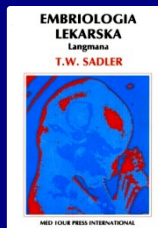
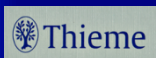




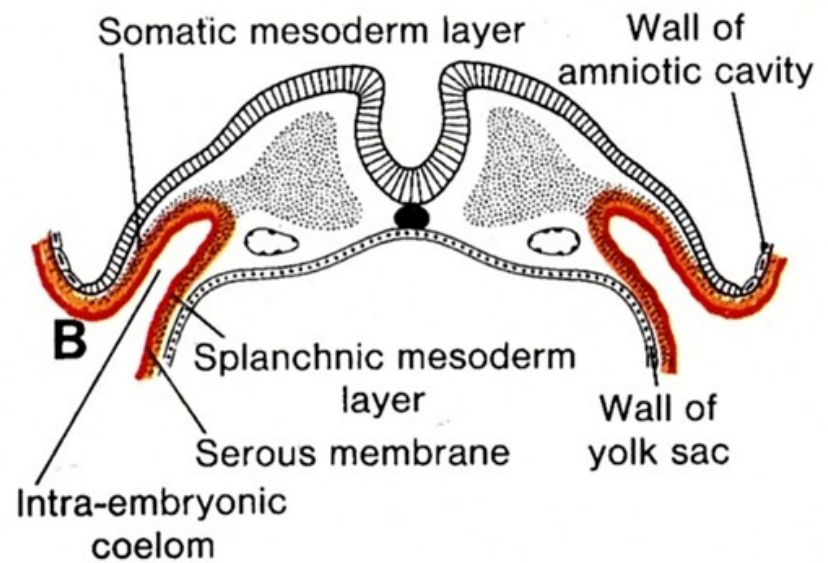
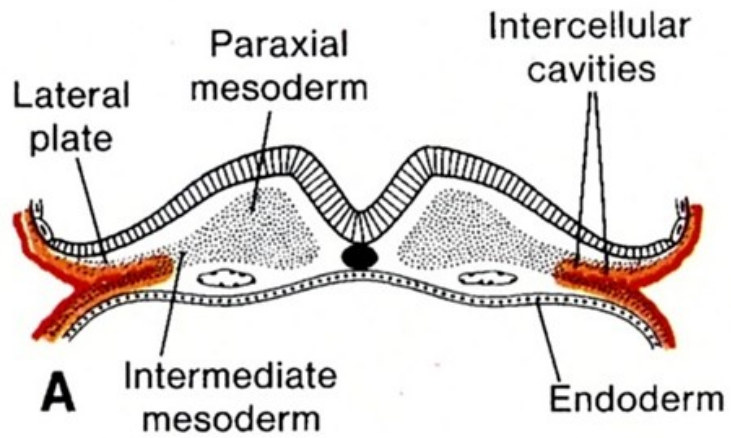
Rozwój jam ciała i układu oddechowego



Przy końcu 3. tygodnia leżąca po obu stronach linii pośrodkowej mezoderma wewnątrzrodkowa różnicuje się na część przyosiową, pośrodkową i boczną.

Z chwilą pojawienia się w mezodermie bocznej rozstępów komórkowych – mezoderma rozdziela się na ścienną i trzewną.

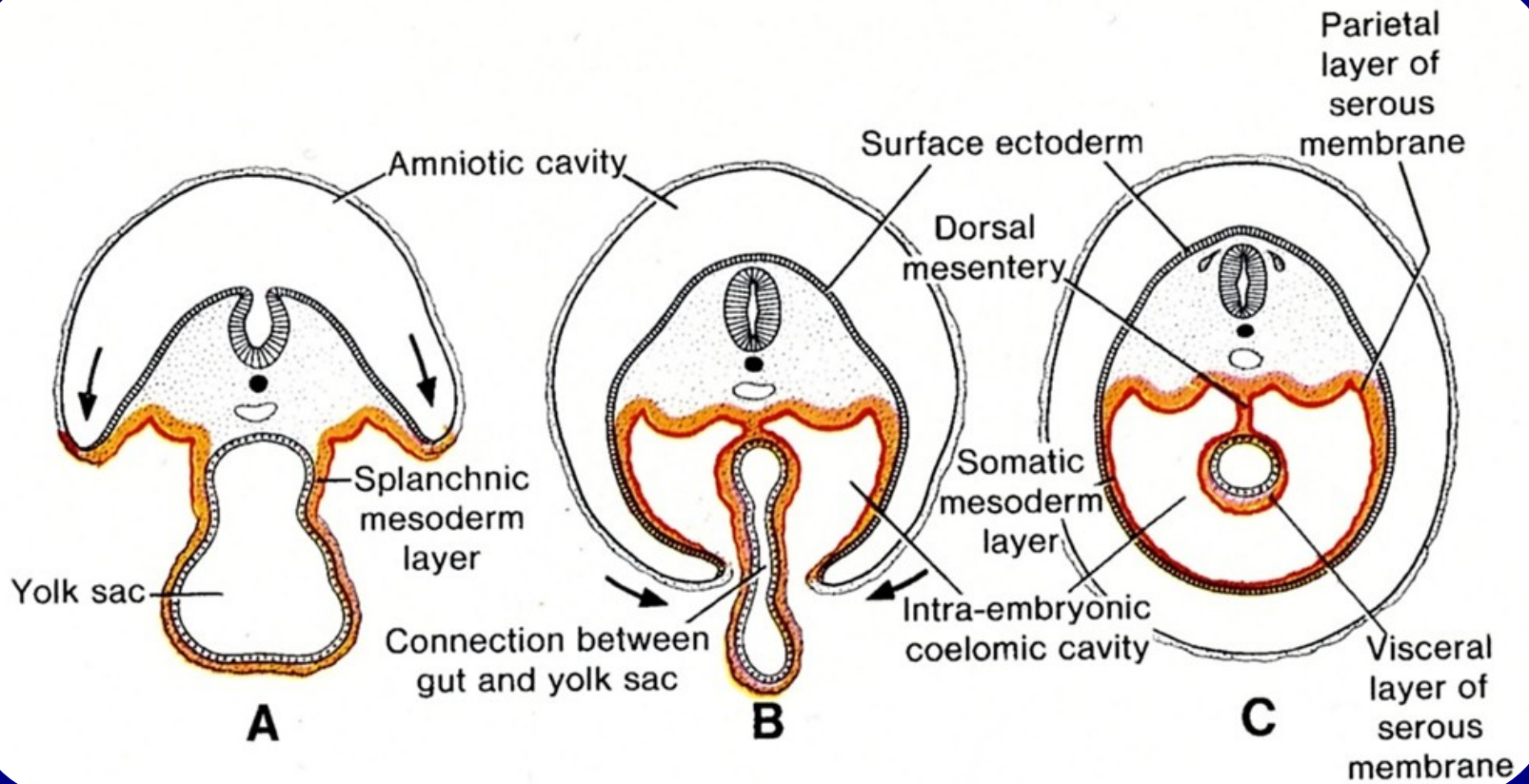
Trzewna przechodzi w mezoderme ściany pęcherzyka żółtkowego.



Przestrzeń między mezoderma trzewną i ścienną
tworzy wewnątrzrodzkową jamę ciała.

Początkowo lewa i prawa strona wewnątrzzarodkowej jamy ciała łączą się na znacznej przestrzeni z zewnątrzzarodkową jamą ciała.

W momencie pofałdowania zarodka w kierunku głowowo-ogonowym oraz poprzecznym, połączenie to zanika.



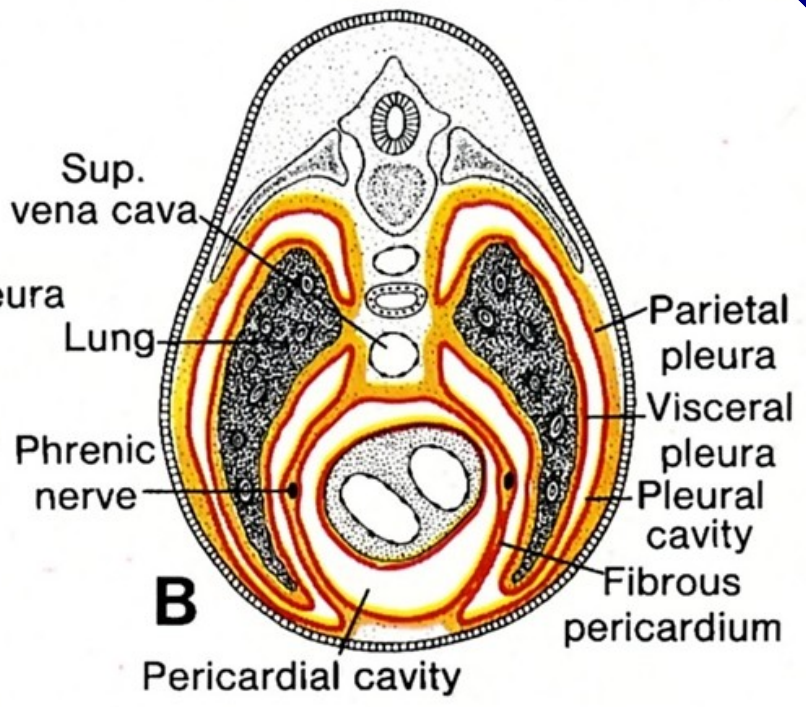
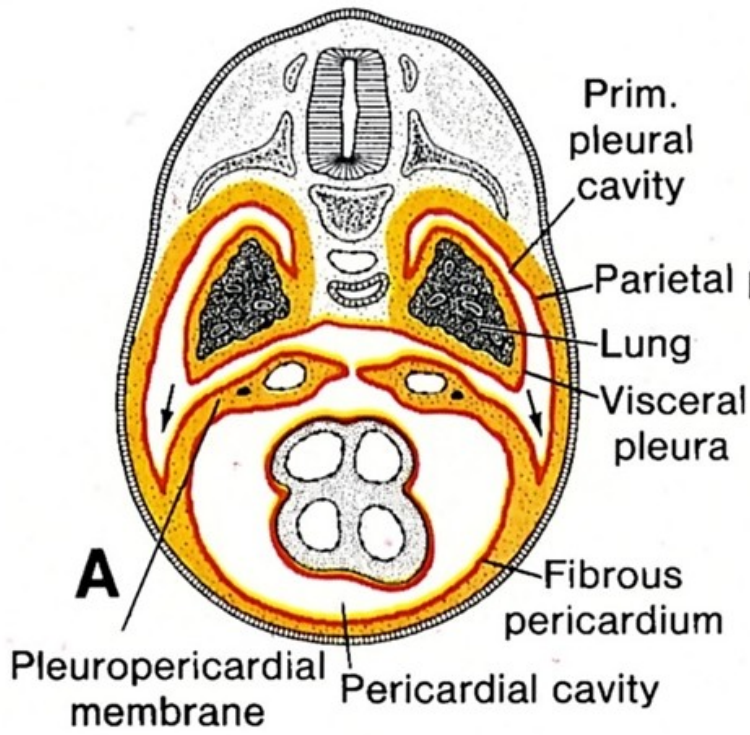
Wyścielające wewnątrzrodzkową jamę ciała komórki przekształcają się w mezotelium (nabłonek surowiczy) i tworzą blaszkę ścienną błon surowiczych, wyścielającą zewnętrzną warstwę jam: otrzewnej, opłucnej i osierdza. Mezoderma trzewna tworzy warstwę wewnętrzną trzewną błon surowiczych.

Przepona dzieli jamę ciała na jamę klatki piersiowej i jamę otrzewnej.

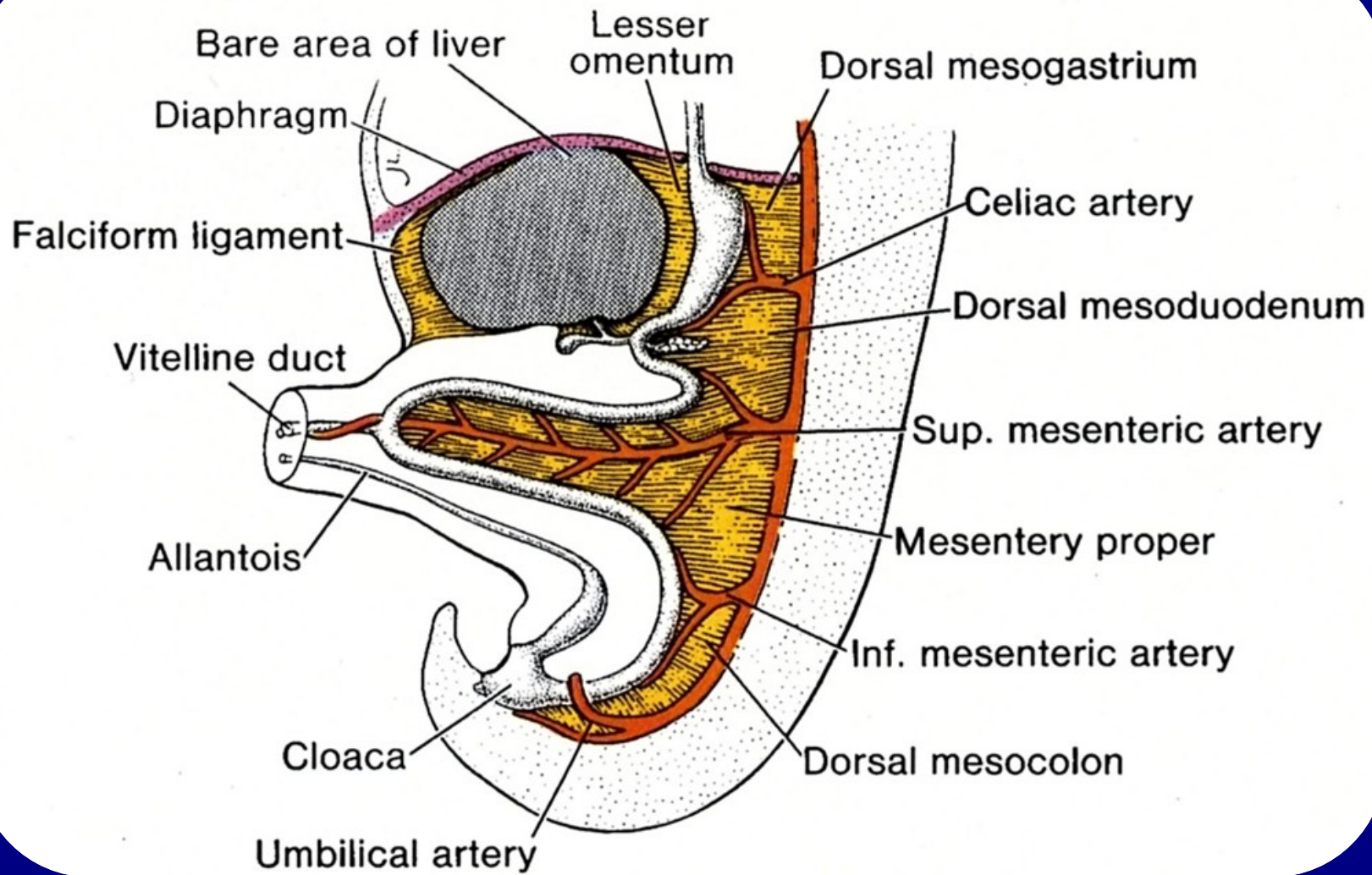
Powstaje z czterech elementów:

- przegrody poprzecznej
- błon otrzewnowo-opłucnowych
- kręzki grzbietowej przęłyku
- komórek mięśniowych ze ściany ciała

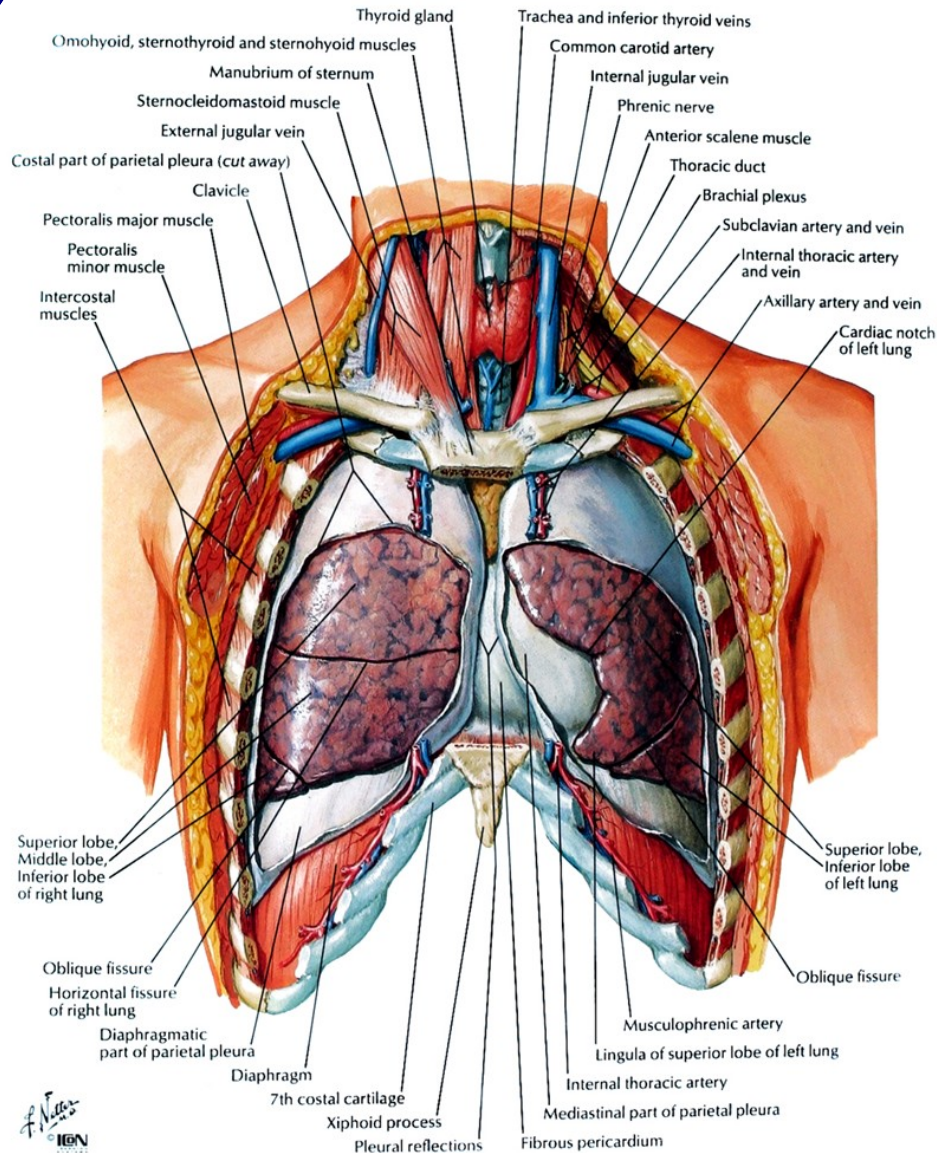
Jama klatki piersiowej zostaje podzielona przez błony opłucnowo-osierdziowe na jamę osierdza i dwie jamy opłucnej.

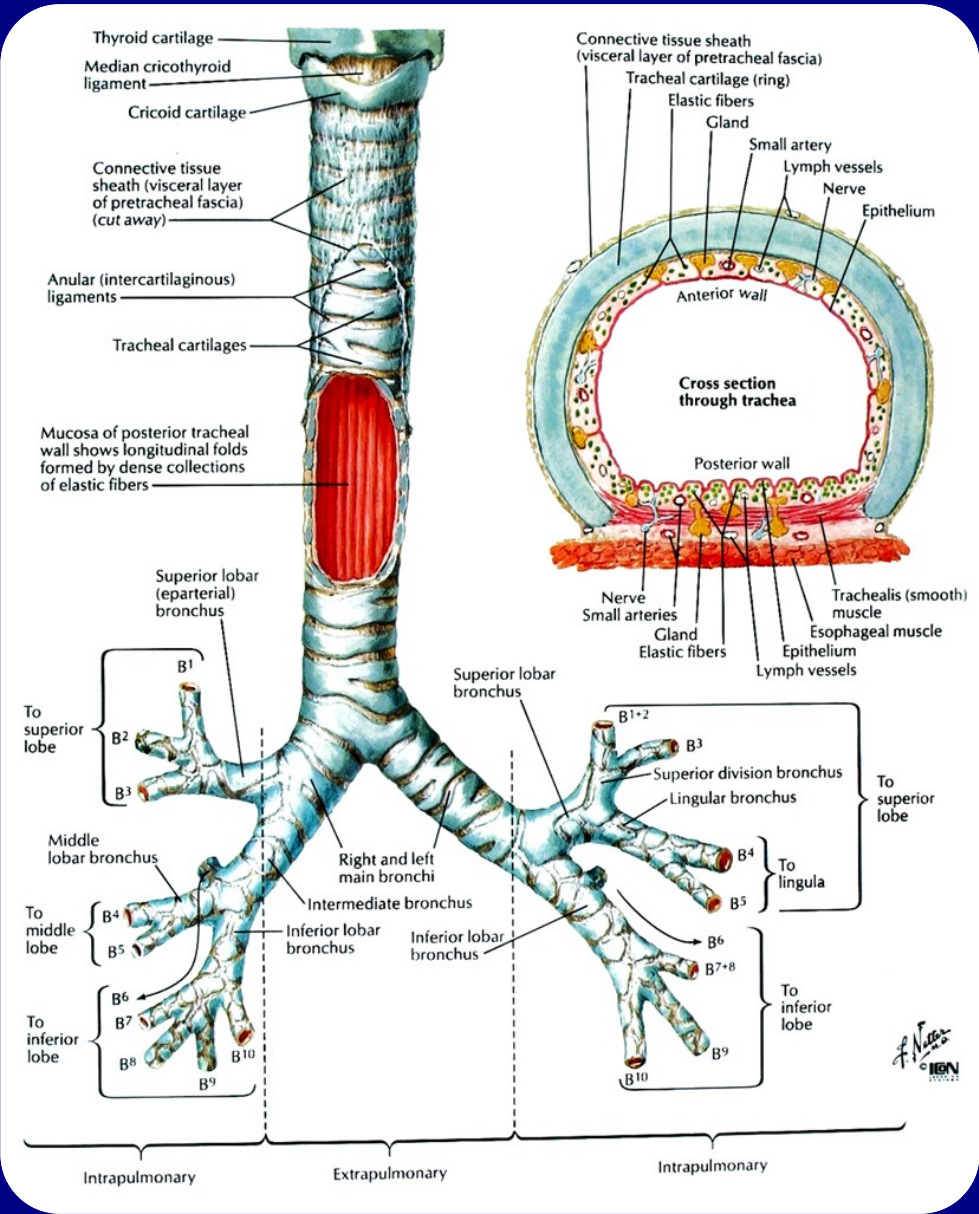


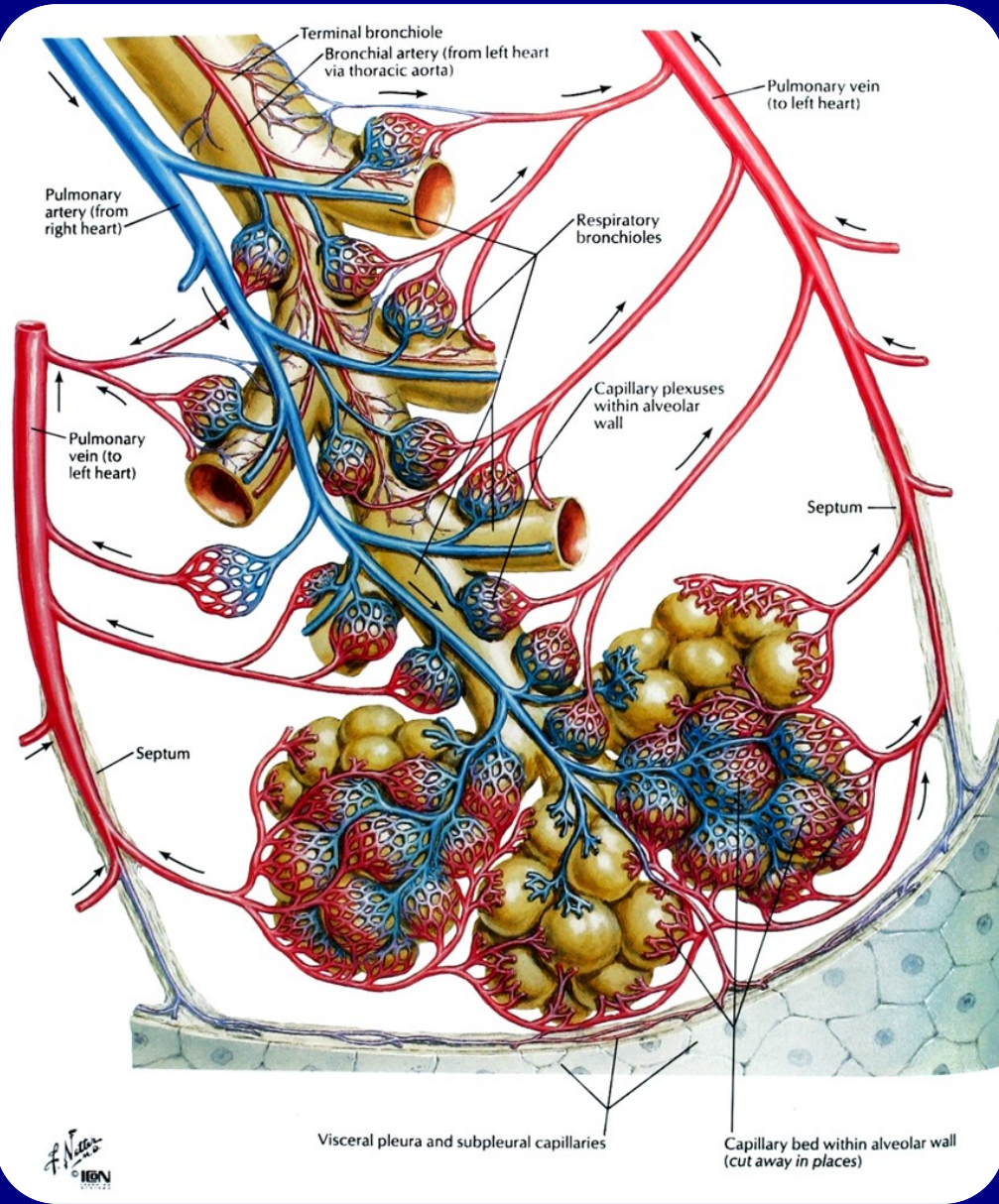
Wraz z fałdowaniem poprzecznym tarczy zarodkowej obie jej części boczne zbliżają się do siebie i łączą w linii pośrodkowej. Prąjelito zostaje otoczone przez mezoderme trzewną.

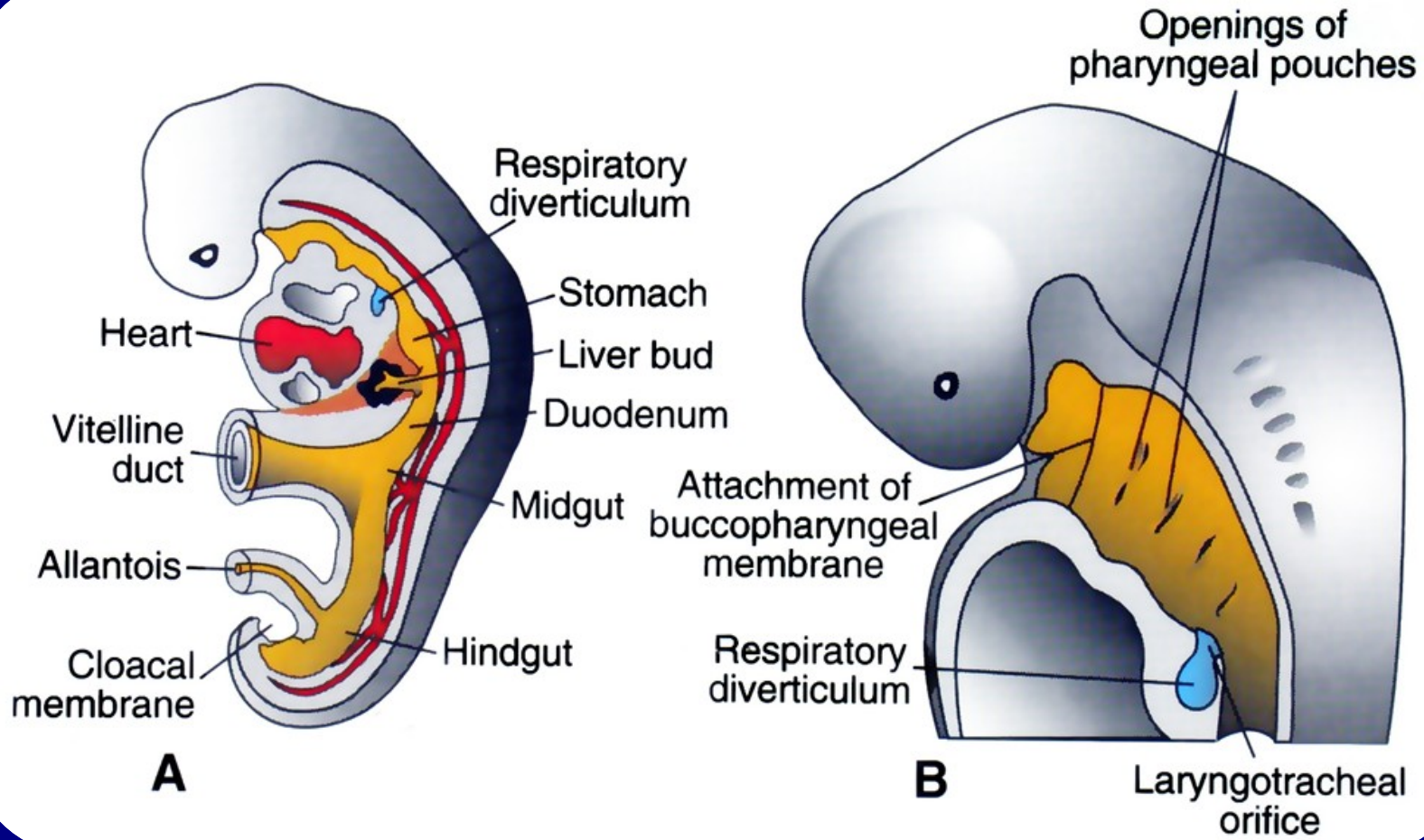


Rozwój układu oddechowego

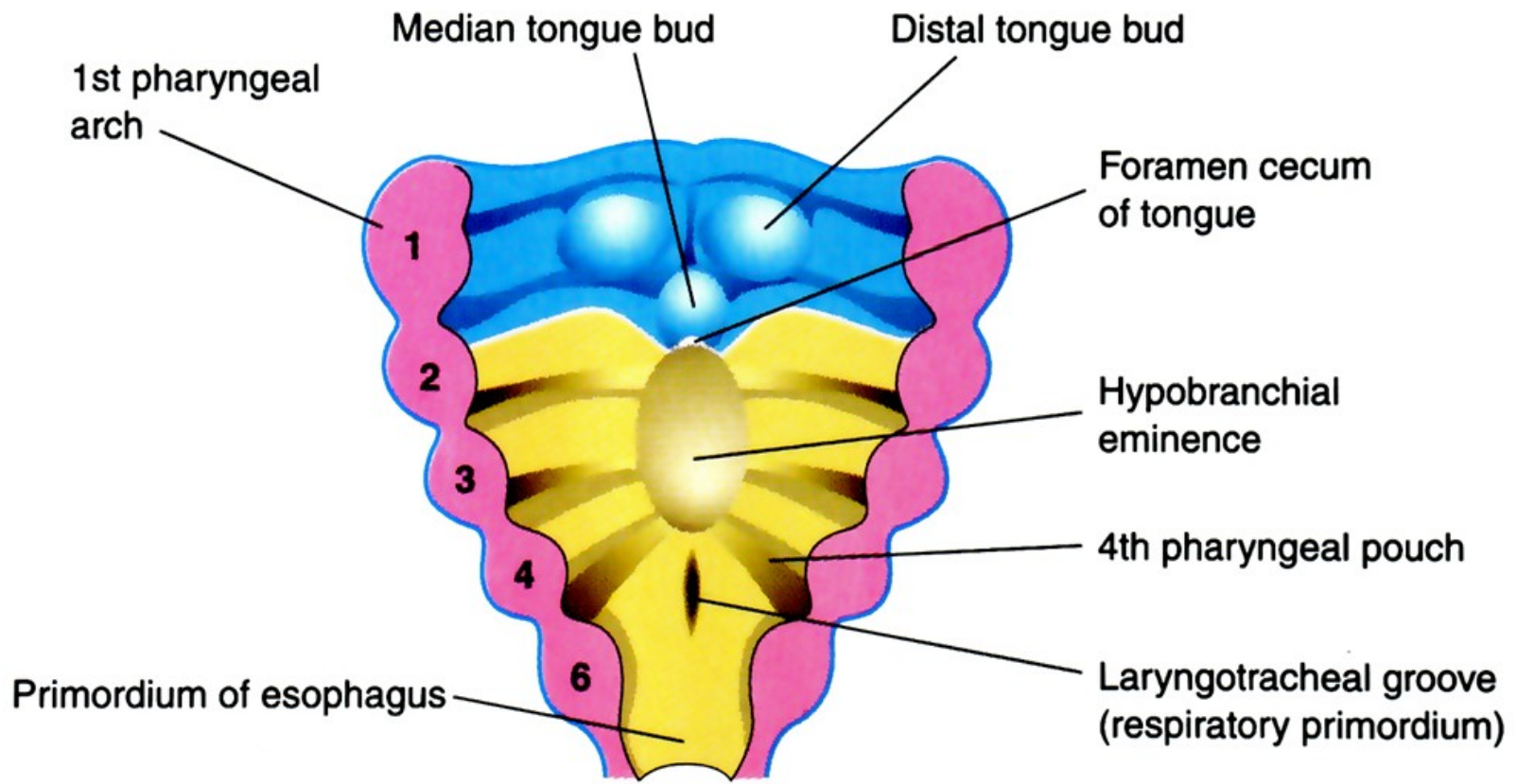






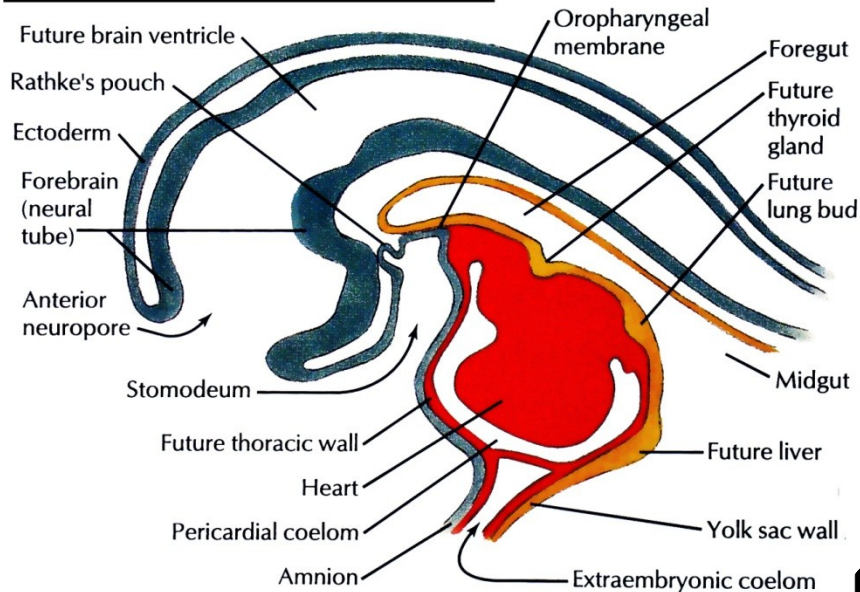


A – Embryo of approximately 25 days gestation showing the relation of the respiratory diverticulum to the heart, stomach, and liver. B – Sagittal section through the cephalic end of 5-week embryo showing the openings of the pharyngeal pouches and the laryngotracheal orifice.

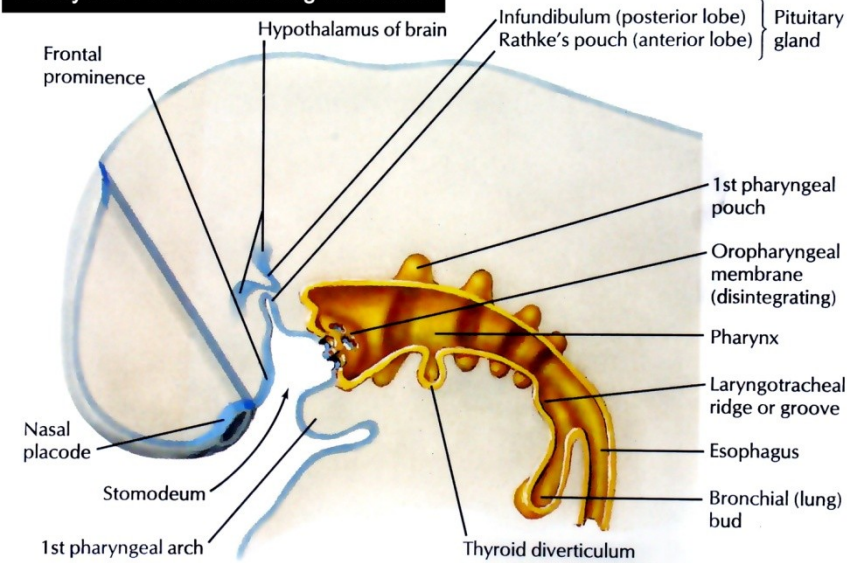


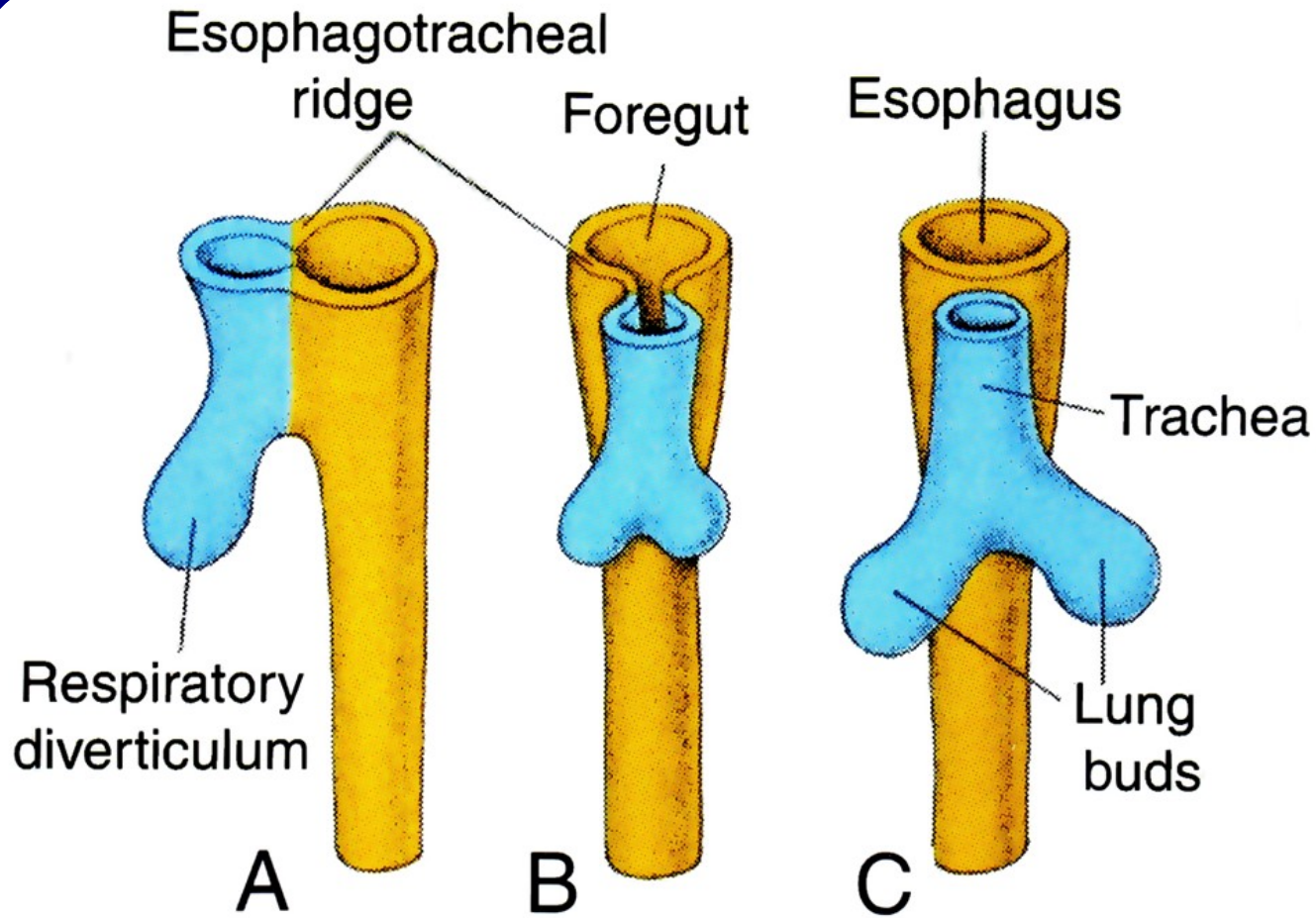
Horizontal section of the embryo, illustrating the floor of the primordial pharynx and the location of the laryngotracheal groove.

Embryo at 3 to 4 weeks. Sagittal section



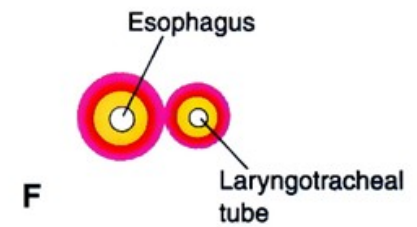
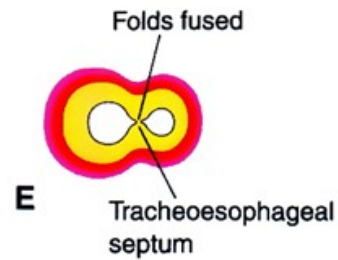
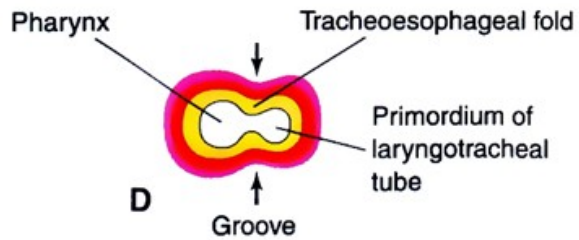
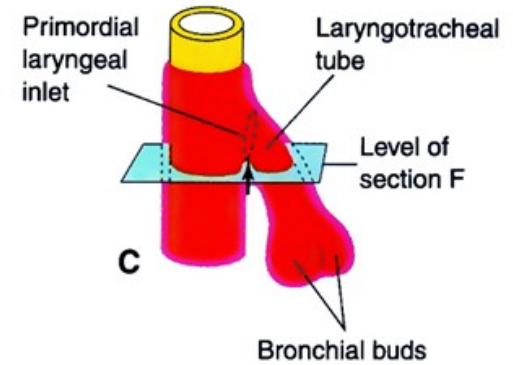
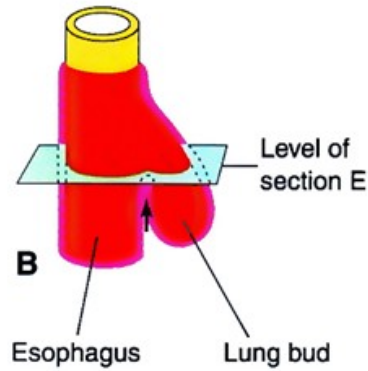
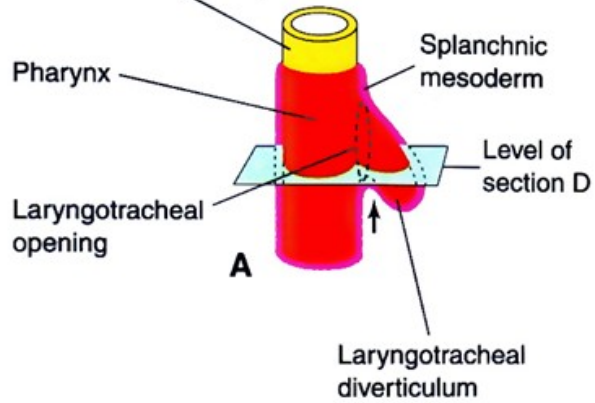
Embryo at 4 to 5 weeks. Sagittal section

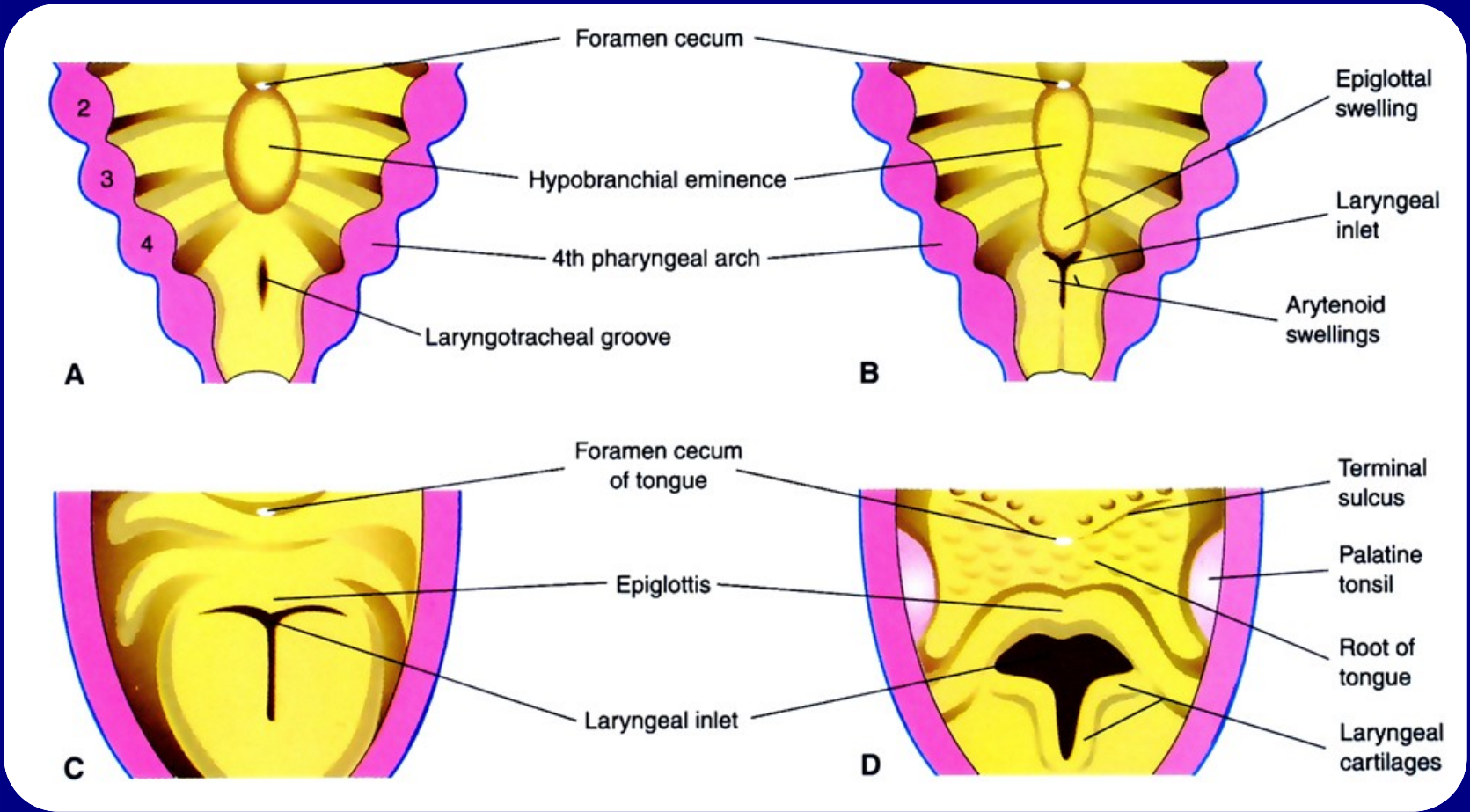




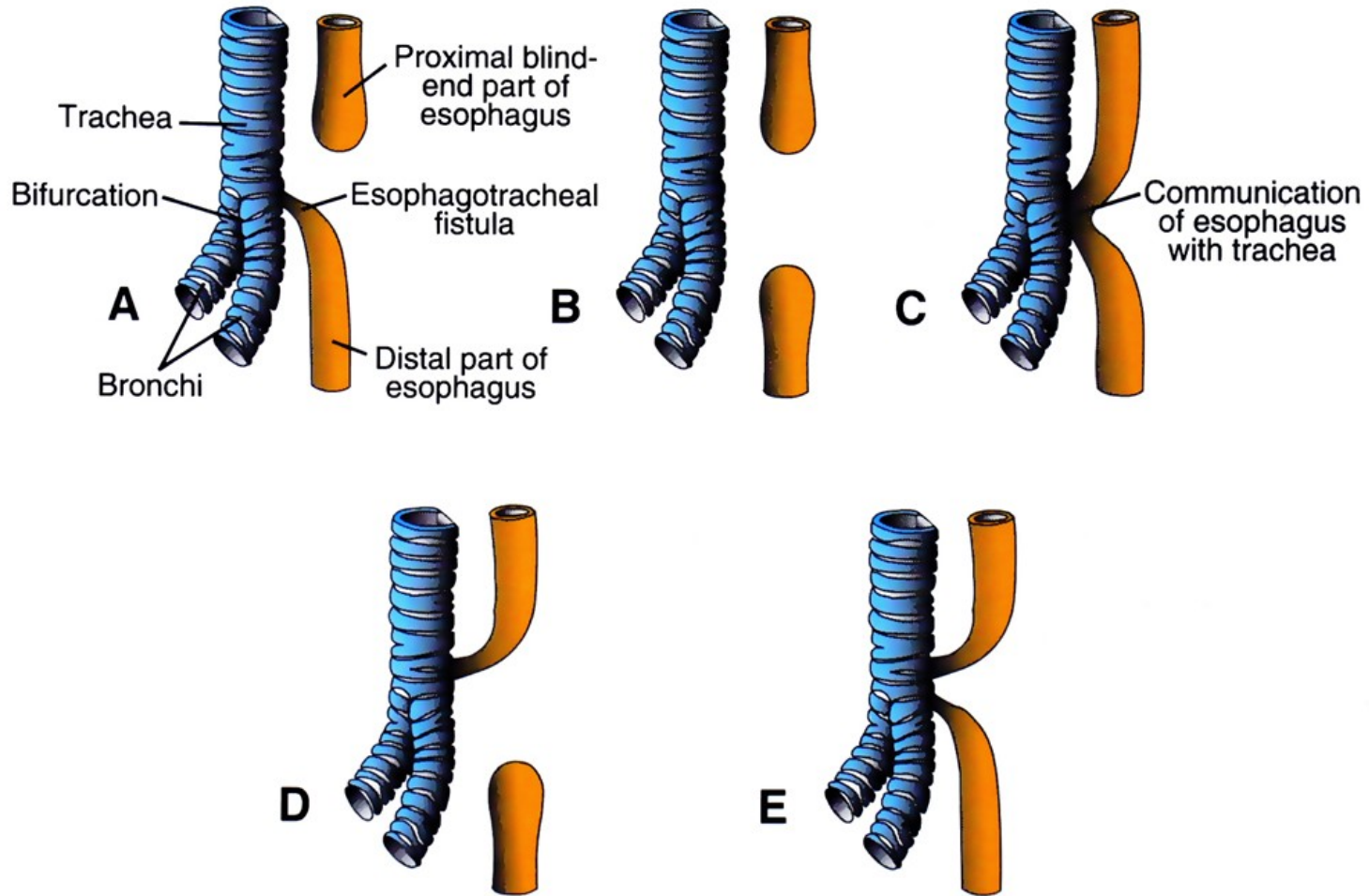
A, B, and C. Successive stages in development of the respiratory diverticulum showing the esophagotracheal ridges and formation of the septum, splitting the foregut into esophagus and trachea with lung buds.

Endoderm
(mesoderm removed)

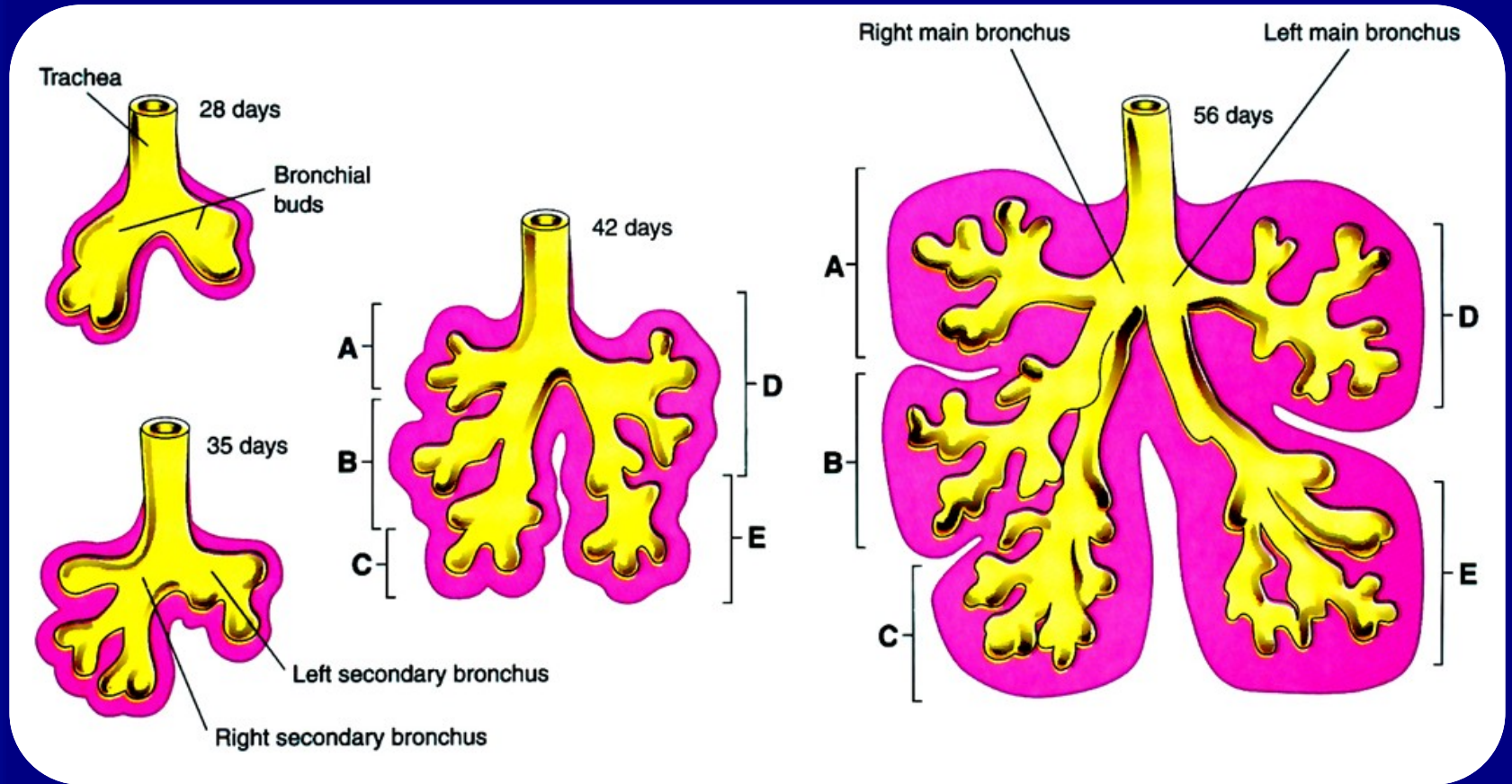




A – 4 weeks, B – 5 weeks, C – 6 weeks, D – 10 weeks



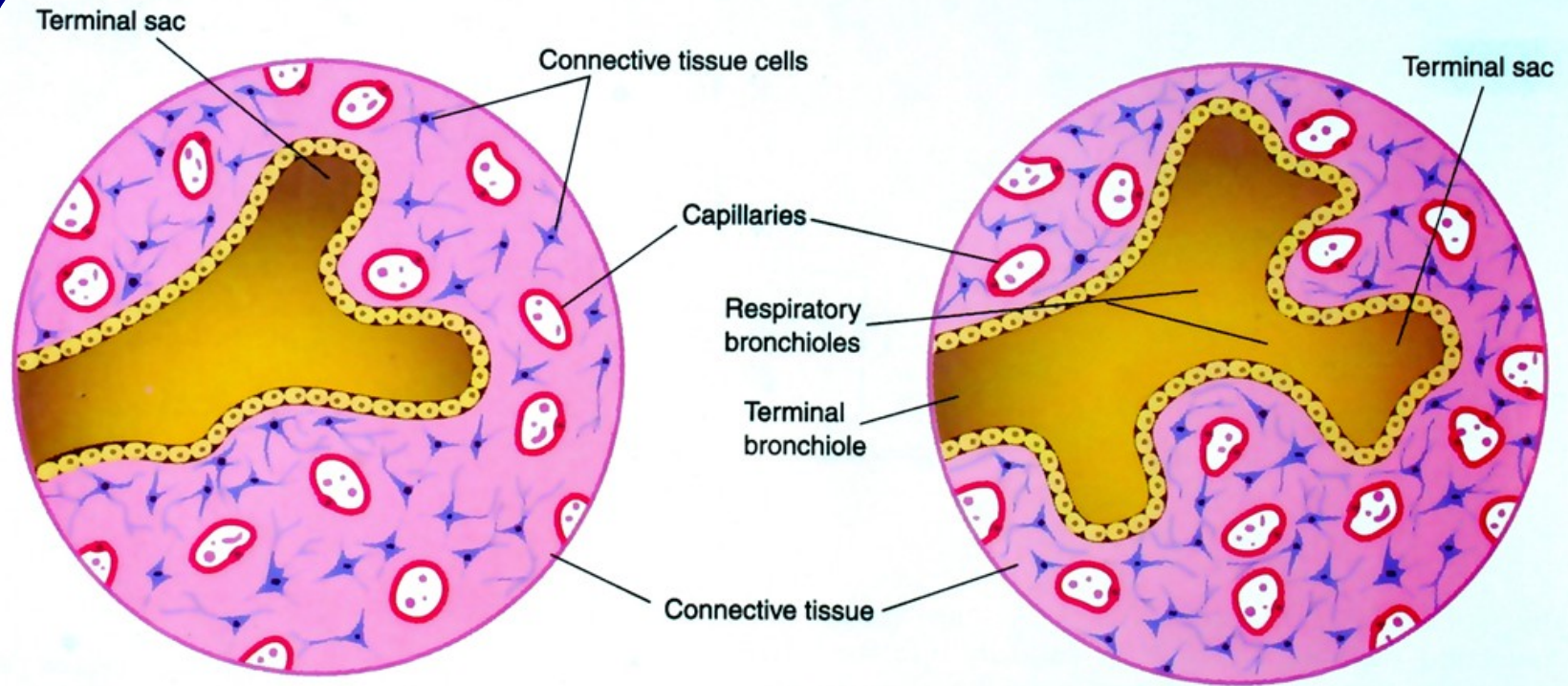
A – The most frequent abnormality (90% cases) occurs with the upper esophagus ending in a blind pouch and the lower segment forming a fistula with the trachea. B – Isolated esophageal atresia (4% of cases). C – H-type tracheoesophageal fistula (4% of cases). D and E – Other variations (each 1% of cases)



A – Right upper (superior) lobe, B – Right middle lobe, C – Right lower (inferior) lobe, D – Left upper (superior) lobe, E – Left lower (inferior) lobe.

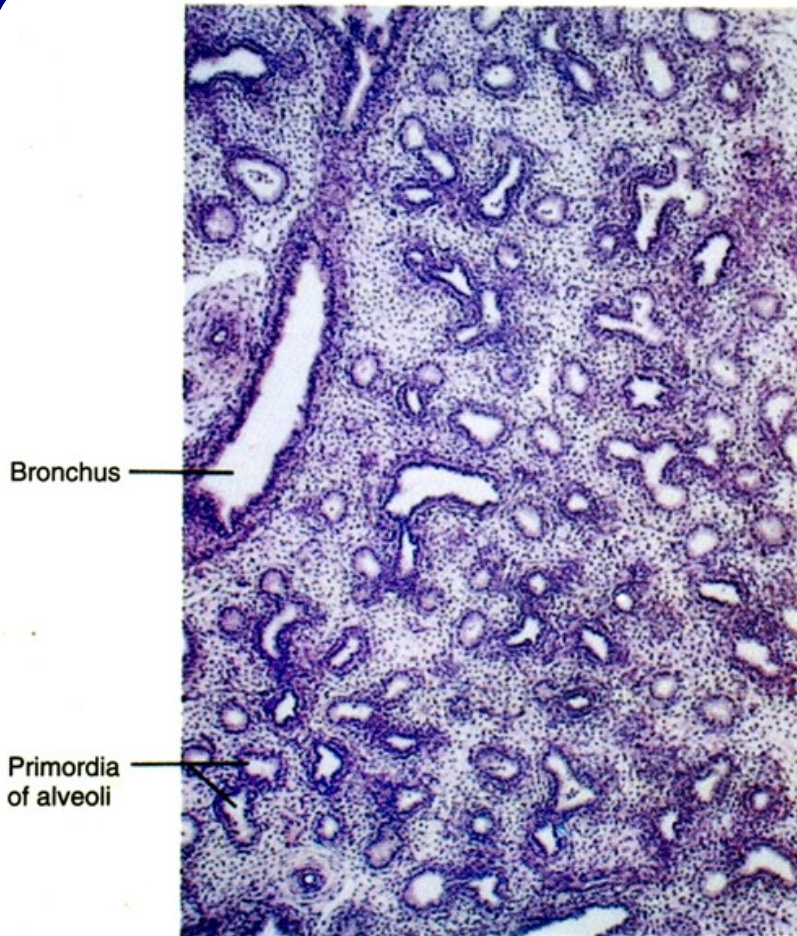
Maturation of the Lungs

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.



A Pseudoglandular period (1-17 weeks)

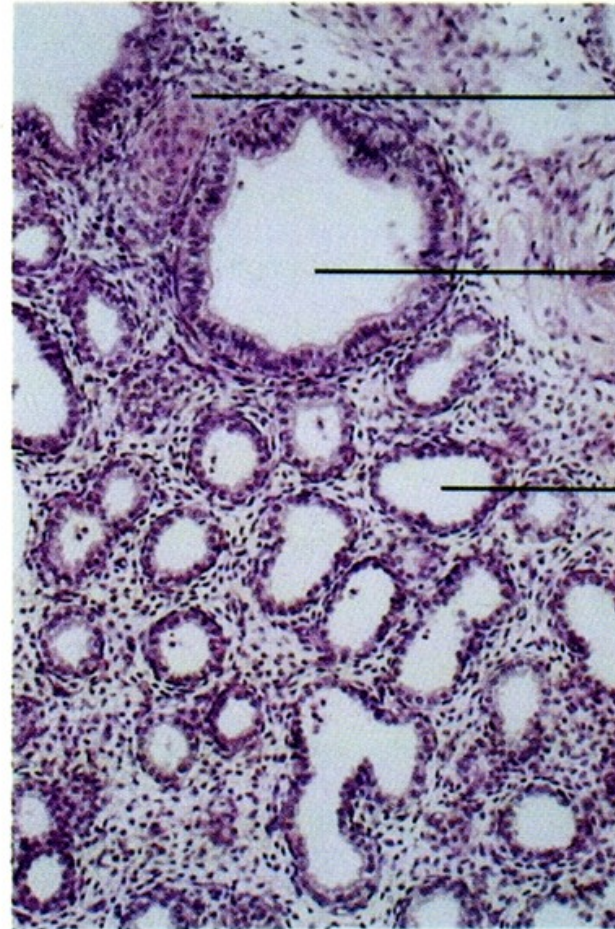
B Canalicular period (16-25 weeks)



Bronchus

Primordia
of alveoli

A

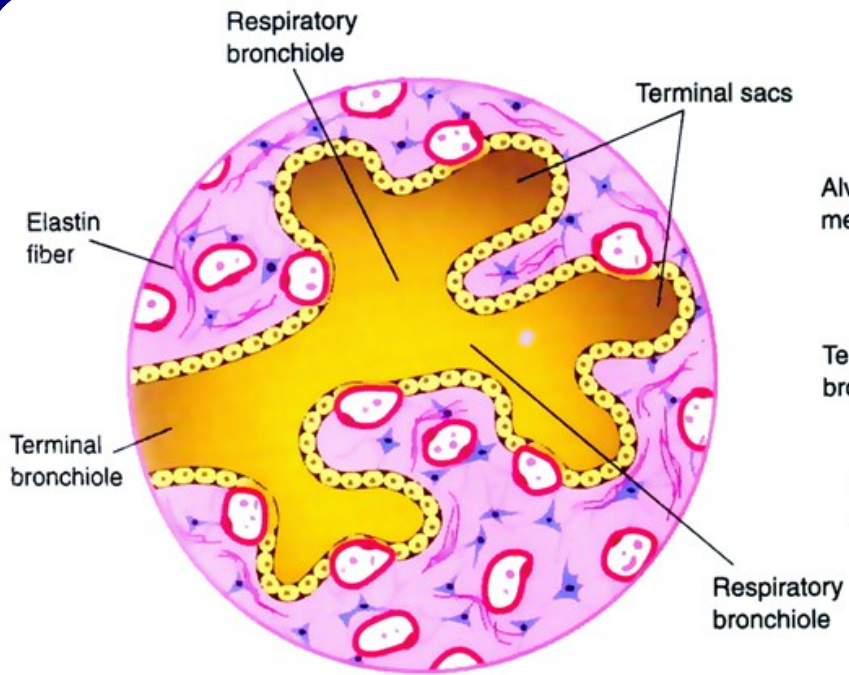


Cartilage
primordium

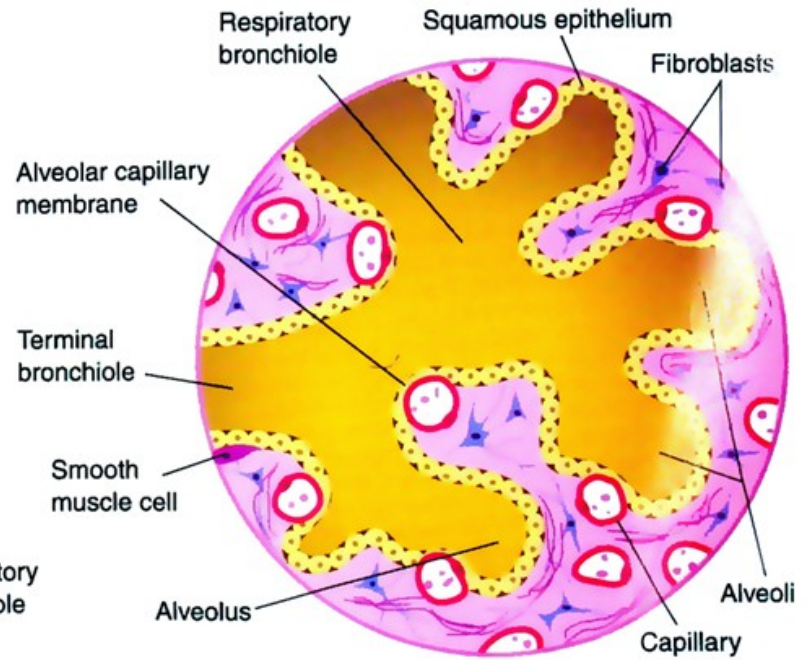
Bronchus

Respiratory
bronchiole

B



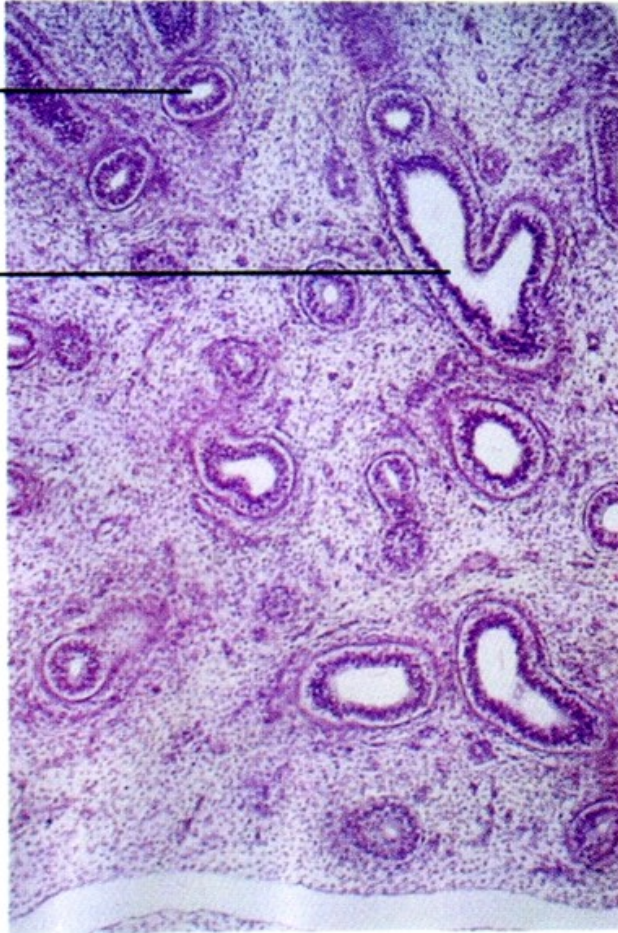
C Terminal sac period (24 weeks–birth)



D Alveolar period (birth–8 years)

Bud for
bronchiole

Stem
bronchus



C

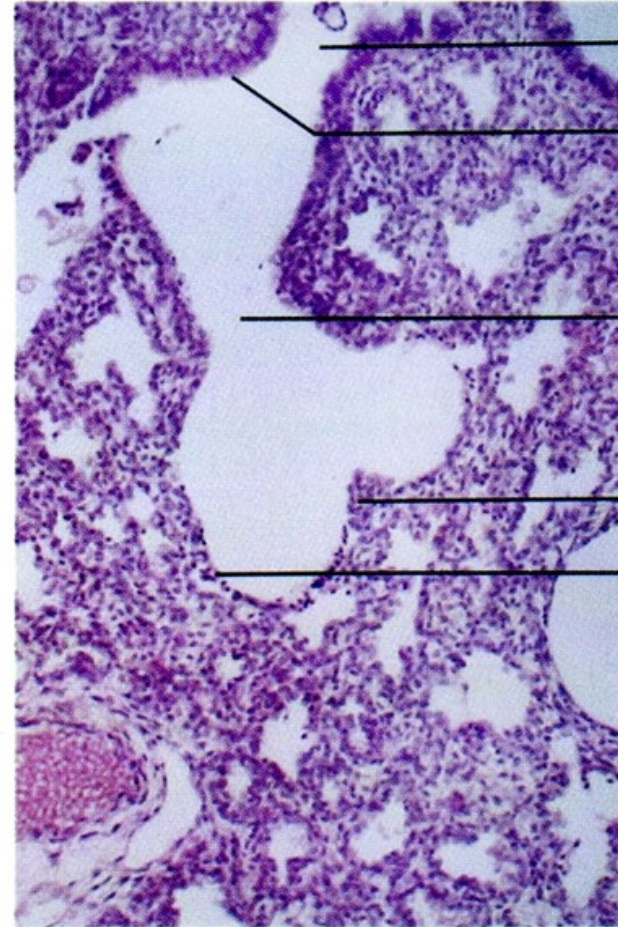
Respiratory
bronchiole

Cuboidal
epithelium

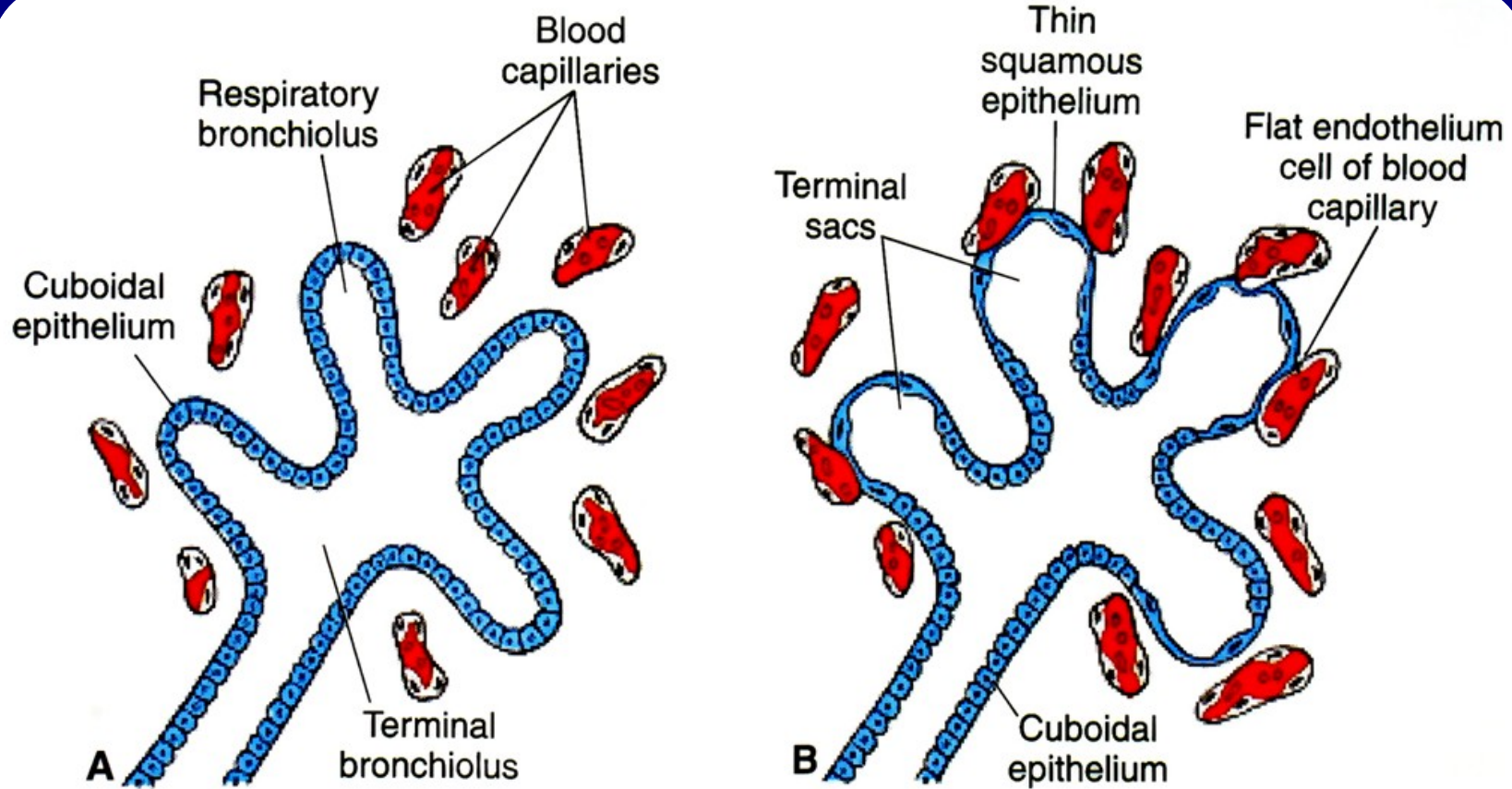
Alveolar
duct

Capillary

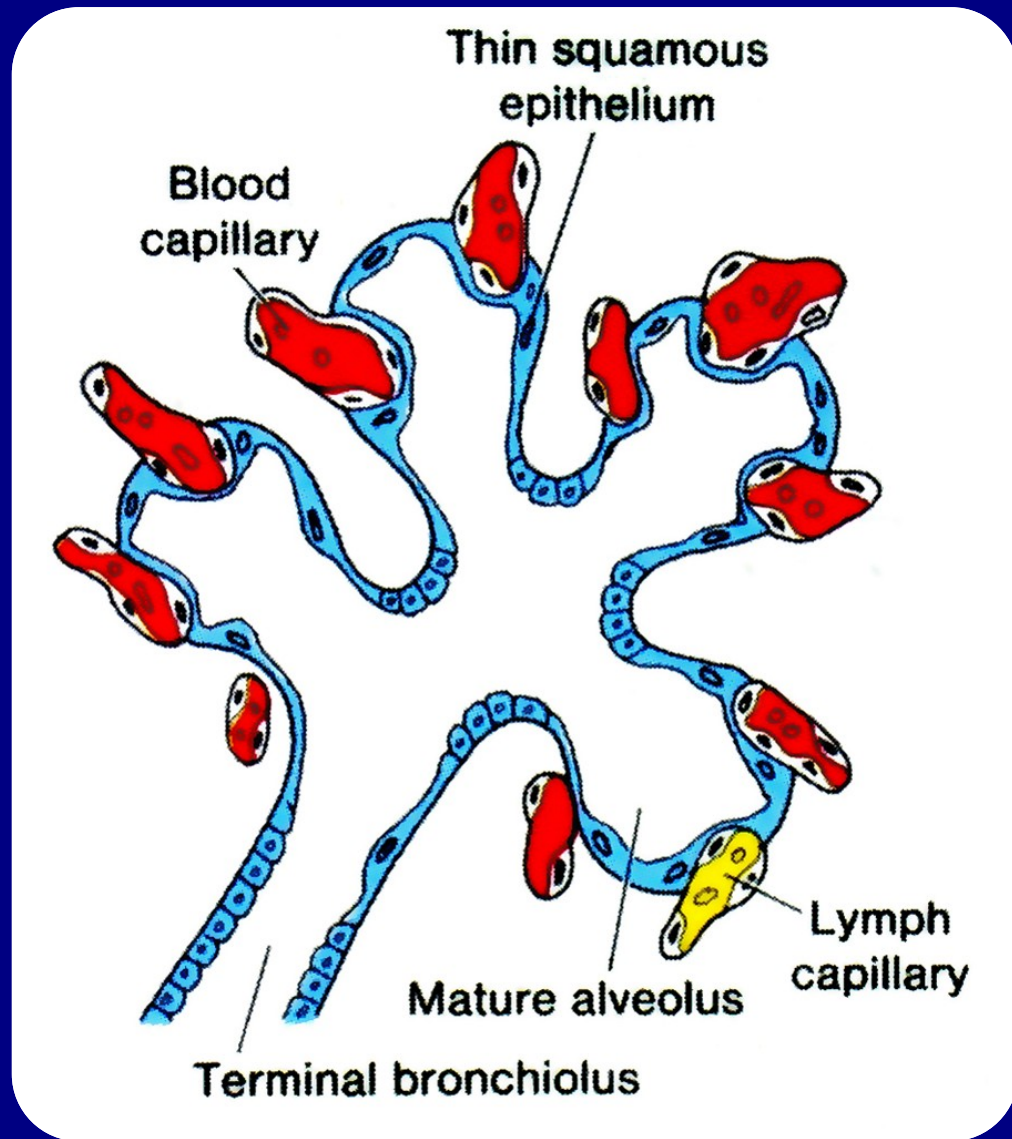
Flattened
epithelium
of alveolus



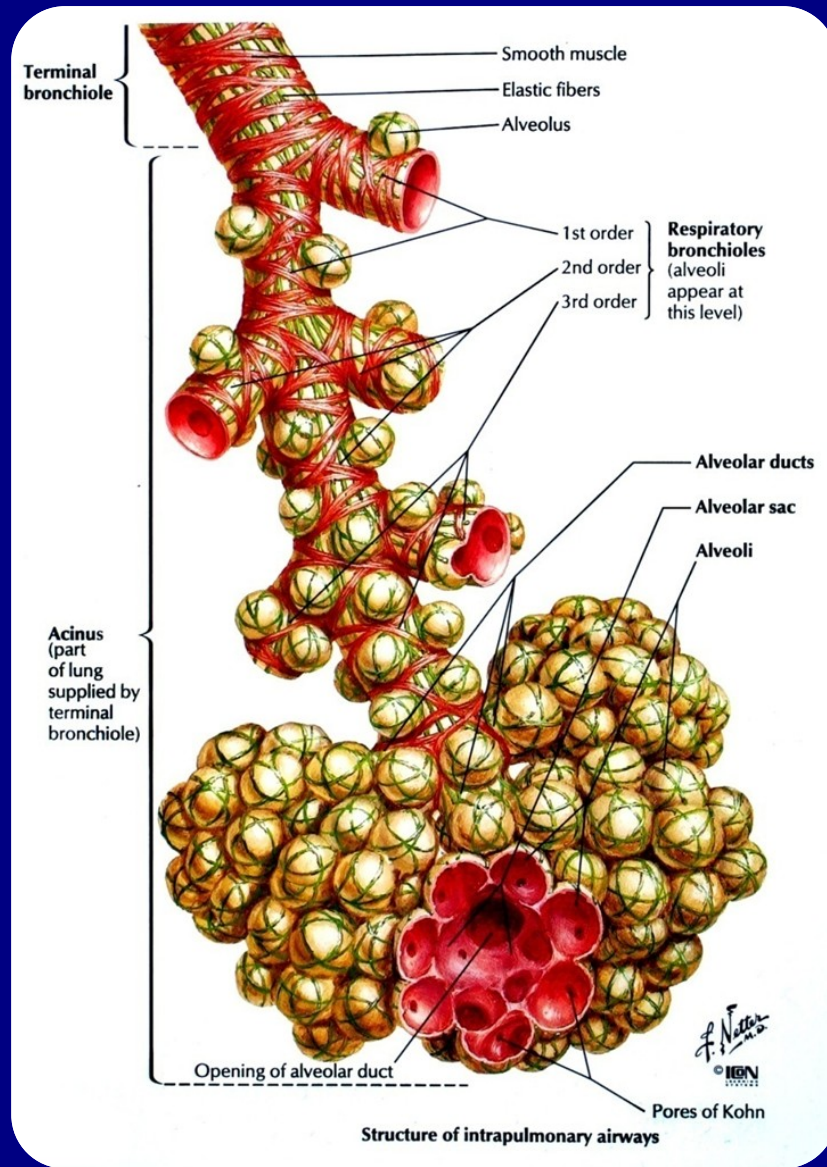
D



A – The canalicular period, B – The terminal sac period.



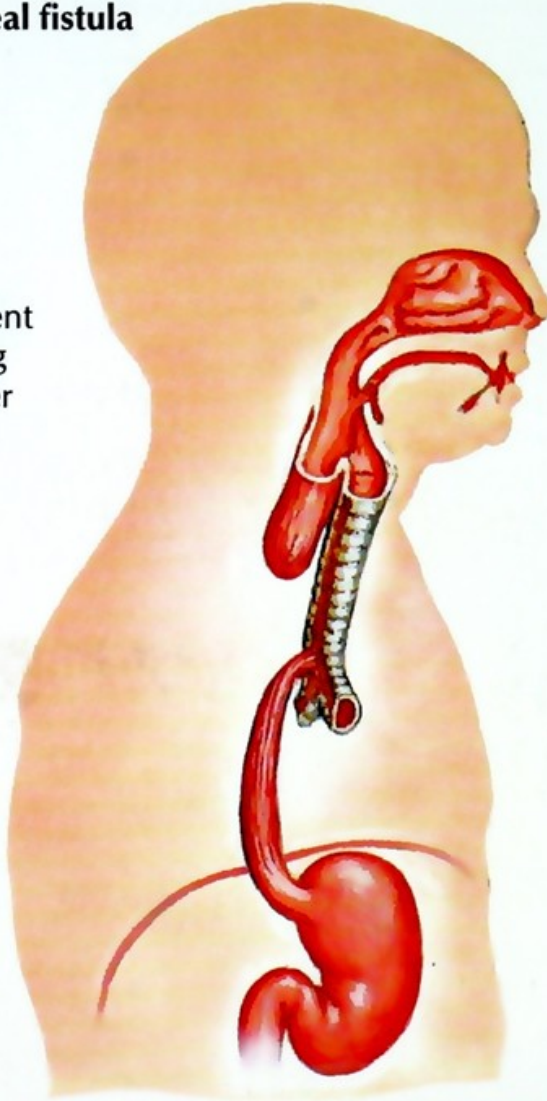
Lung tissue in a newborn



Development of bronchioles and alveoli

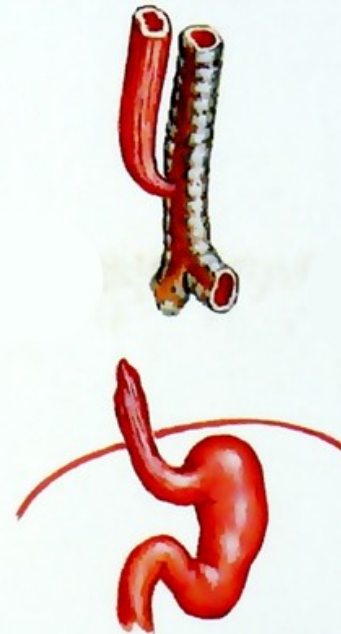
A. Tracheoesophageal fistula

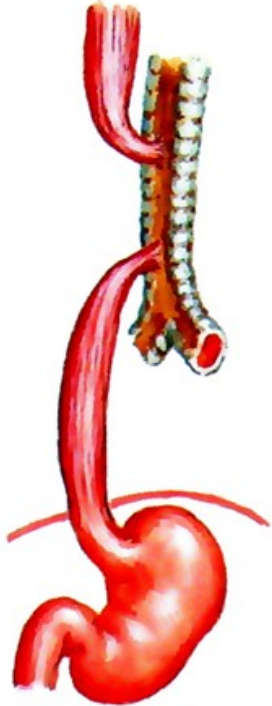
Most common form (90% to 95%) of tracheoesophageal fistula. Upper segment of esophagus ending in blind pouch; lower segment originating from trachea just above bifurcation. The two segments may be connected by a solid cord



B. Variations of tracheoesophageal fistula and rare anomalies of trachea

Upper segment of esophagus ending in trachea; lower segment of variable length

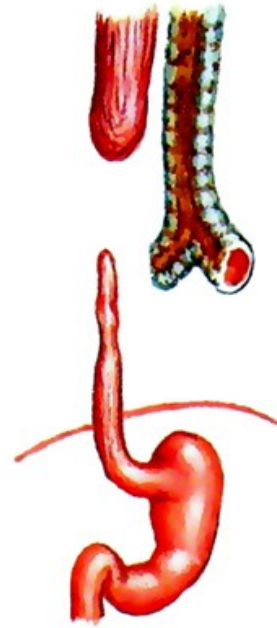




C. Double fistula



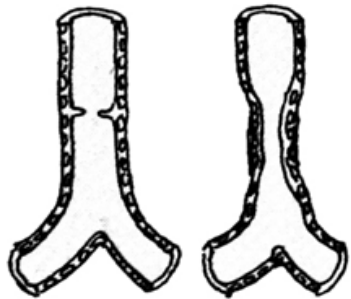
D. Fistula without esophageal atresia



E. Esophageal atresia without fistula

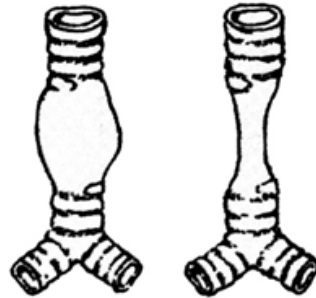


F. Aplasia of trachea (lethal)



Web Hourglass

G. Stricture of trachea

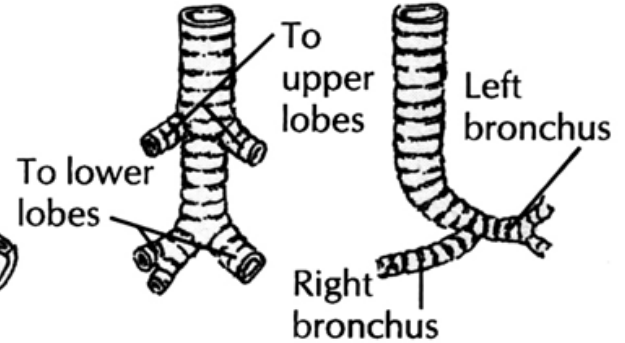


Inspiration Expiration

H. Absence of cartilage

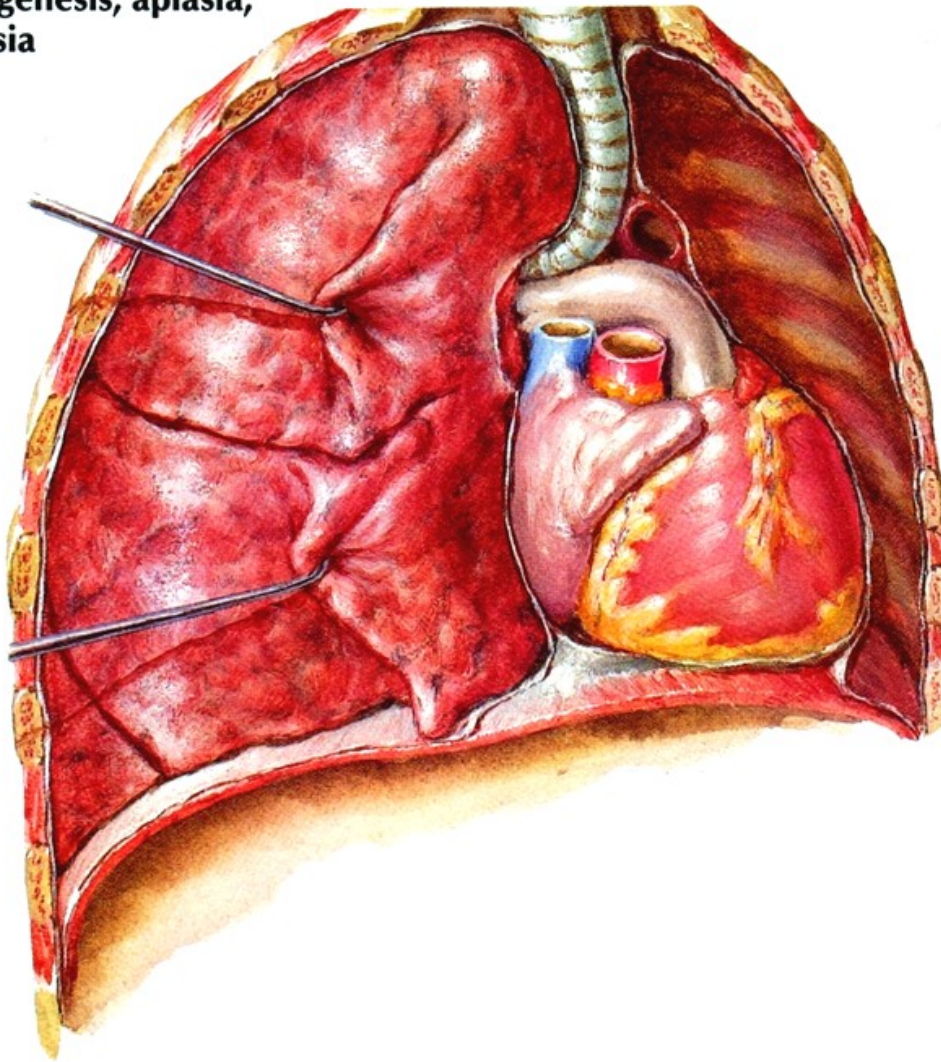


I. Deformity of cartilage



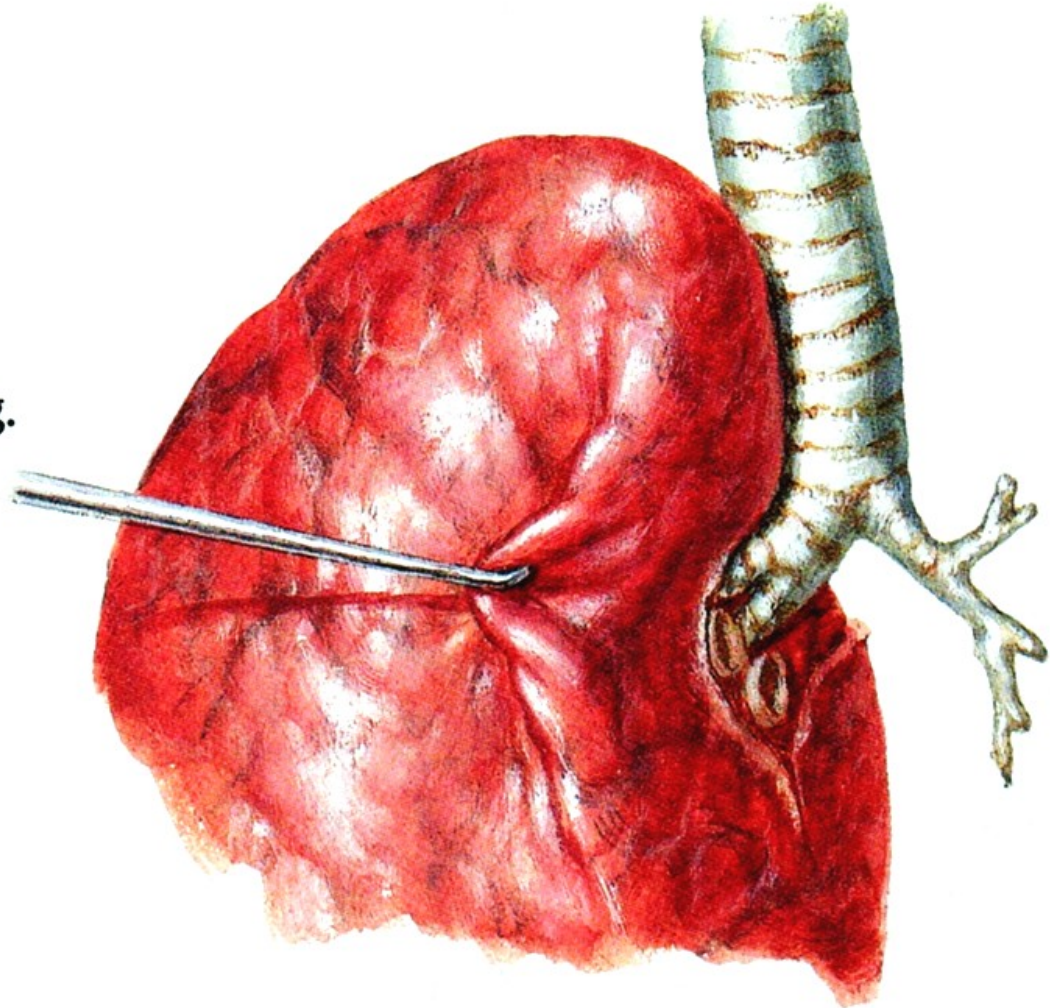
J. Abnormalities of bifurcation

**Pulmonary agenesis, aplasia,
and hypoplasia**

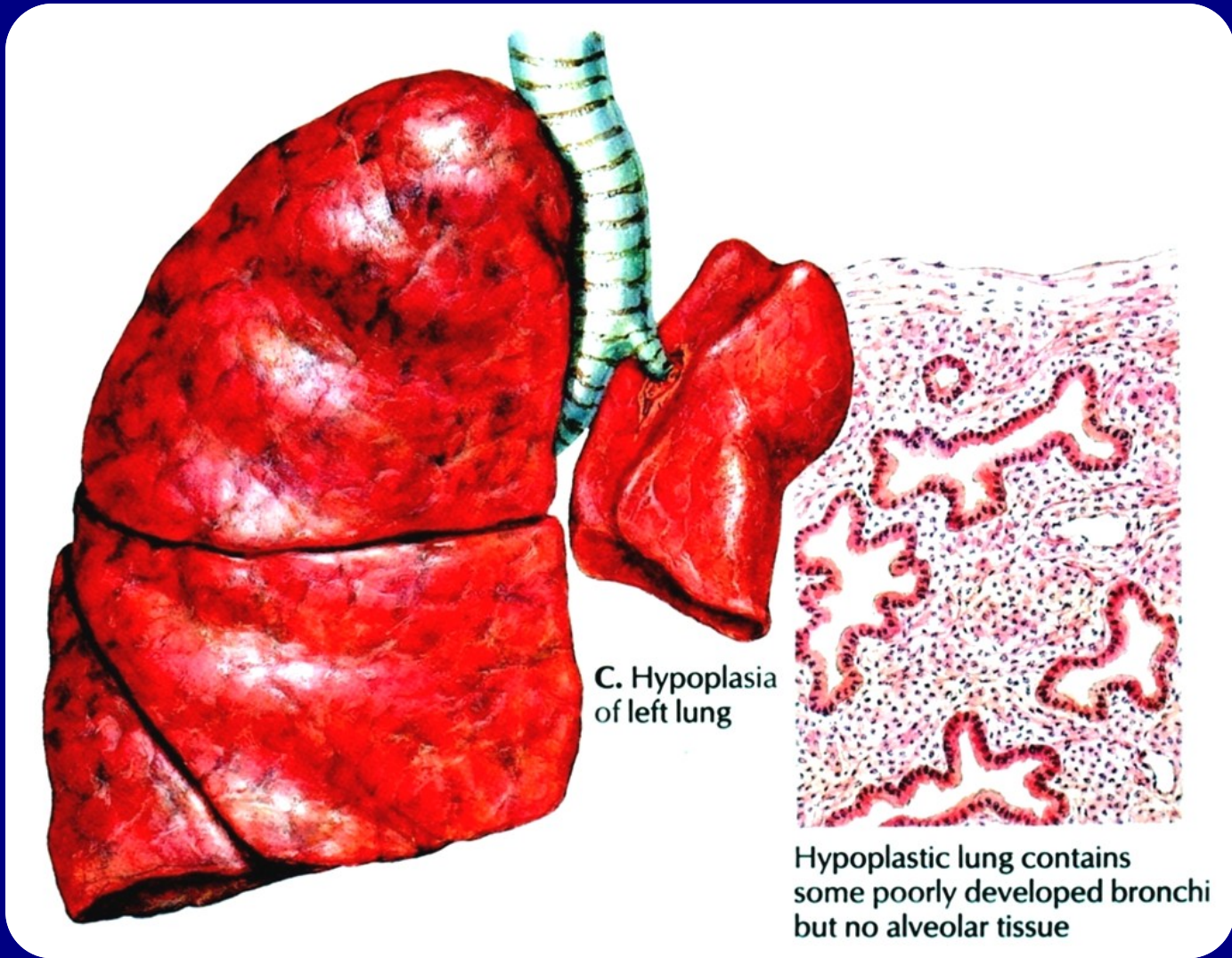


A. Complete unilateral agenesis. Left lung and bronchial tree are absent. Right lung is greatly enlarged with resultant shift of mediastinum to left, elevation of left diaphragm, and approximation of ribs on that side

B. Aplasia of left lung.
Only rudimentary
bronchi on left side,
which end blindly



Airway branching anomalies



Airway branching anomalies