DEV. OF INT., RECTUM & ANAL CANAL



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embryonic disc, about 16 days after fertilization

REV.



REV.



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

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

Site:-Connect the gut to anterior abdominal wall extent:- from lower end of the esophagus to 1st 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 1st inch Mesentery proper:-opposite the jejunum &ileum will form the mesentery mesocolon :- opposite the colon will disappear except transverse mesocolon,

sigmoid mesocolon, mesoappendix



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 and opening of bile duct in second part of duodenum
- □ 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
- loop rotate 90 o to Rt due to rotation of stomach convexity become to Rt
- □ Fixation: Mesoduodenum degenerate
- □ duodenum become retroperitoneal except 1st inch



DUODENUM

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





Extent:

from ant. intestinal portal

- (site of liver bud)
- (in adult opening of CBD in duodenum)
- to post. intestinal portal
- (In adult junction of Rt 2/3 & Lt 1/3 of transverse colon). $\frac{Cloacal}{membrane}$
- it is connected with yolk sac by vitelline (vitellointestinal) duct **Derivatives (fate):**
- It gives
- lower half of duodenum caudal to opening of CBD_{Duodenum} -jejunum, ilium, appendix, caecum,

ascending colon & Rt 2/3 of tr. colon





Hindaut

Development:

□ primary midgut (intestinal) loop:

midgut grows rapidly in length formation of

- U shaped loop that is formed of
- 1- cranial limb
- 2- apex connected to vitelline duct
- 3- caudal limb with swelling (close to the apex) that will form caecum

4- superior mesenteric art. along its axis(in its dorsal mesentery)

physiological umbilical hernia:

-at 6th week of dev., the rapidly elongating loop herniate into umbilical cord through umbilical orifice

- -herniation is due to inability of abd. cavity
- to accommodate rapidly growing mid gut due to
- 1-slow growth of abd. cavity
- 2-development of liver & kidney



MIDGUT **Development:** □ While the loop in the umbilical cord -The cranial limb form the lower 1/2 of duodenum, jejunum and greater part of ileum -The caudal limb also form the distal part of ileum, caecum, appendix, ascending colon and Rt 2/3 of transverse colon



- **Development:**
- □ rotation of the intestinal loop
- -Due to further elongation of loop.
- -Total 270 anticlockwise

around its long axis formed by SMA.

- -While in umbilical cord, it rotates:
- 90→ caudal limb become to Lt & cranial limb become to Rt., then rotate
- 90 →caudal limb become cranial & cranial limb become caudal
- As the gut returns to abd cavity, it rotates additional
- 90 → the caudal limb (tr. Colon) become to Rt &crosses (become superficial)
- to the cranial limb (2nd part of duodenum)







MIDGUT Development:

return of the loop:

- -at 10^{th} week of dev., the abd. cavity become wide enough to accommodate the intestine _____ return of physiological hernia
- -jejunum is the 1st part to return
- into abd cavity &lies on the Lt side
- -caecum is the last part to return
- into abd cavity & lies with the appendix
- on the Rt side, below and in contact with Rt lobe of liver.
- □ caecum and appendix descend
- to Rt iliac fossa due to elongation
- of the segment () caecum & tr. Colon
- to form Rt colic flexure and ascending colon





Development

□ change the site of opening of appendix

- from apex of caecum to
- its posteromedial wall
- by differential growth
- □ Fixation of intestine:
- The mesentery of duodenum,
 ascending colon and descending colon
 fuse with peritoneum of post. abd wall &
 these organs become retroperitoneal
 The other mesenteries persist
- □ At 2nd month the vitelline duct
- is obliterated,
- fibrosed and degenerate



- **Congenital Anomalies:**
- of intestine:
- A-of intestinal loop
- <u>**1- atresia**</u> (due to failure of recanalization)
- **<u>2-stenosis</u>** (due to defect in recanalization)
- **<u>3-Diverticulosus</u>**:- due to week wall
- **4-Duplication** is common in **ileum**
- **B- congenital umbilical hernia (omphalocele)**
- -herniation of intestinal loop into umbilical cord
- -due to failure of return of
- physiological umbilical hernia
- or wide umbilical orifice



- MIDGUT Congenital Anomalies:
- of intestine:
- **C- of rotation:**
- **<u>1- excessive rotation more than 270</u>**
- leads to congenital volvulus
- 2-incomplete rotation 90 anticlockwise only:
- caecum & colon become in Lt side of abdomen while
- duodenum, jejunum and ileum become in Rt side
- **3-reversed rotation, 90 in reverse direction (clockwise)**
- duodenum lies in front transverse colon



Volvulus

- **Congenital Anomalies:**
- of intestine:
- **D- of vitelline duct**

<u>1- vitelline (umbilical faecal) fistula:</u>

- -due to persistence of vitelline duct
- with faecal discharge at the umbilicus

2- Meckel's diverticulum:

- -due to persistence of the proximal part of vitelline duct
- -it has the following features:
- \Box in 2% of people , 2 inches (5 Cm) long ,2 feet from ileocaecal valve
- $\hfill\square$ Attach to ant mesenteric border of ileum
- $\hfill\square$ Attached to umbilicus by a fibrous cord
- $\hfill\square$ May contain ectopic gastric or pancreatic tissue
- \Box May cause pain confused with the pain from appendicitis



Vitelline fistula



Meckel's diverticulum

- **Congenital Anomalies:**
- of intestine:
- **D- of vitelline duct**
- <u>**3-vitelline sinus:**</u> due to persistence of distal part of vitelline duct
- **<u>4-vitelline cyst:</u>** due to persistence of middle part of vitelline duct

5-fibrous cord:

- -due to failure of degeneration of
- the obliterated, fibrosed vitelline duct
- -a loop of intestine may become wrapped around it causing intestinal obstruction



- **Congenital Anomalies:**
- of caecum and appendix:
- **1- Abnormal position:**
- e.g. sub hepatic, or Rt lumbar caecum and appendix:
- Due to failure of descend or arrest during descend
- **2-** Retention of fetal shape with apical appendix.





HINDGUT

DEF.:-

the part of the primitive gut which is enclosed in the tail fold of the embryo.

Extent:

- It extends from the posterior intestinal portal
- until the cloacal membrane.
- Fate(derivatives):
- -left 1/3 of transverse colon.
- -left colic flexure, descending colon ,
 sigmoid colon, rectum.
 -upper 1 /2 of anal canal.
 - Rectum Sigmoid colon

siverae colors

Descending color



RECTUM

Development source:

endodermal cloaca of hindgut.

<u>allantois:</u>

- diverticulum projecting ventrally from hindgut into umbilical cord endodermal cloaca:
- dilatation in hindgut just distal to origin of allantois,
- closed caudally by
- cloacal membrane
- (that separate the cavity of
- hind gut from the surface)



RECTUM Development:

□ the mesoderm at the angle () hindgut and allantois proliferate and invaginate the endoderm forming urorectal septum that grows inferiorly in a coronal plane dividing the endodermal cloaca into:

-primitive recto anal canal (dorsal)

Form rectum

upper part of anal canal.

-primitive urogenital sinus (ventral)

Form urinary bladder & urethra vagina.



- -anal membrane (dorsal) -urogenital membrane (ventral) primitive perineum (at site of fusion)
- $\hfill\square$ the muscle of rectum develop from surrounding mesoderm



ANAL CANAL Development:

upper half: from rectoanal canal (endodermal)

- This part of anal canal is lined
- by a mucous membrane.
- **lower half:** from proctodeum (ectodermal) as follows:
- mesoderm around anal membrane
- proliferate to form anal tubercles,
- and by the 9th week of development,
- the anal membrane comes to lie
- at the bottom of a depression called the proctodeum.
- This part of anal canal is lined by stratified squamous epithelium (skin).



ANAL CANAL Development:

- □ rupture of anal membrane
- results in continuity()upper & lower 1/2s
- N.B: remnants of anal membrane
- are represented in adult by
- anal valves & pectinate line
- The two parts of anal canal differs
- in A.S., N. S.& V.D., L.D..
- □ muscles of anal canal
- develop from surrounding mesoderm.
- In the upper part form
- int. anal sphincter (involuntary), while
- In the lower part form
- ext. anal sphincter (voluntary)



ANAL CANAL Congenital anomalies of rectum and anal canal:

1-rectal atresia: obliteration of lower part of rectum

2-rectal fistulae: recto vesical,

recto urethral,

recto vaginal fistulae

- communication () rectum and either the urinary bladder, urethra or vagina
- due to incomplete growth of urorectal septum

3-imperforate anus:

- anal membrane fails to rupture
- and persist as a diaphragm stretching across the anal canal at level of anal valves





ANAL CANAL

Congenital anomalies of rectum and anal canal:

4-primary megacolon

(Hirchsprung's disease, aganglionic colon)-in the 1st few days after birth ,the child fails to pass meconiumand the abdomen become distended

- Rectum & anal canal are constricted
- &sigmoid colon is greatly distended
 Due to failure of migration of neural crest cells from neural folds
 to form parasympathetic ganglia
 in wall of bowel



