




- Development of the mucous membrane of the anterior $2 / 3$ of the tongue:
- At $4^{\text {th }}$ week 3 endodermal swellings appears from $1^{\text {st }}$ pharyngeal arches
- 1- A median swelling called the tuberculum impar
- 2- Two lateral lingual swellings proliferate and grow medially.
- In the midline, they fused together at median sulcus and completely covered tuberculum impar forming mucous membrane of anterior $2 / 3$ of the tongue.
- So the anterior $2 / 3$ is supplied by Lingual nerve from posterior division of mandibular nerve.
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- Development of the mucous membrane of the posterior $1 / 3$ of the tongue:
- It develops as a large midline mass (copula of His) derived from endoderm of 2nd, $3^{\text {rd }}$ and 4th pharyngeal arches.
- The part of the $3^{\text {rd }}$ arch proliferates and migrates forward overlying the $2^{\text {nd }}$ arch and forms posterior $1 / 3$ of the tongue that fused with anterior $2 / 3$ by sulcus terminalis. - So the posterior $1 / 3$ is supplied by the glossopharyngeal nerve.
- Development of the mucous membrane of the Root of the tongue;
- From the part of the copula derived from the $4^{\text {th }}$ pharyngeal arch
- So it is supplied by the vagus nerve (internal laryngeal nerve).


## - Development of the muscles of the tongue:

- The muscles of the tongue are derived from the occipital myotomes except palatoglossus muscle that develop from the mesoderm in situ.
- So the muscles are supplied by the hypoglossal nerve except palatoglossal muscle supplied by pharyngeal nerve plexus


## Separation of the tongue

- At first the tongue is adherent to the floor of the mouth, then a horse-shoe (C-shaped) groove called alveololingual groove separates the anterior $2 / 3$ of tongue from the floor of the mouth except in the midline where the tongue is connected to the floor by the frenulum of the tongue.

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** Congenital anomalies of the tongue

- A glossa: failure of development of the tongue,
- Macroglossia: large sized tongue which protrudes from the mouth (as in mongolism and cretinism).
- Microglossia: small sized tongue
- Bifid tongue: due to failure of fusion of the 2 lingual swellings.
- Tongue-tie (Ankyloglossia): the tongue is adherent to the floor of the mouth.
- Short frenulum: due to incomplete separation of tongue (common).
- Long frenulum: due to excess separation of the tongue. It causes the tongue to fall back and close the pharynx and larynx (suffocation).


 from the ventral wall of the duodenal loop by the 4th week of development
- The diverticulum grows ventrally and cranially into ventral mesentery.
- The diverticulum divides into 2 parts:
- Cranial part called pars hepatica
- Caudal part called pars cystica.



## II- Development of the gall bladder (Pars cystica)

a- Distal part is dilated and forms gall bladder.
b- Proximal part remains narrow and forms cystic duct.

## III- Development of the common bile duct:

- The proximal part of the hepatic diverticulum forms the common bile duct.
** At first, the common bile duct opens in the ventral wall of the duodenum.
- After rotation of the duodenal loop $90^{\circ}$ (clockwise) and unequal growth of its walls,
a) The opening shifts to the dorsomedial wall of the $2^{\text {nd }}$ part of the duodenum.
b) The common bile duct passes behind the first part of the duodenum


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** Development of ligaments of liver:

- Development of the liver between the layers of the ventral mesentery divides it into:
a- Ventral part connects liver to anterior abdominal wall (falciform ligament).
b- Dorsal part connects liver to stomach (lesser omentum)
c- Cranial part forming triangular and coronary ligaments
orst Lesser omentum:
- Hepatogastric ligament

** Congenital Anomalies of the liver and biliary system:
I. Agenesis or hypo-genesis of the liver: due to failure of formation of the hepatic diverticulum or due to incomplete development of the hepatic bud.
II. Abnormal number of the liver lobes: due to abnormal division of the pars hepatica.
III. Agenesis of the gall bladder: failure of development of the cystic bud
IV. Double gall bladder: abnormal division of the cystic bud into 2 parts.
V. Mobile gall bladder: the gall bladder is completely separated from the liver and completely covered with peritoneum.
VI. Atresia (narrowing) of the biliary ducts: due to failure of their canalization. It is associated later with congenital jaundice


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** Congenital abnormalities of the esophagus

- Esophageal atresia (obstruction): associated with polyhydramnios due to failure of swallowing of the amniotic fluid
- Esophageal stenosis (narrow): due to a- Posterior displacement tracheoesophageal septum.
b- Mechanical factors push posterior wall of the tube forward.


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- Tracheoesophageal fistula:
- An abnormal opening between esophagus and trachea caused by failure of complete closure of the tracheoesophageal septum.

1. Proximal part of esophagus ends as a blind sac and distal part continues with the trachea.

2- Proximal part of esophagus continues with trachea and distal part ends as blind sac.


Most
3- Proximal and distal parts of continue with trachea by single tube .

4- Proximal and distal parts of esophagus continue with trachea separately by double tubes.



[^0]:    https://www.youtube.com/@ProfDrYoussefHusseinAnatomy/playlists

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