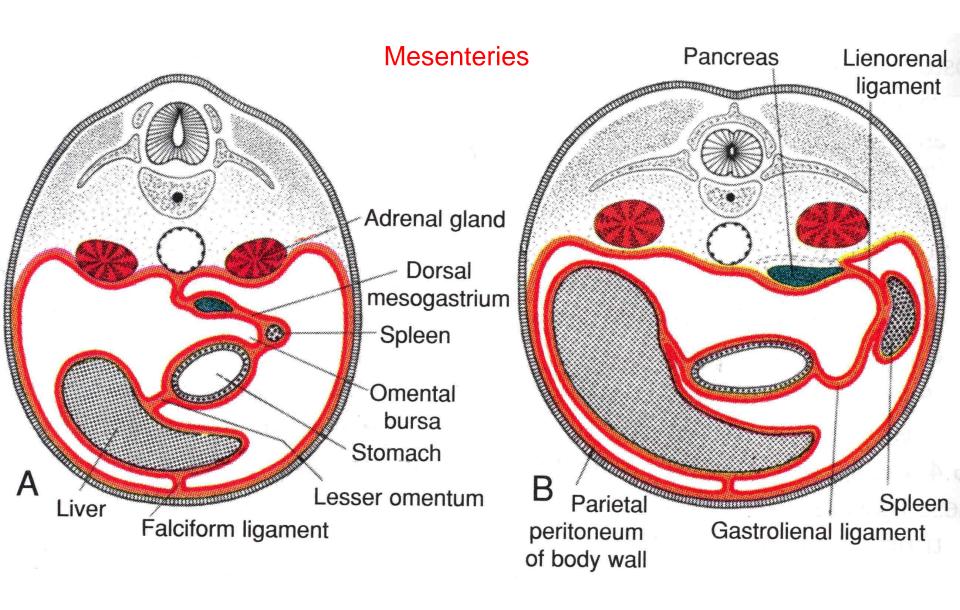
Pyloric stenosis



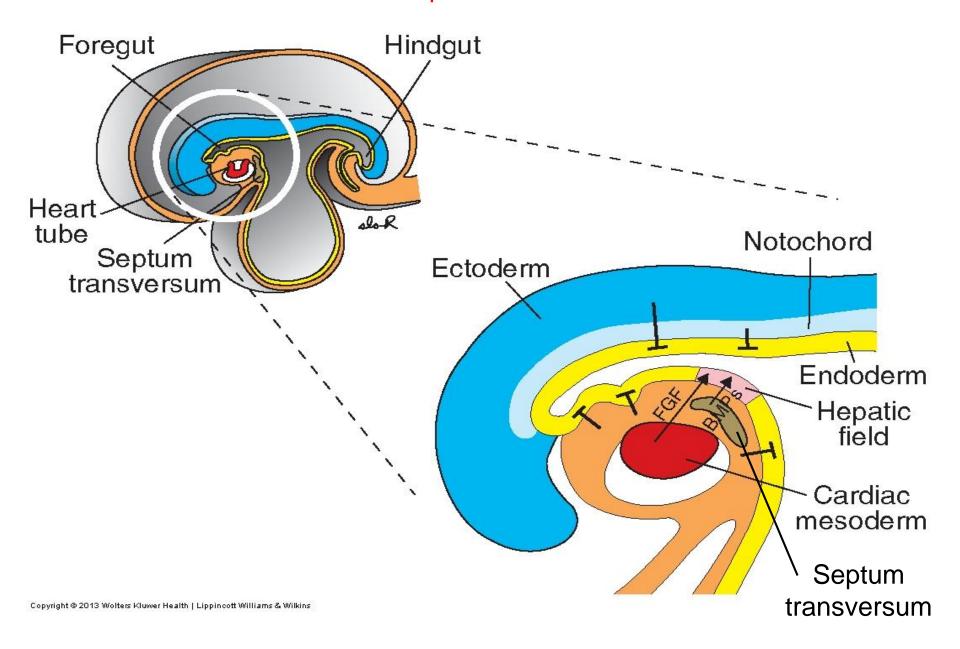


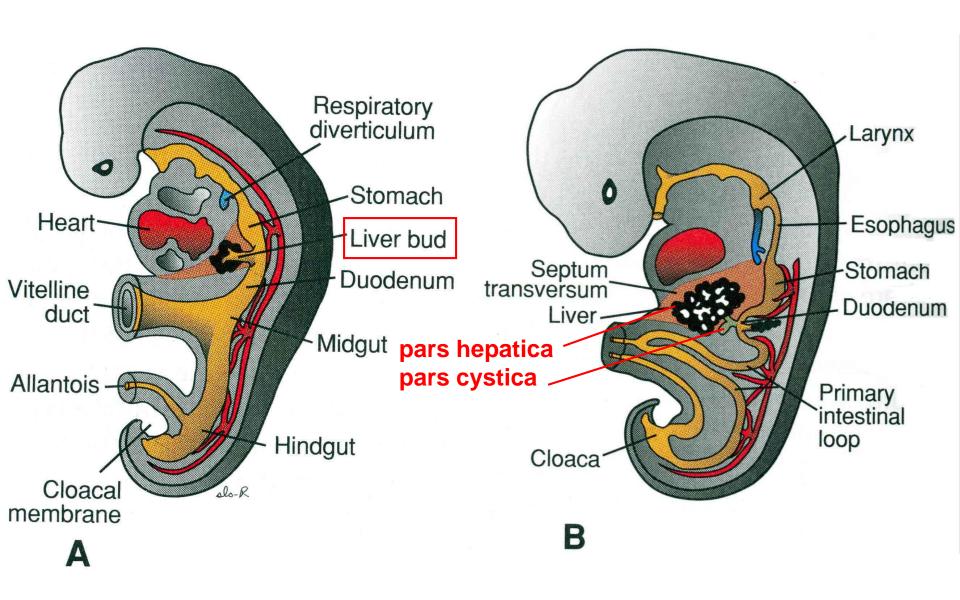
functional defect – hypertrophy of m. sphincter pylori males 1:150, females 1:750 projectile vomiting without the presence of bile

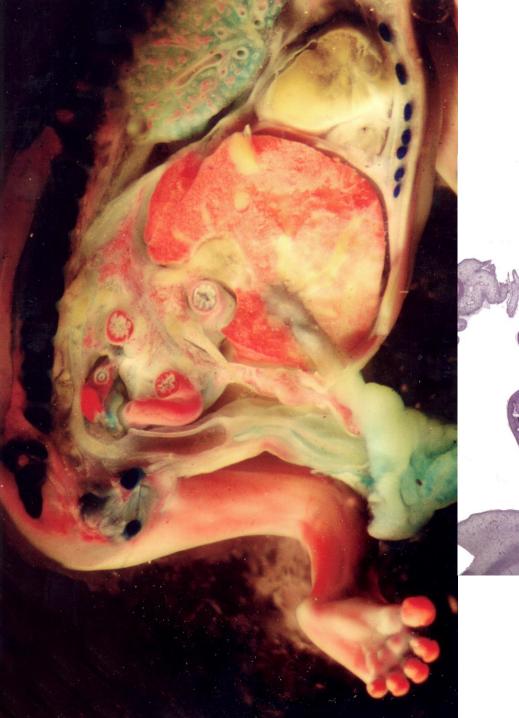




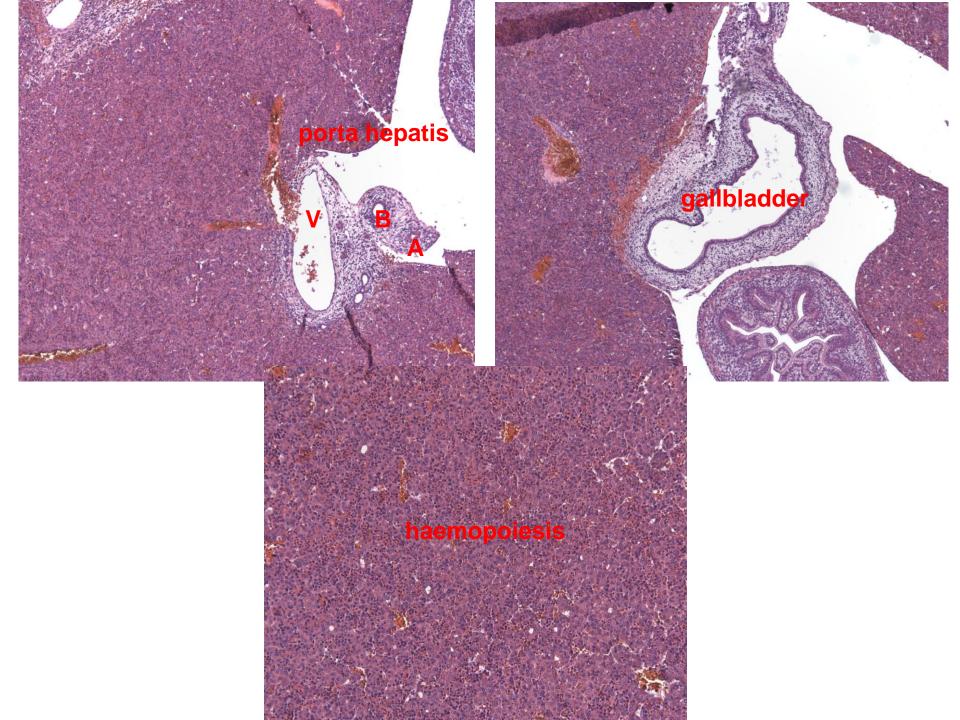
Development of liver



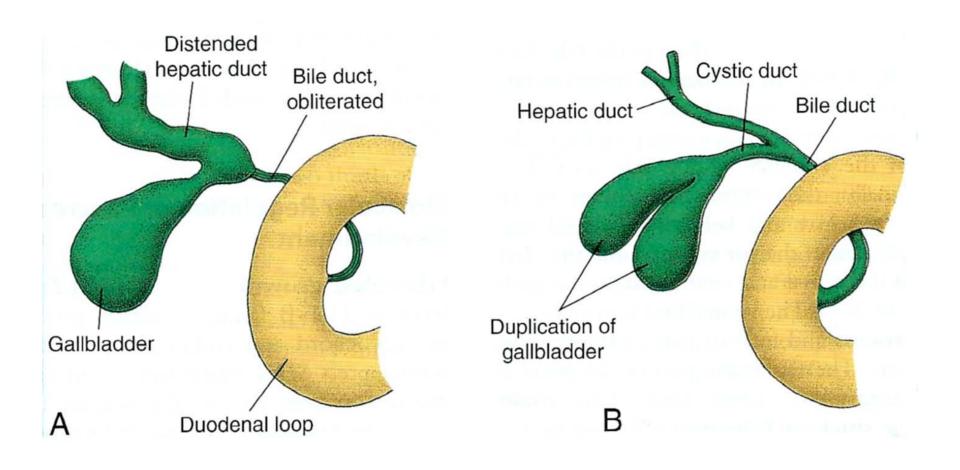


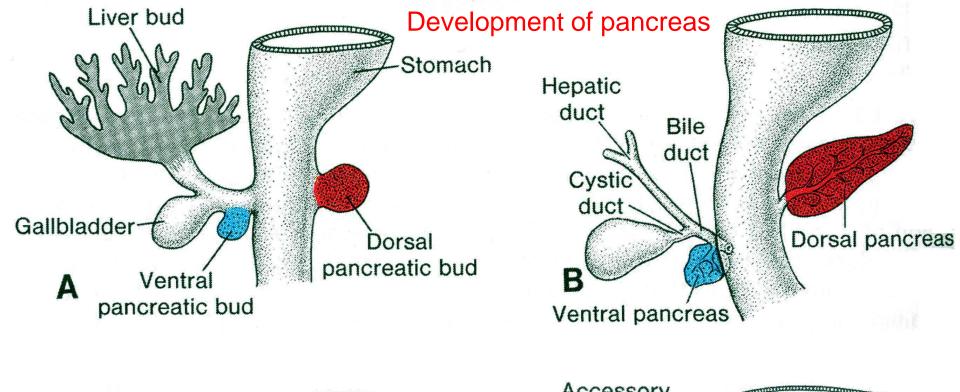


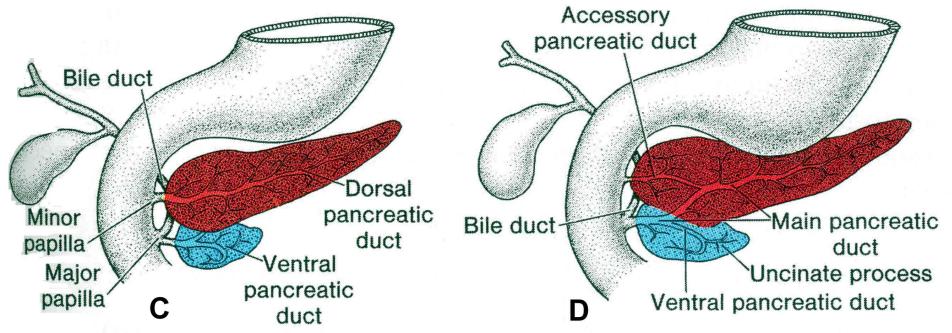


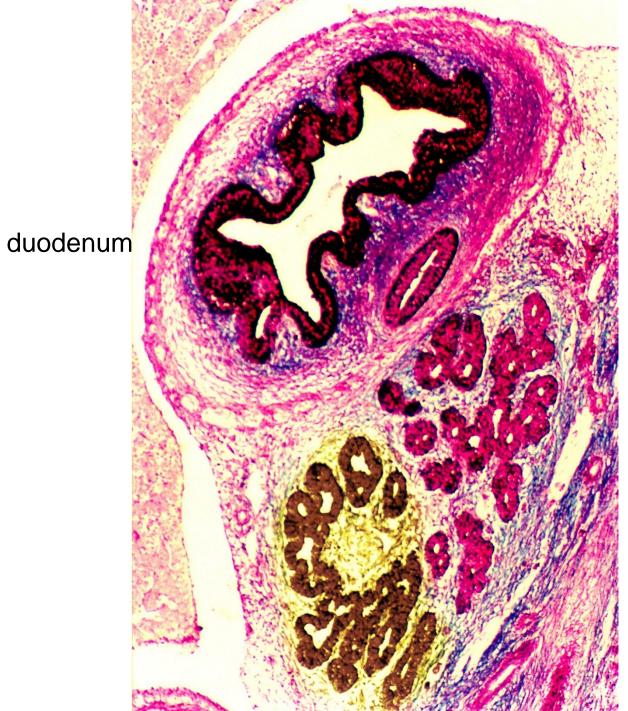


Biliary atresia, duplication of gallbladder









pancreas

Aberrant pancreatic tissue (stomach, duodenum, Meckel's diverticulum)



Fig. 1 Enhanced computed tomography showing a 4×4 cm heterogeneous solid submucosal tumor (*arrowheads*) arising from the posterior wall of the pyloric antrum

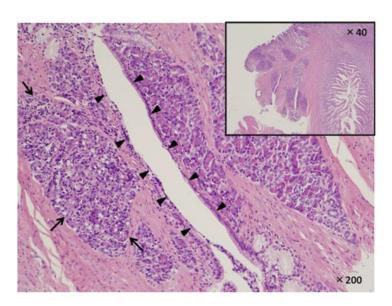
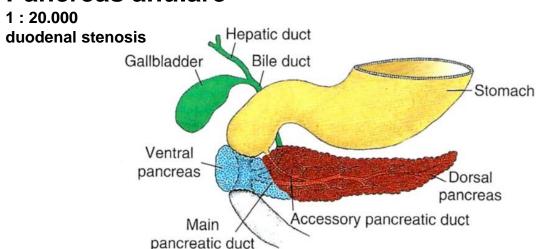


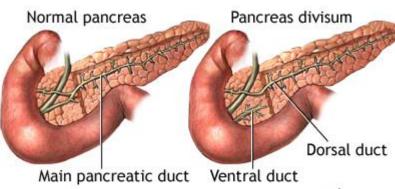
Fig. 2 Histopathology shows aberrant pancreatic tissue (*arrows*) with acini and ductal components (*arrowheads*)

Pancreas anulare



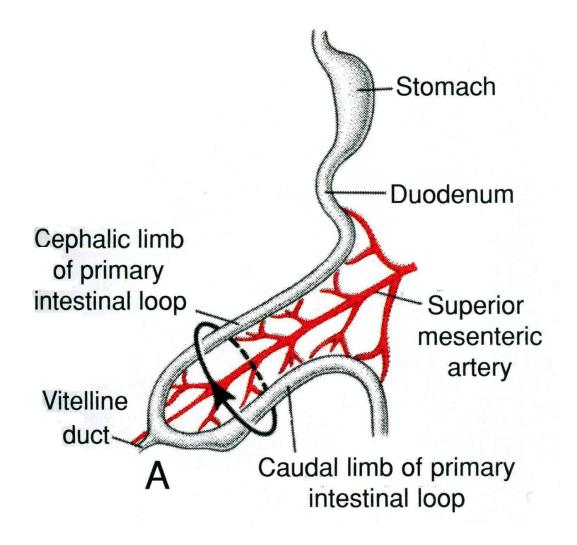
Pancreas divisum

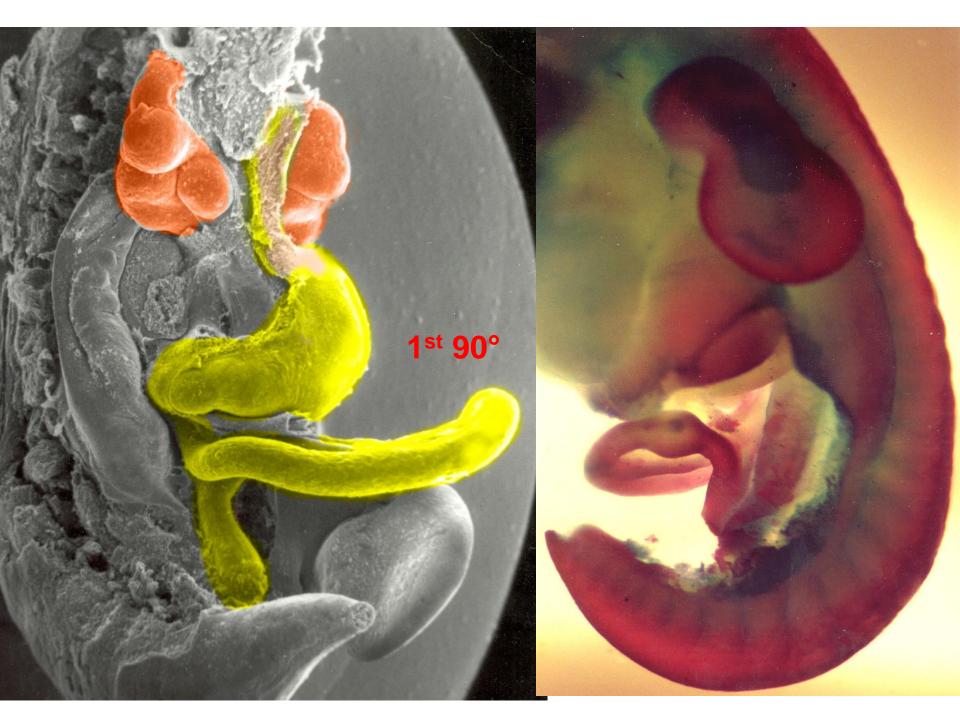
5 – 8 % asymptomatic



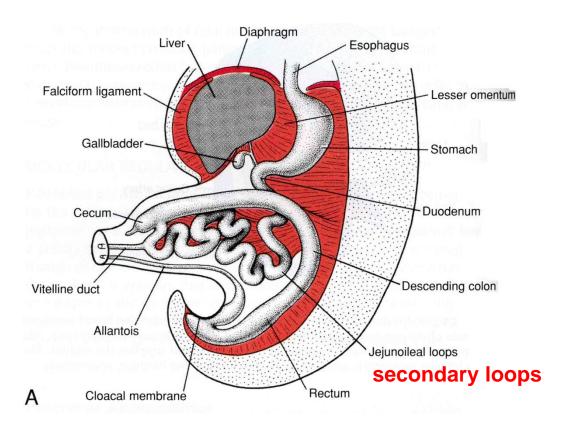


Development of intestines



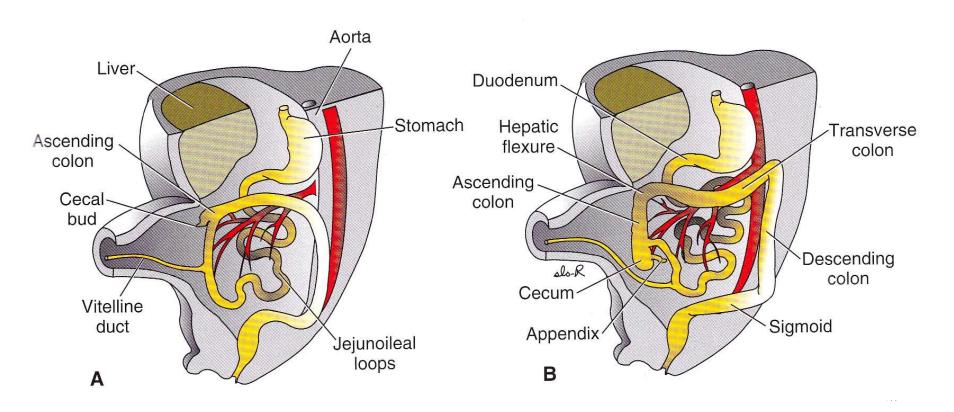


2nd 90°



3rd 90°

cecal bud descent



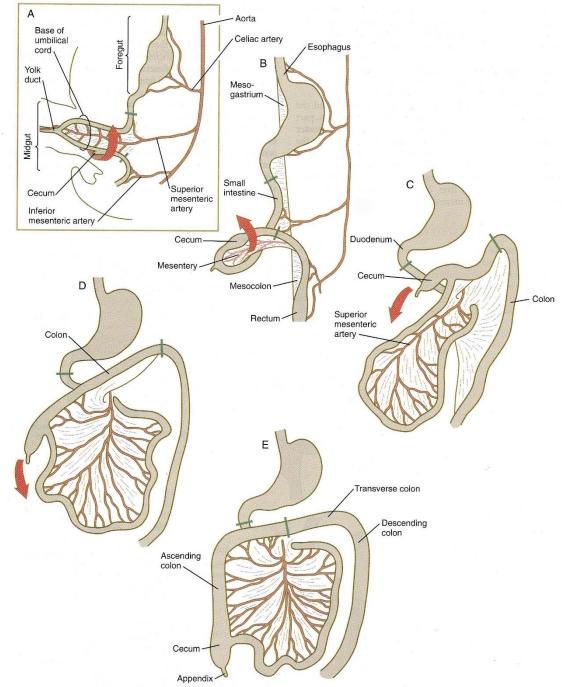
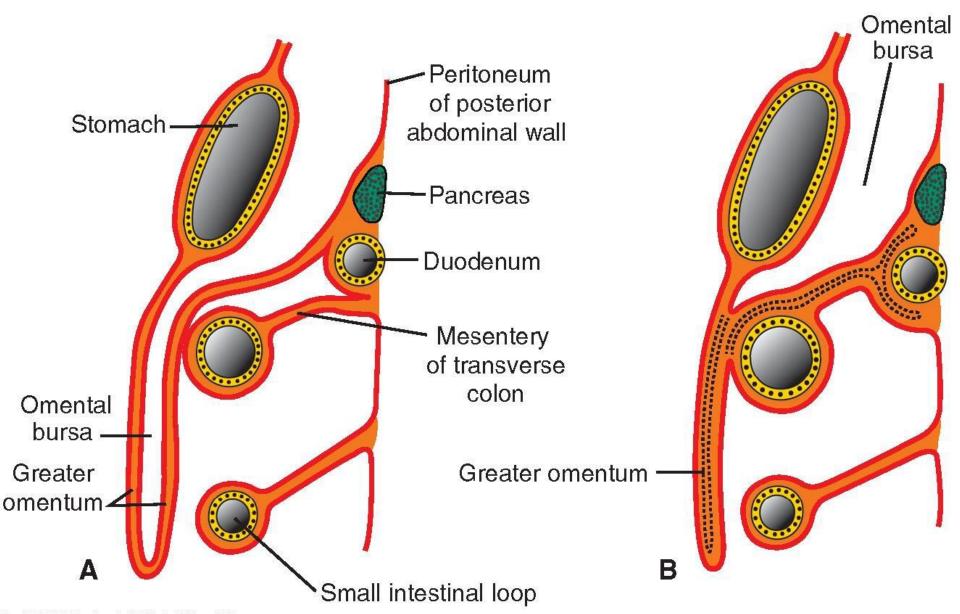
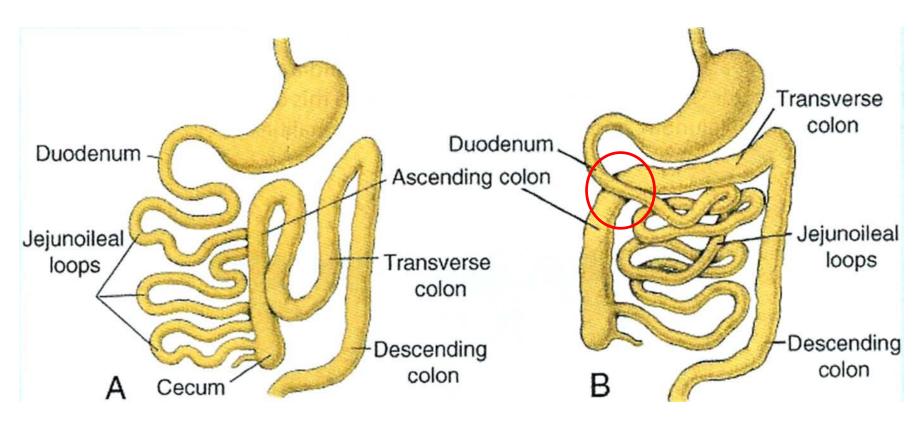


Fig. 15.7 Stages in the development and rotation of the gut. A, At 5 weeks. B, At 6 weeks. C, At 11 weeks. D, At 12 weeks. E, Fetal period. Areas between the *green lines* represent the midgut, which is supplied by the superior mesenteric artery.



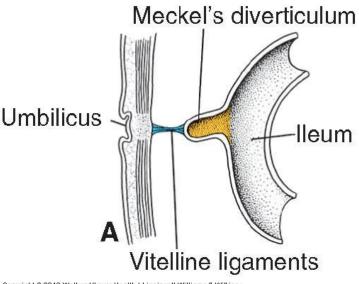
Abnormal rotation of the umbilical loop

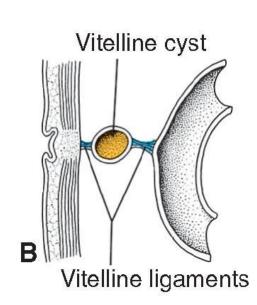


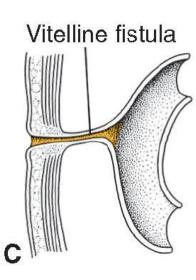
insufficient rotation

reversed rotation

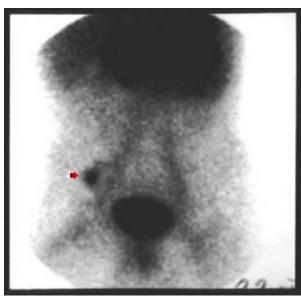
Anomal cloasure of ductus ophaloentericus







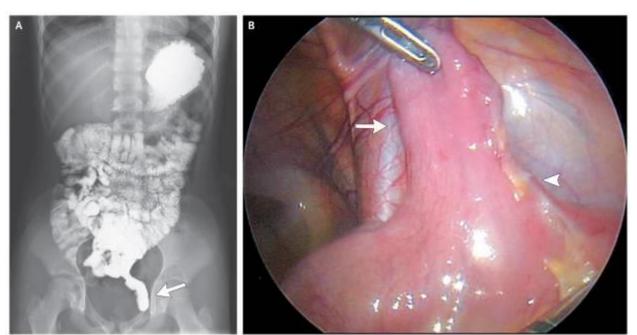
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http://www.surgical-tutor.org.uk/default-home.htm?tutorials/meckels.htm~right

Diverticulum ilei Meckeli

- 2%
- 0-100 cm from ostium ileocaecale



http://www.nejm.org/doi/full/10.1056/NEJMicm1001158



http://www.learningradiology.com/archives2009/COW%20378-Meckels%20Tic/caseoftheweek378page.htm



Histogenesis of intestine

early primitive gut lined with simple columnar epithelium lying on splanchnic mesoderm

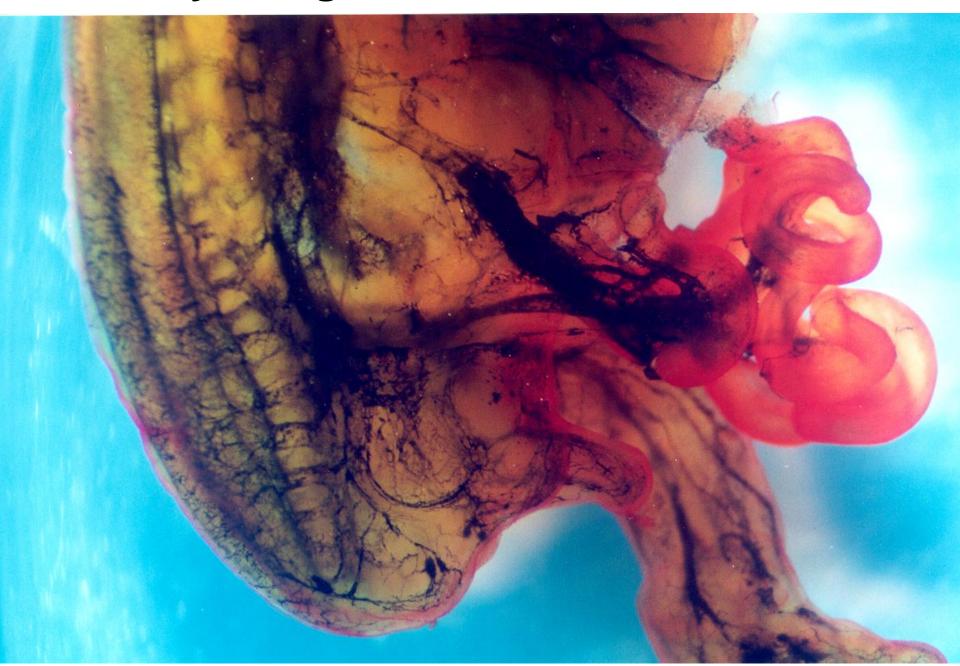
fast proliferation of epithelium temporarily obliterates lumen (6th – 7th week)

during several weeks, recanalization proceeds to form definitive intestinal lumen – lack of recanalization can cause atresia or stenosis

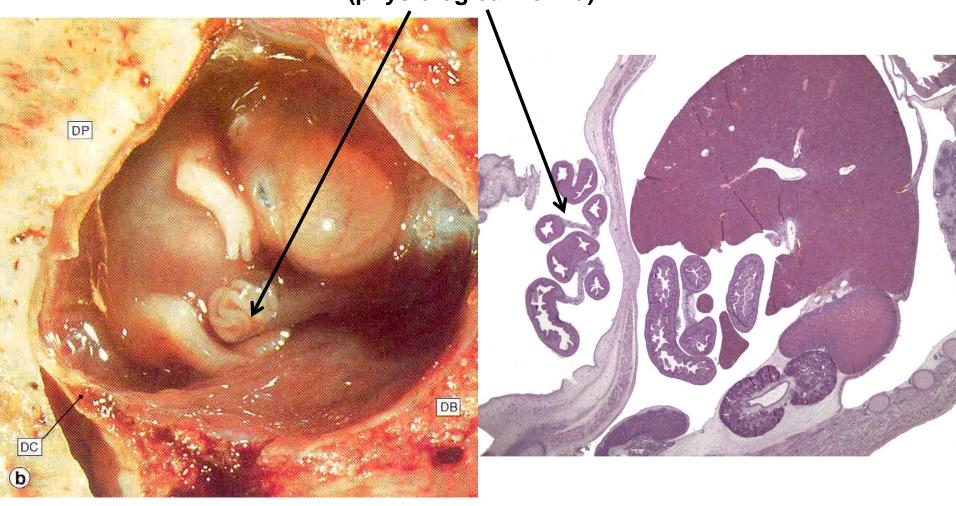
small secondary lumina lined with stratified epithelium small aggregates of mesoderm within epithelium as a background of villus stroma intestinal villi develop lined with simple columnar epithelium crypts develop lined with intestinal stem cells

goblet cells and enteroendocrine cells in the second trimester

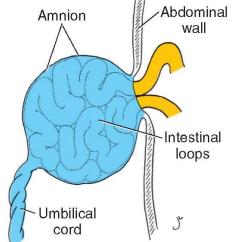
Physiological umbilical herniation



intestinal loops in the umbilical coelom (physiological hernia)



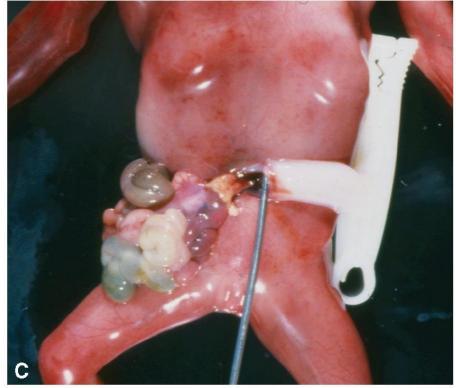
Omphalocele





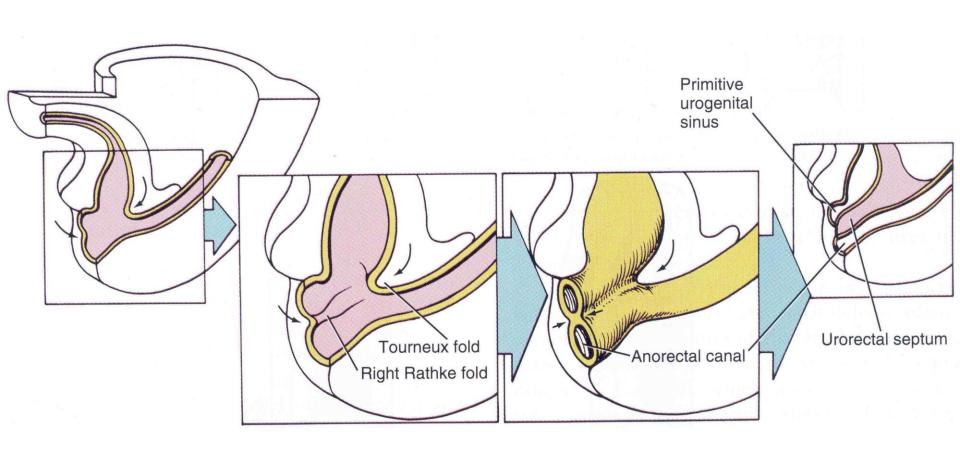
Gastroschisis





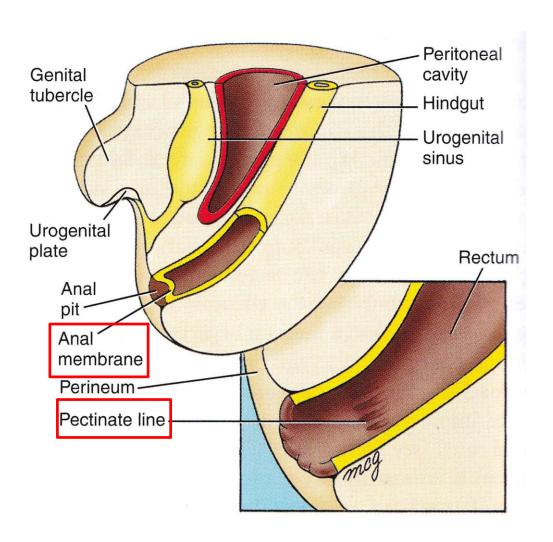
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Cloaca, formation of urorectal septum



4th week 5th week 6th week 7th week

8th week

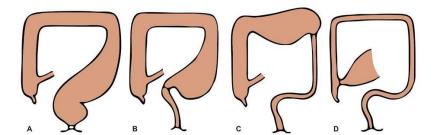


fetus

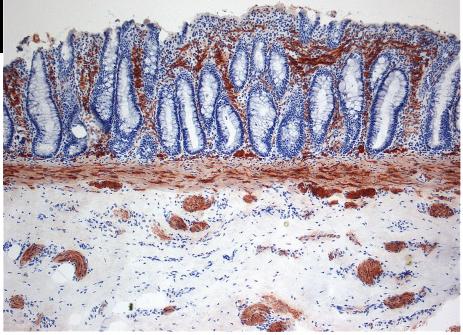
Megacolon congenitum (Hirschprung's disease)



Hirschprung's disease (types)

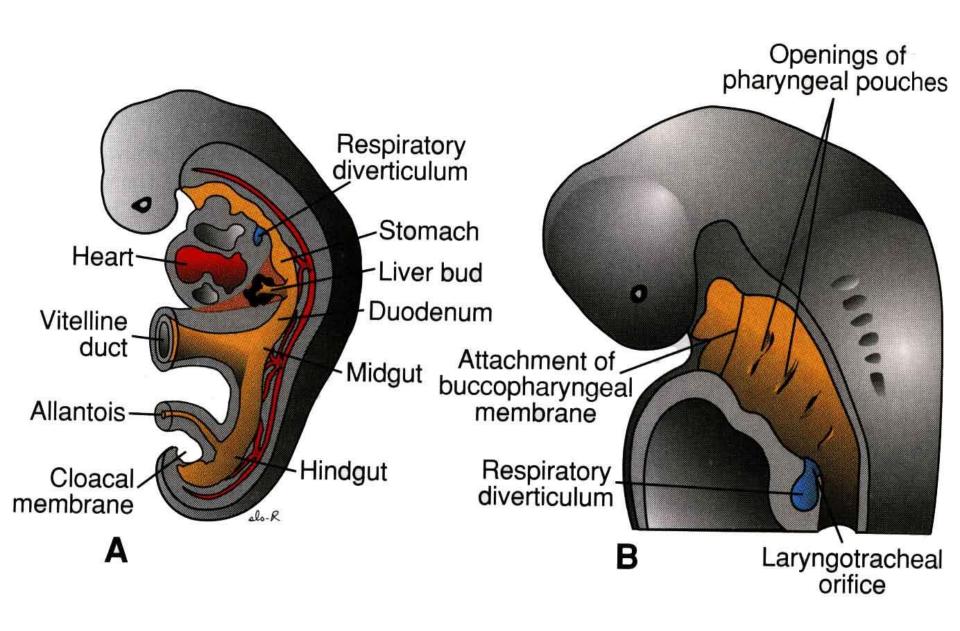


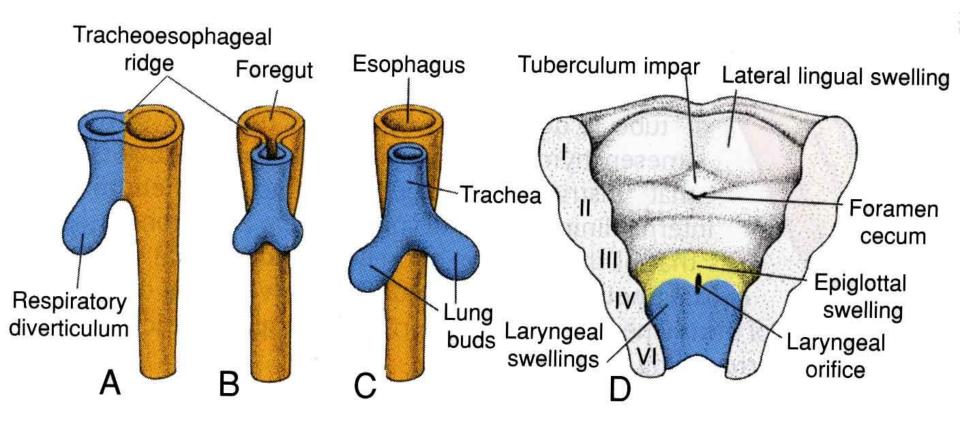
- inborn agangliosis of large intestine
- defect of neuroblast migration from neural crest
- multiplication of atypical nerve endings with acetylcholinesterase activity

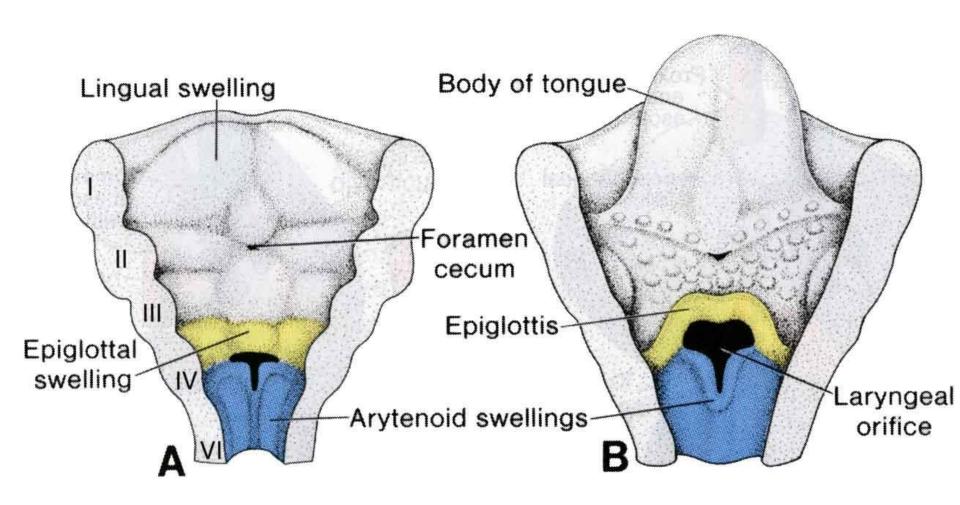


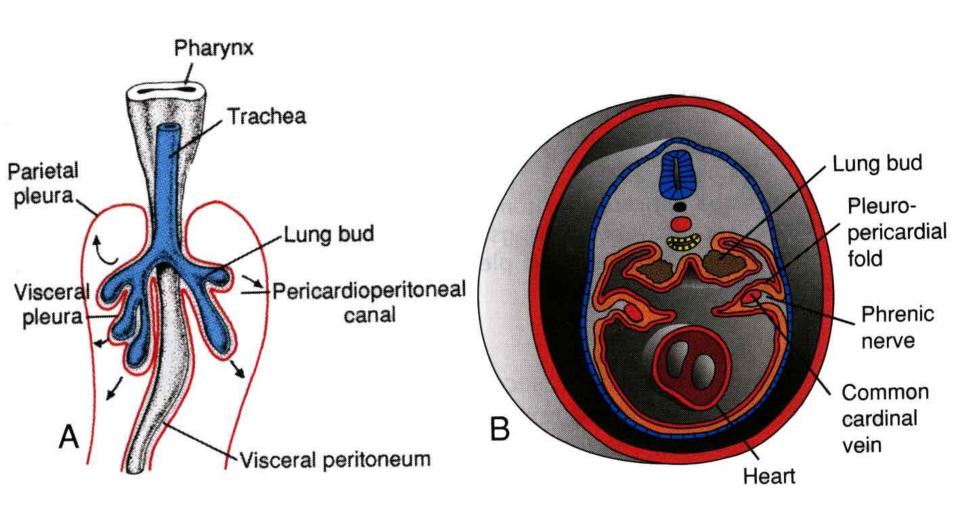
proof of acetylcholinesterase

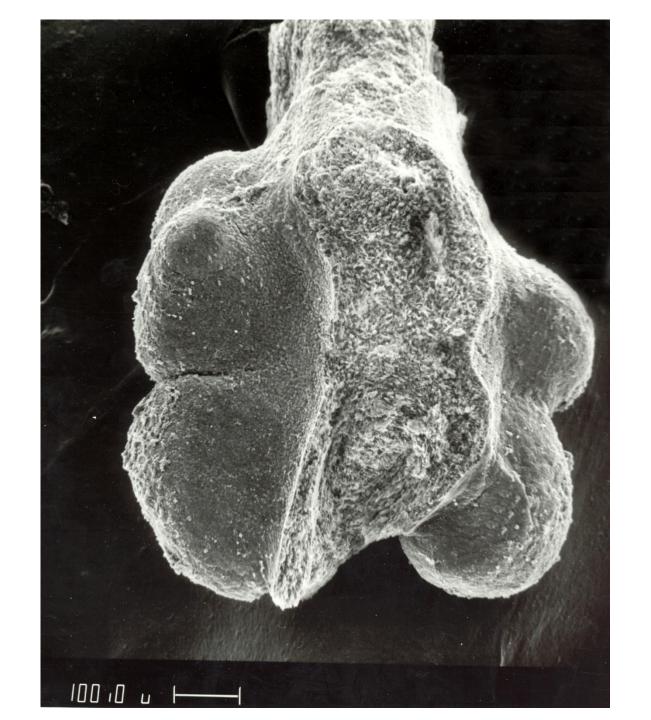
DEVELOPMENT OF THE RESPIRATORY SYSTEM

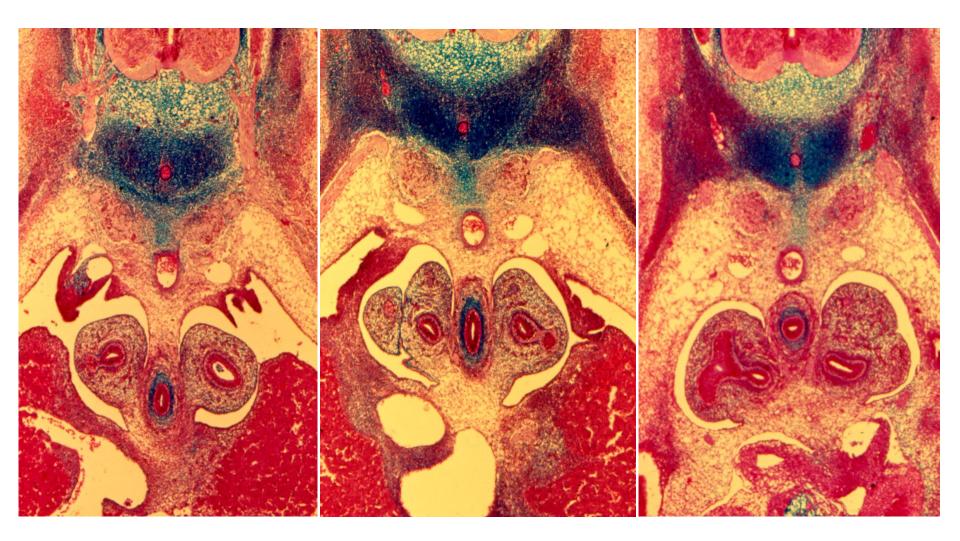


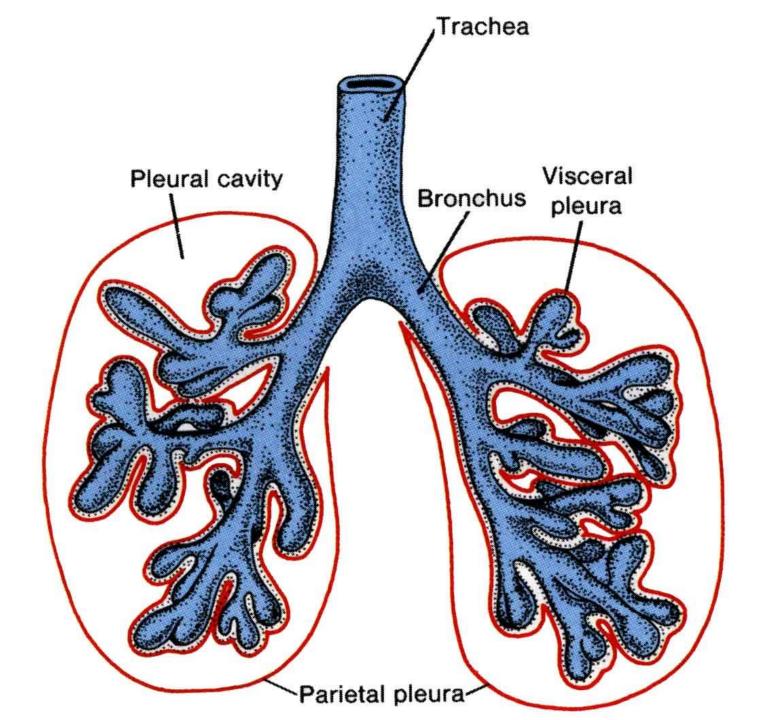


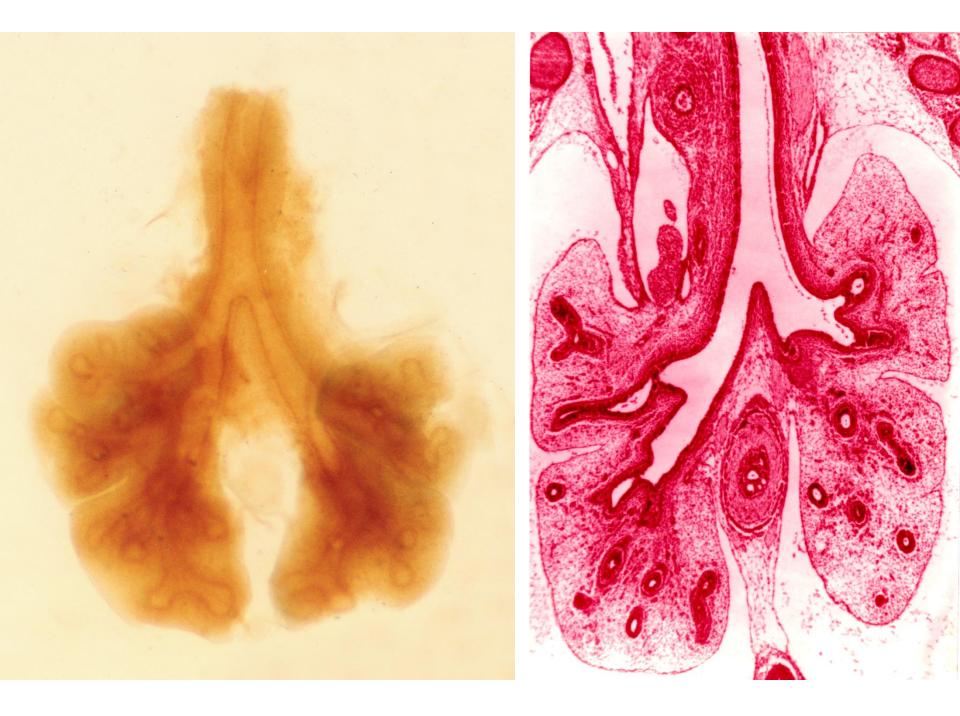








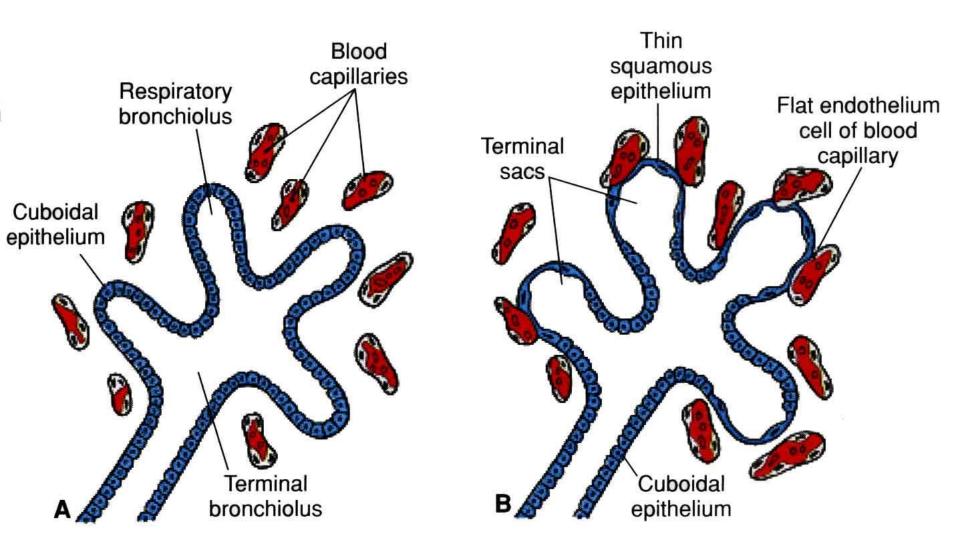


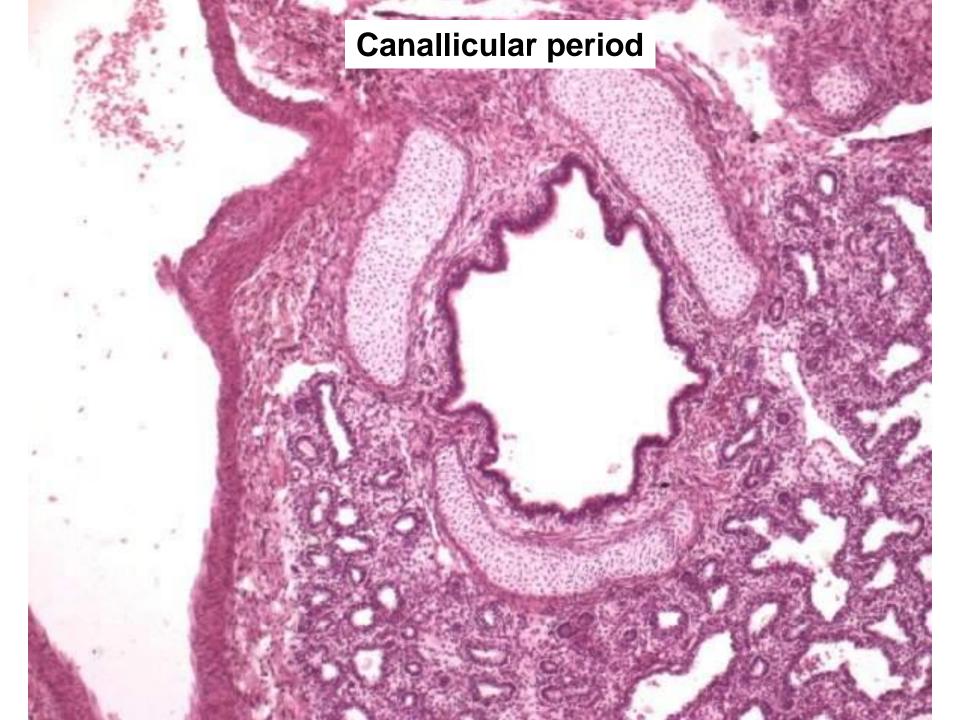


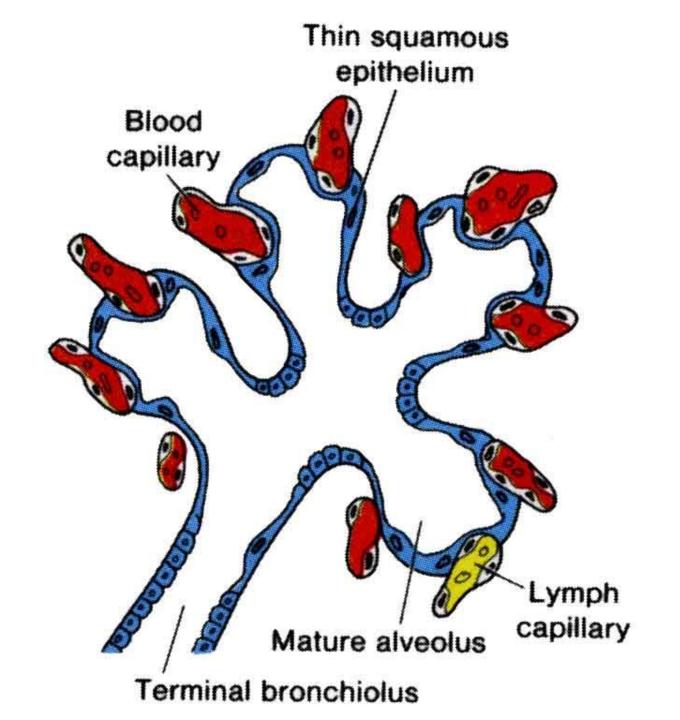
Maturation of the Lungs

Pseudoglandular period	5-16 weeks	Branching has continued to form terminal bronchioles. No respiratory bronchioles or alveoli are present.
Canalicular period	16-26 weeks	Each terminal bronchiole divides into 2 or more respiratory bronchioles, which in turn divide into 3-6 alveolar ducts.
Terminal sac period	26 weeks to birth	Terminal sacs (primitive alveoli) form, and capillaries establish close contact.
Alveolar period	8 months to childhood	Mature alveoli have well-developed epithelial endothelial (capillary) contacts.

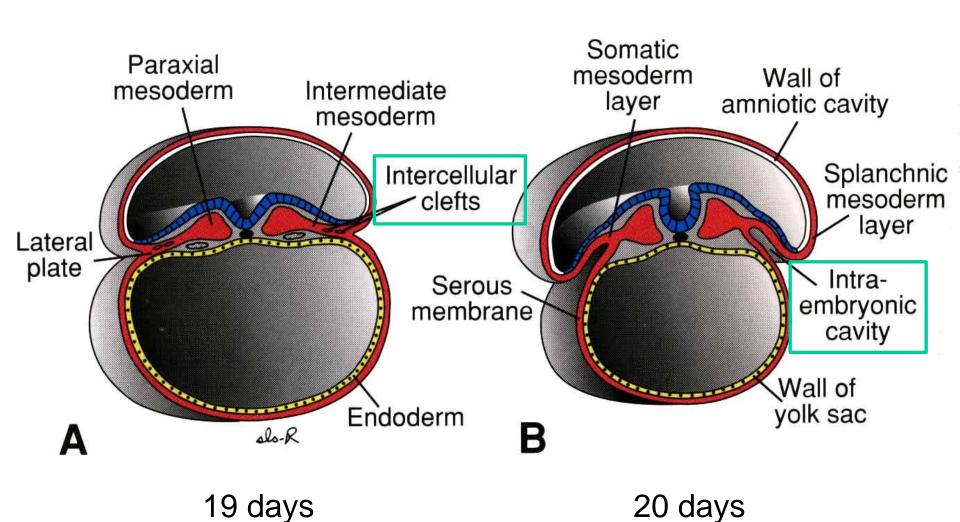


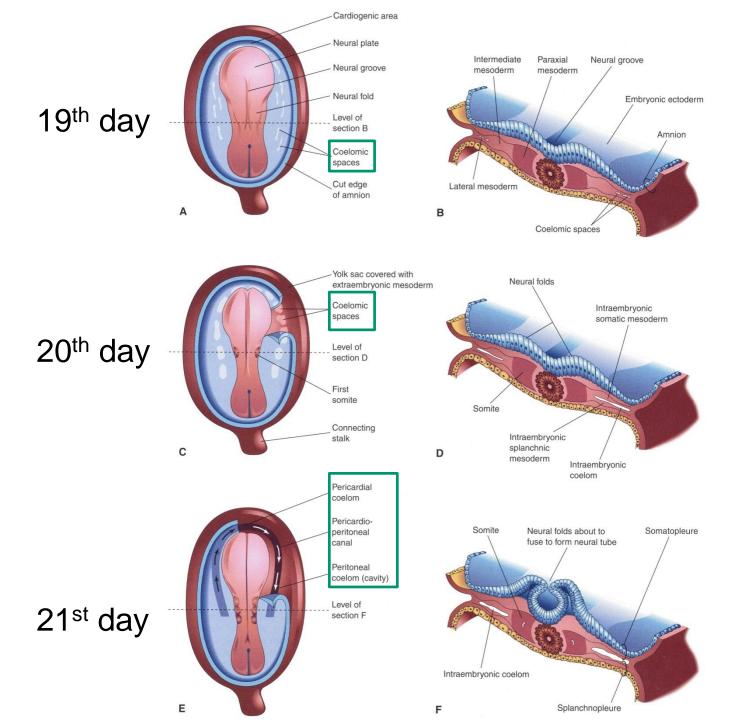




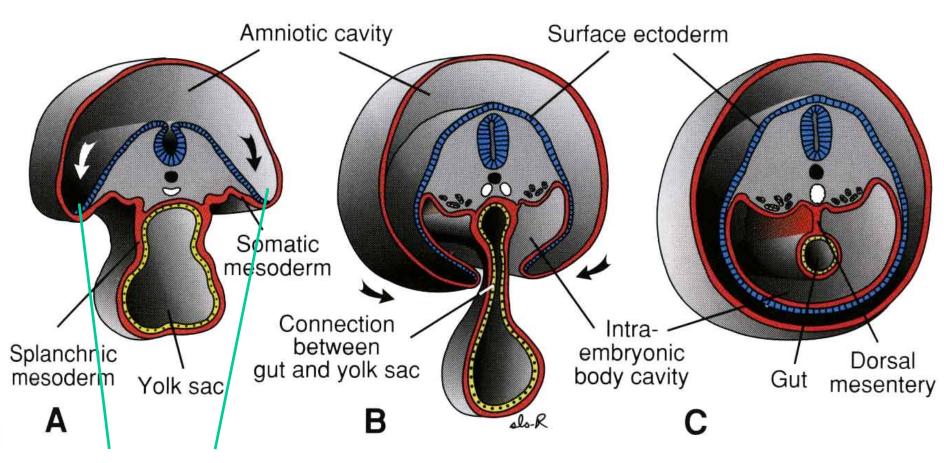


DEVELOPMENT OF THE COELOM AND DIAPHRAGM





Lateral folding (rolling up)

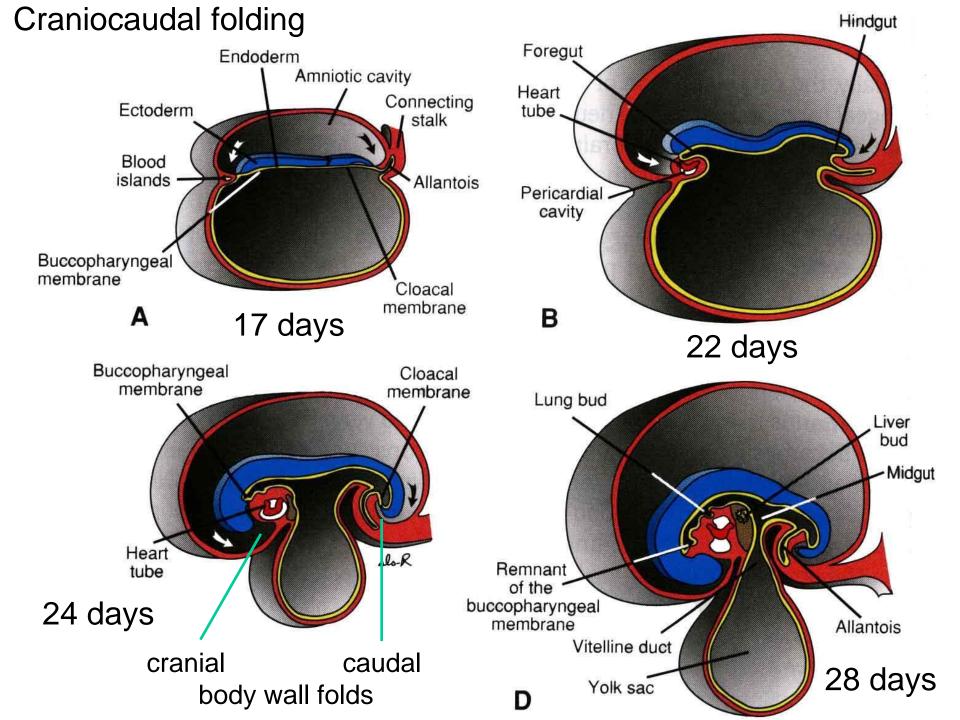


lateral body wall folds

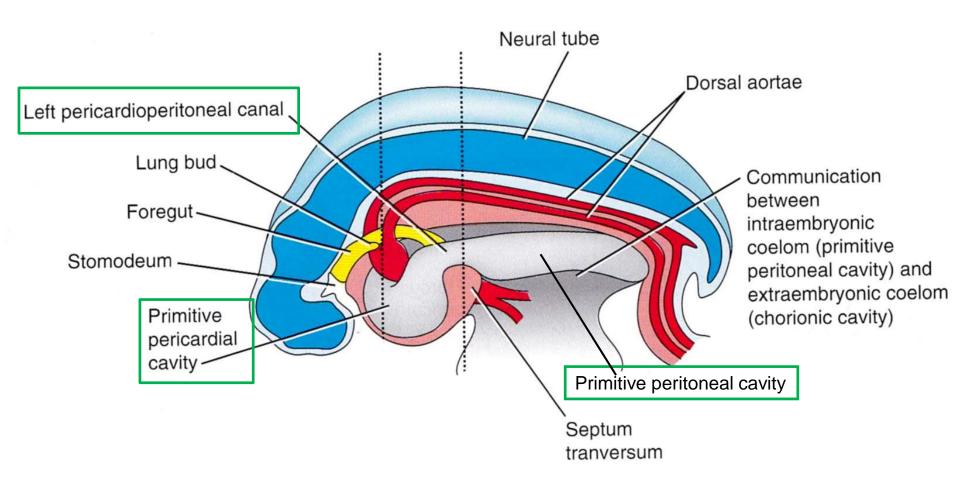
21 days

24 days

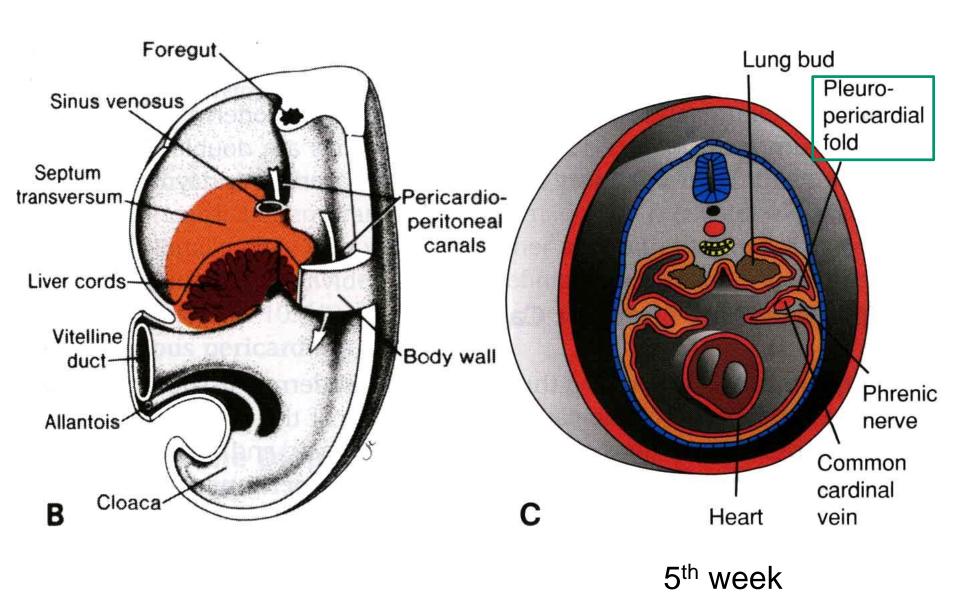
28 days

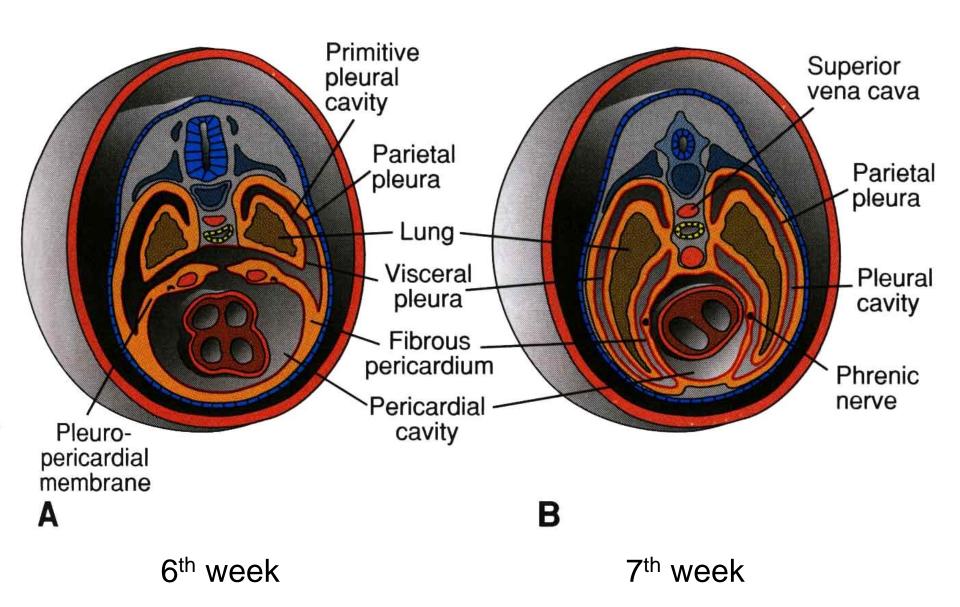


Intraembryonic coelom (24th day)

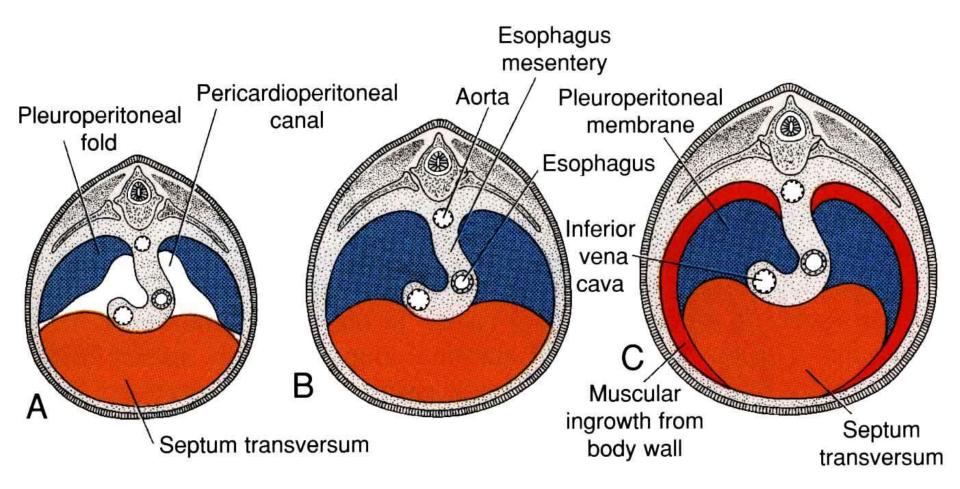


Separation of pericardial cavity from pericardoperitoneal canals





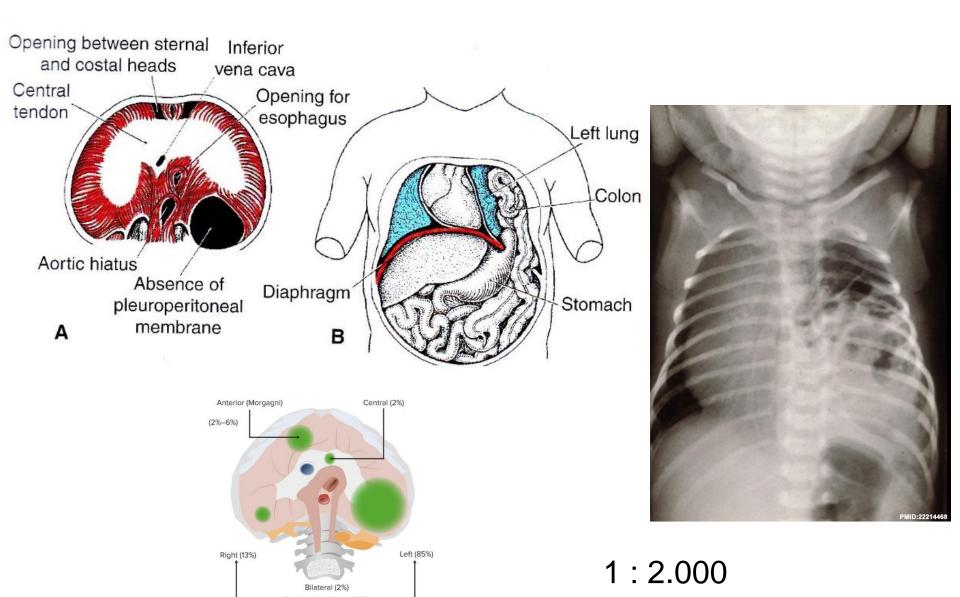
Separation of peritoneal cavity from pericardoperitoneal canals



4 anlages:

- pleuroperitoneal folds (somatopleura)
- septum transversum (splanchnopleura)
- mesentery (splanchnopleura)
- abaxial muscles of cervical somites via the body wall

Congenital diaphragmatic hernia



(Bochdalek) hernias