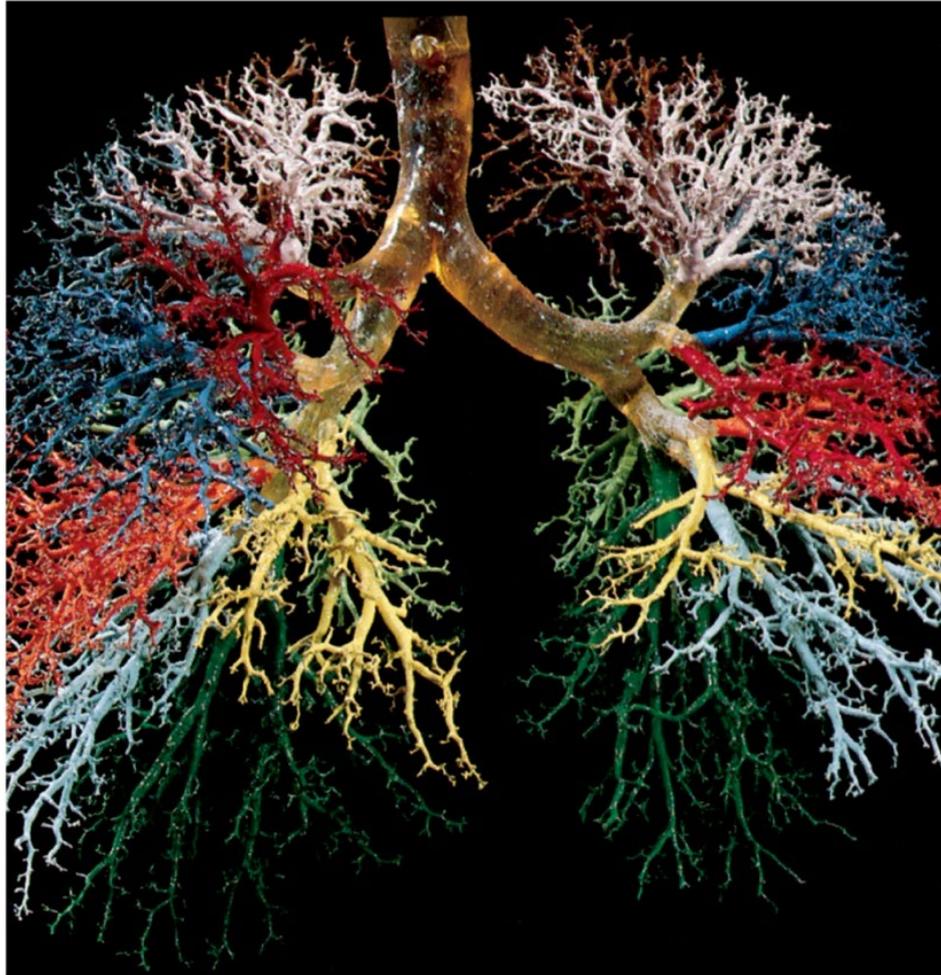


Chapter 22.1

The Respiratory System



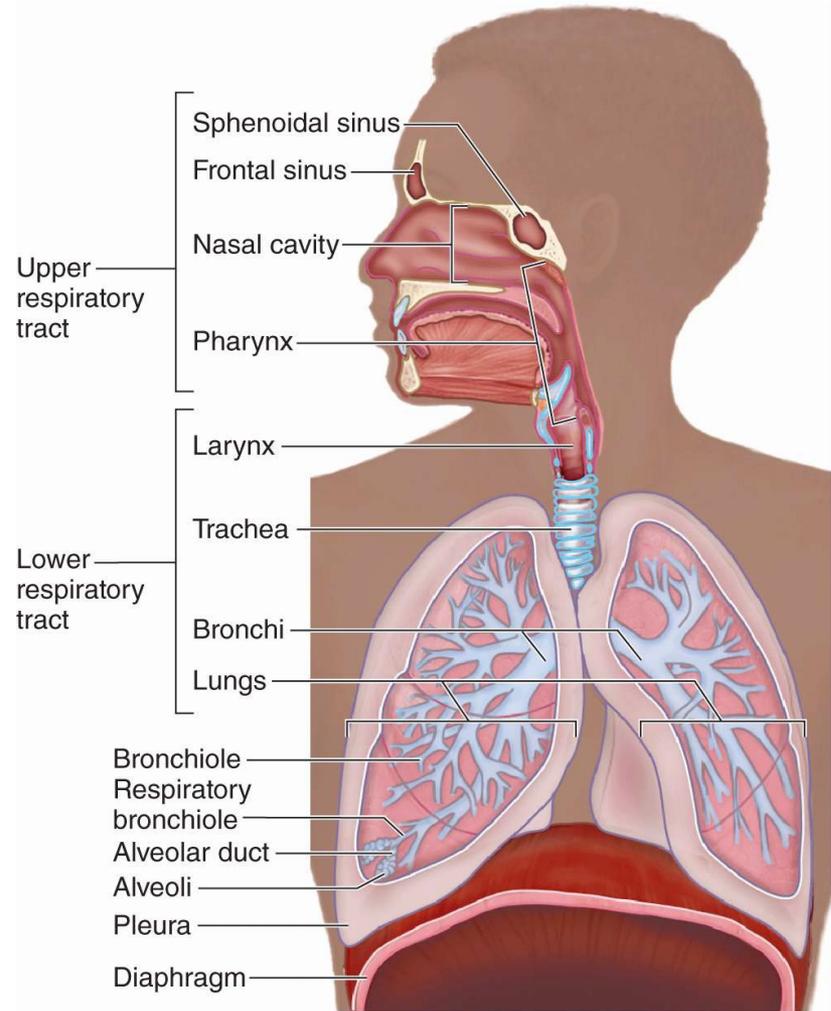
Divisions of the Respiratory System

Upper respiratory tract

- located in head and neck
- nose through laryngopharynx

Lower respiratory tract

- organs of the thorax
- larynx through alveoli



Conducting Division VS Respiratory Division

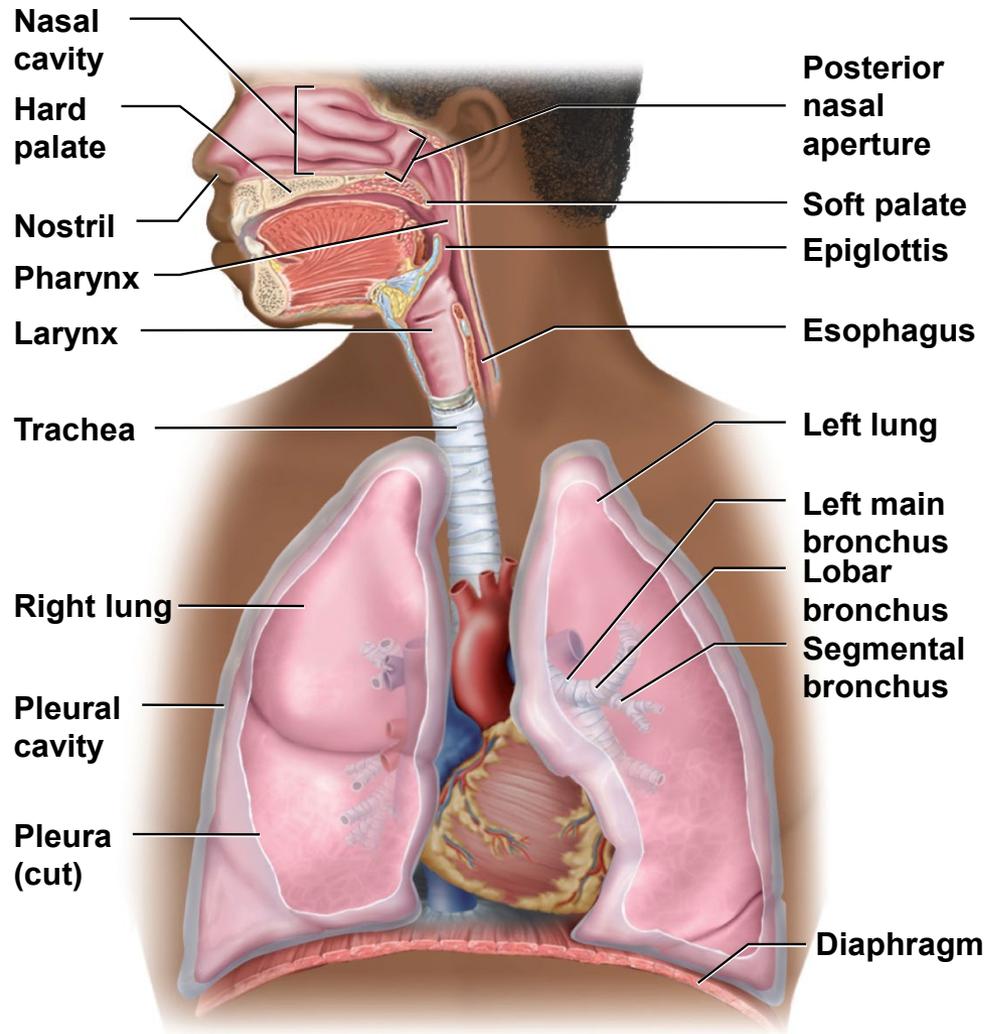
Conducting division of the respiratory system

- > those passages that serve only for airflow
- > no gas exchange
- > nostrils through major bronchioles

Respiratory division consists of alveoli and other gas exchange regions

- respiratory membrane // space between alveoli and capillary
- incoming air stops at the alveoli of the lungs
- 300 millions alveoli (150 million per lung)
- gases inside alveoli are exchanged with the gases in the bloodstream
- gas molecules (O₂ and CO₂) diffuse “down” concentration gradients
- 70 and 80 square meters of surface area

Organs of Respiratory System



nose, pharynx, larynx, trachea, bronchi, lungs

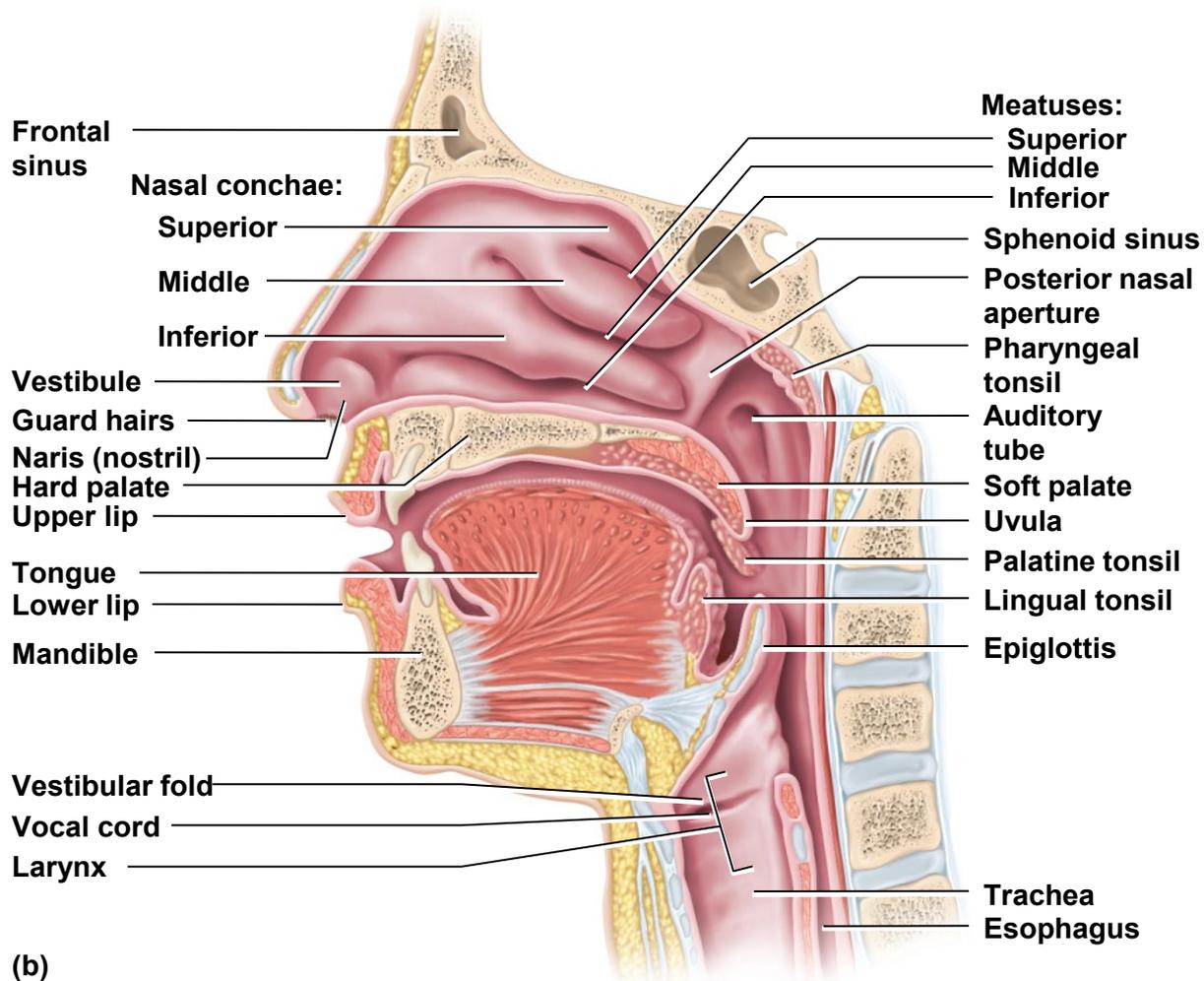
Nasal Cavity

Nasal conchae

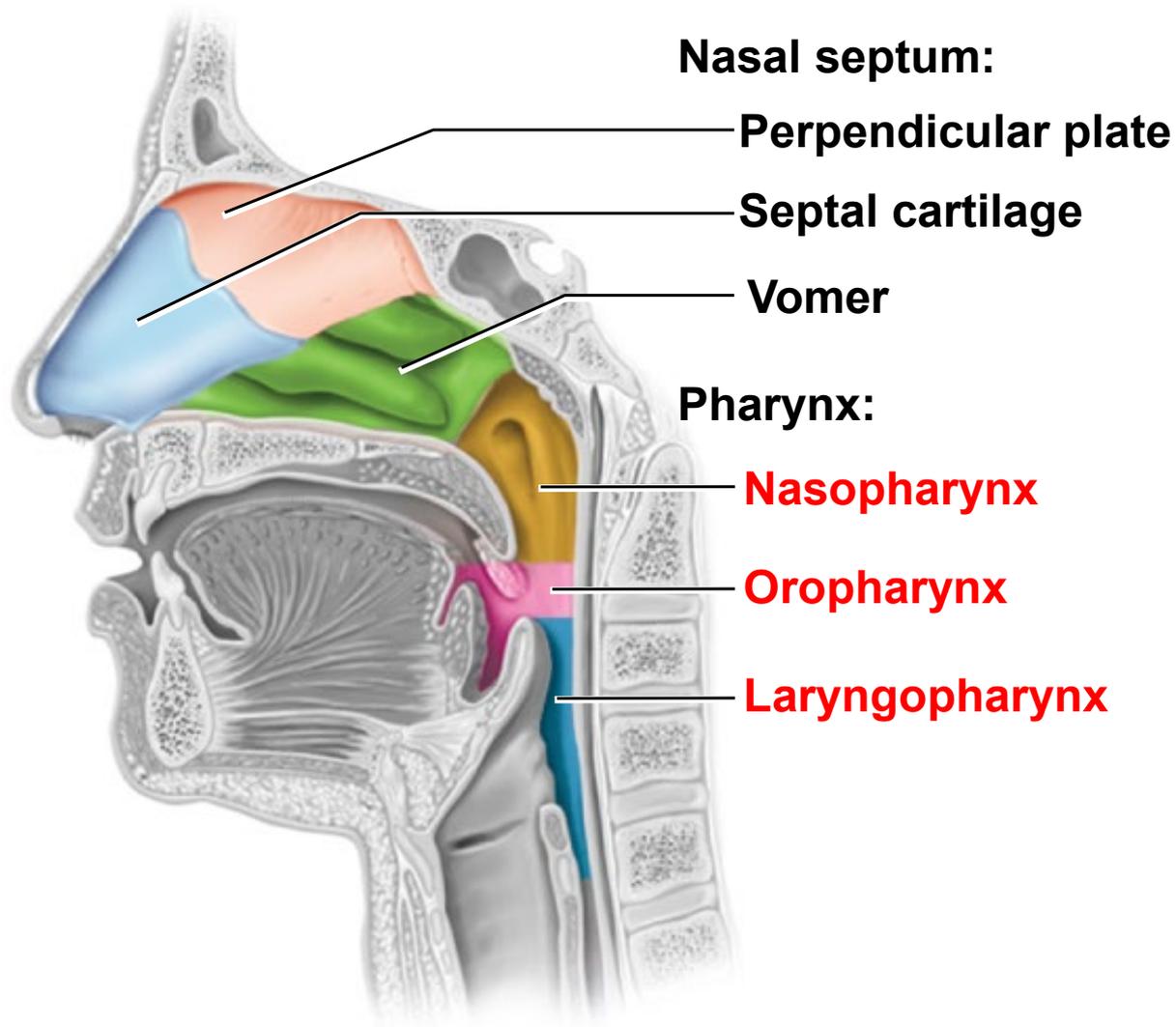
- nasal cavity occupied by three folds of tissue
- **superior, middle, and inferior nasal conchae** (turbinates) // project from lateral walls toward septum
 - **meatus** – narrow air passage beneath each concha // narrowness and turbulence ensure that most air contacts mucous membranes
- **meatus function = cleans, warms, and moistens the air**

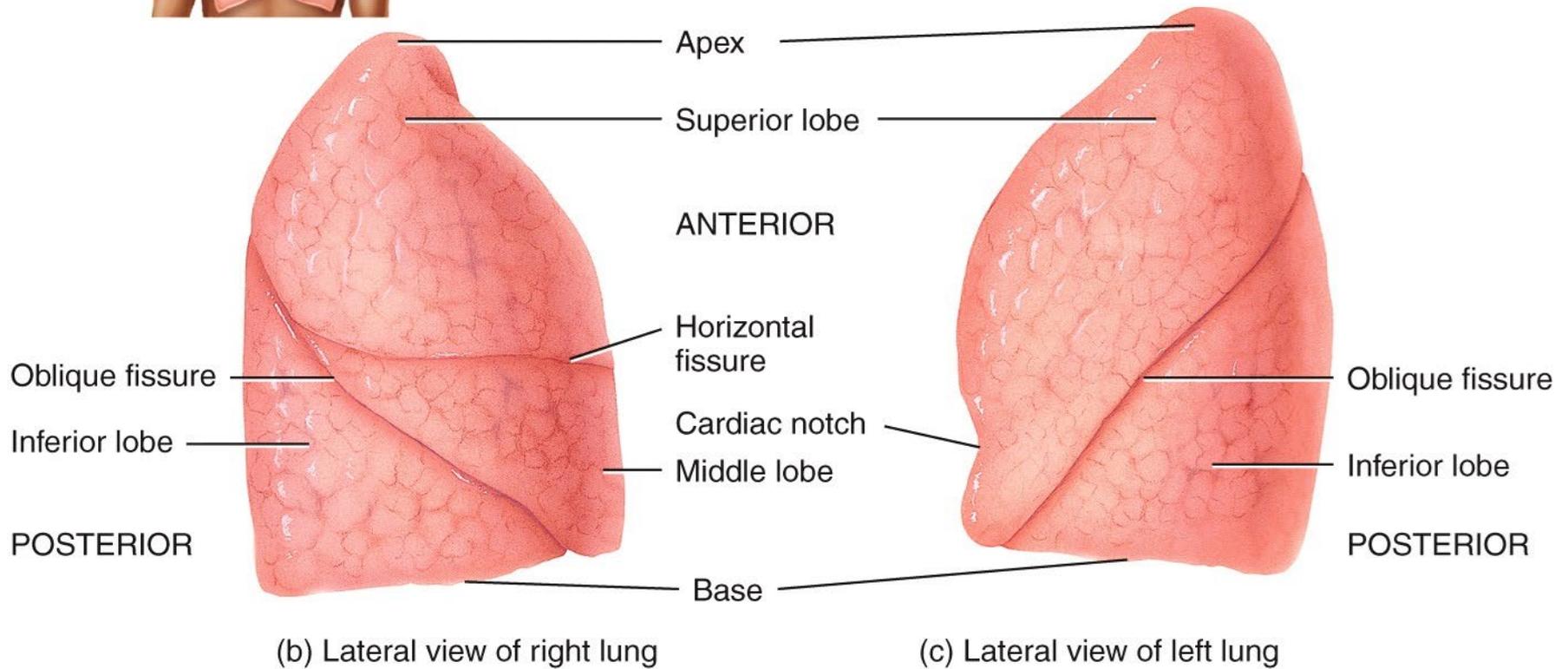
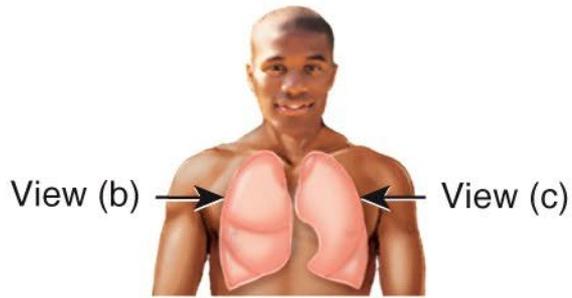
Upper Respiratory Tract

(respiratory organs in the head and neck / stop at entrance to larynx)



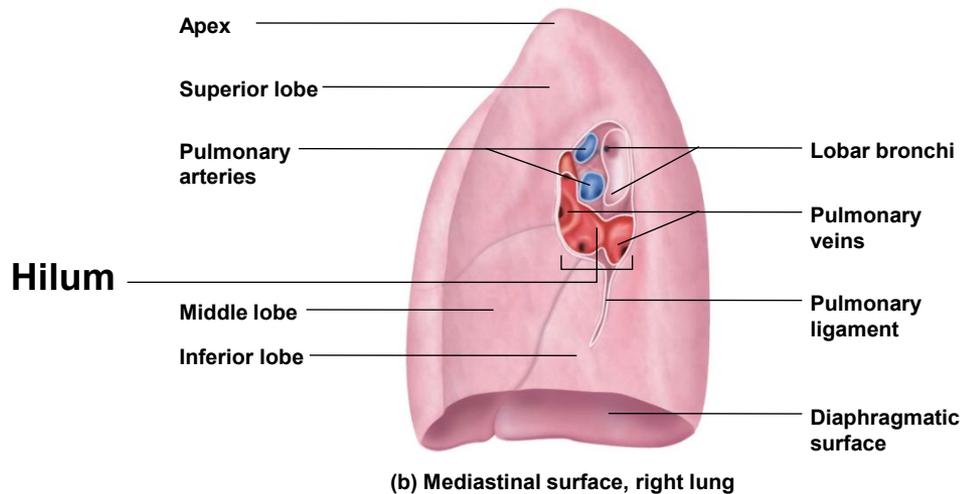
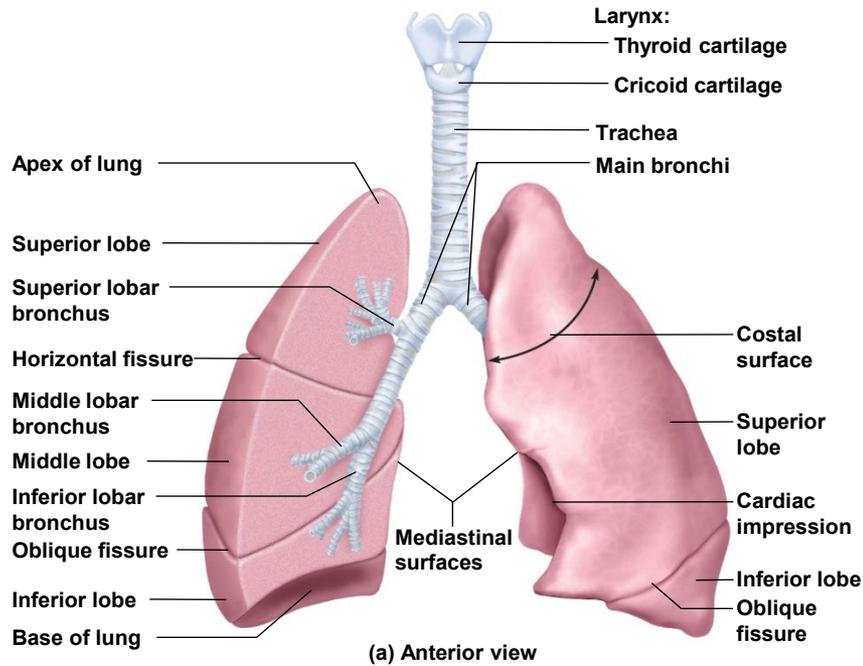
Regions of Pharynx





- Right lung three lobes and left lung two lobes
- Note: oblique fissures important lab learning objectives

Lungs - Surface Anatomy



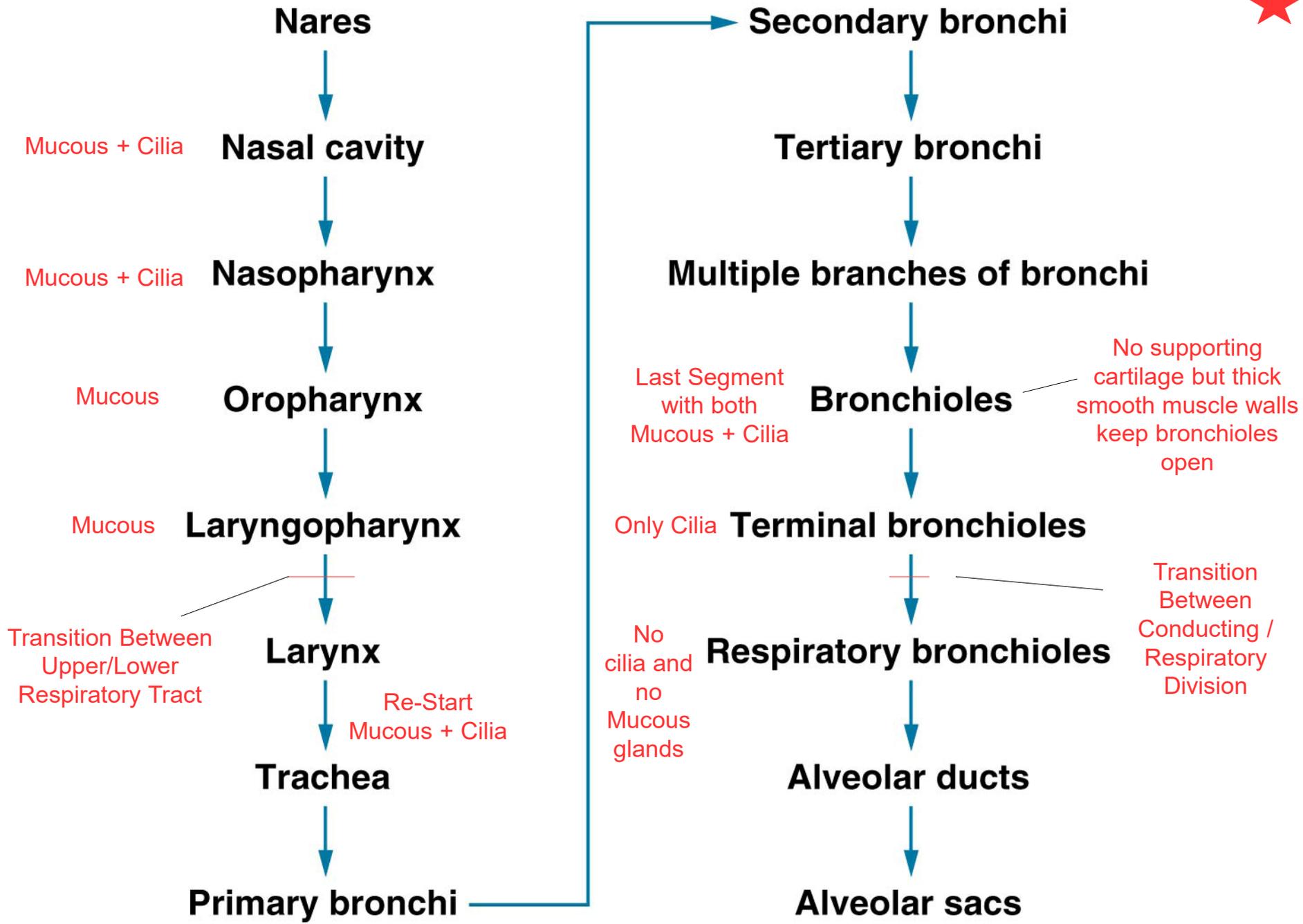
Path of Air Flow

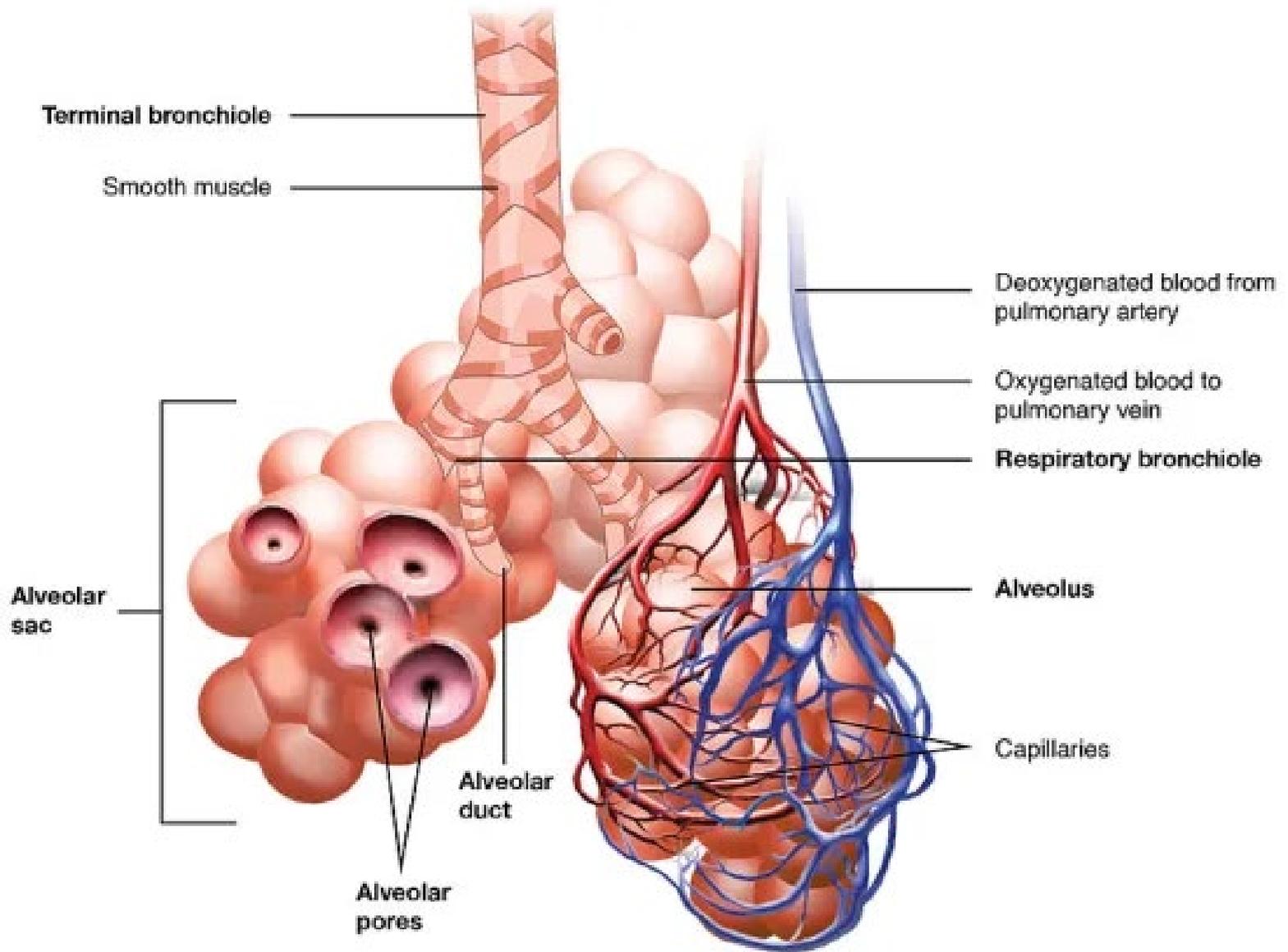
Conducting Division

nasal cavity
pharynx
larynx
trachea
main bronchus
lobar bronchus
segmental bronchus
Bronchiole
Terminal bronchiole

Respiratory Division

respiratory bronchiole
alveolar duct
atrium
alveolus





- Respiratory membrane formed by union of alveolar and capillary epithelial cells. These cells share a common basement membrane.

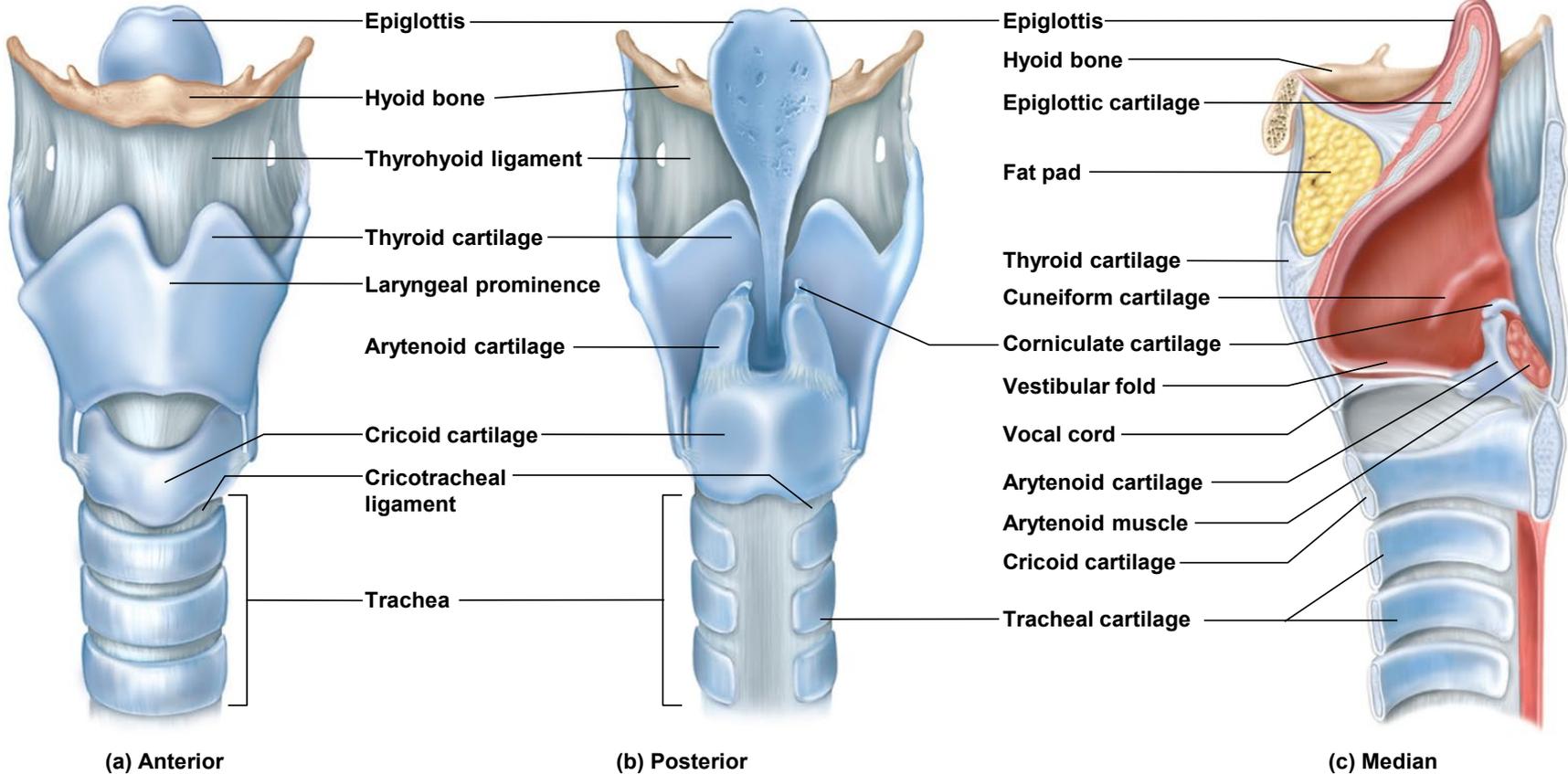
Larynx

Larynx (voice box) – cartilaginous chamber about 4 cm (1.5 in.)

- primary function is to keep food and drink out of the airway
- has evolved additional role – phonation – production of sound
- epiglottis – flap of tissue that guards the superior opening of the larynx
 - at rest, stands almost vertically
 - during swallowing, extrinsic muscles of larynx pull larynx upward
 - tongue pushes epiglottis down to meet it
 - closes airway and directs food to the esophagus behind it
 - **vestibular folds** of the larynx play greater role in keeping food and drink out of the airway VS **vestibular ligaments (also called vocal cords)**

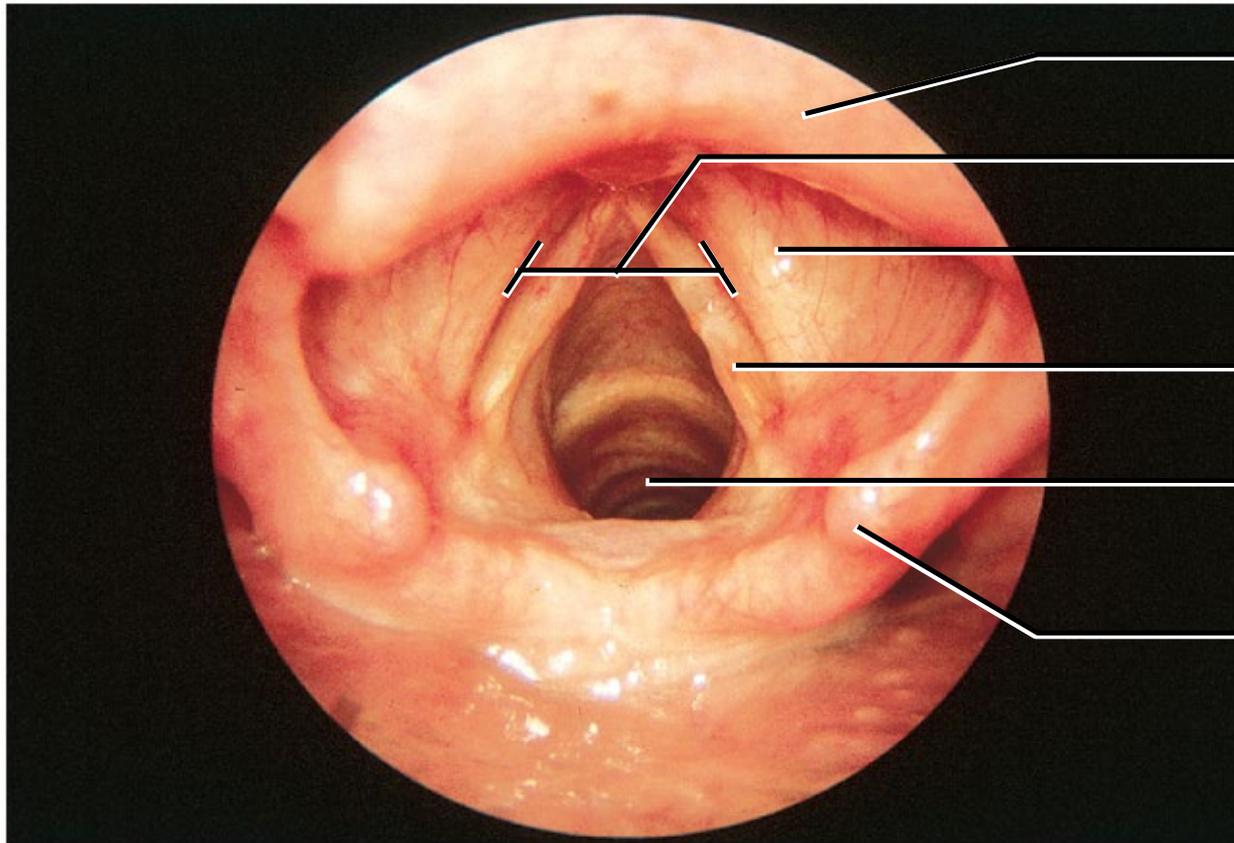
Views of Larynx

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Endoscopic View of the Larynx

Anterior



Epiglottis

Glottis

Vestibular fold

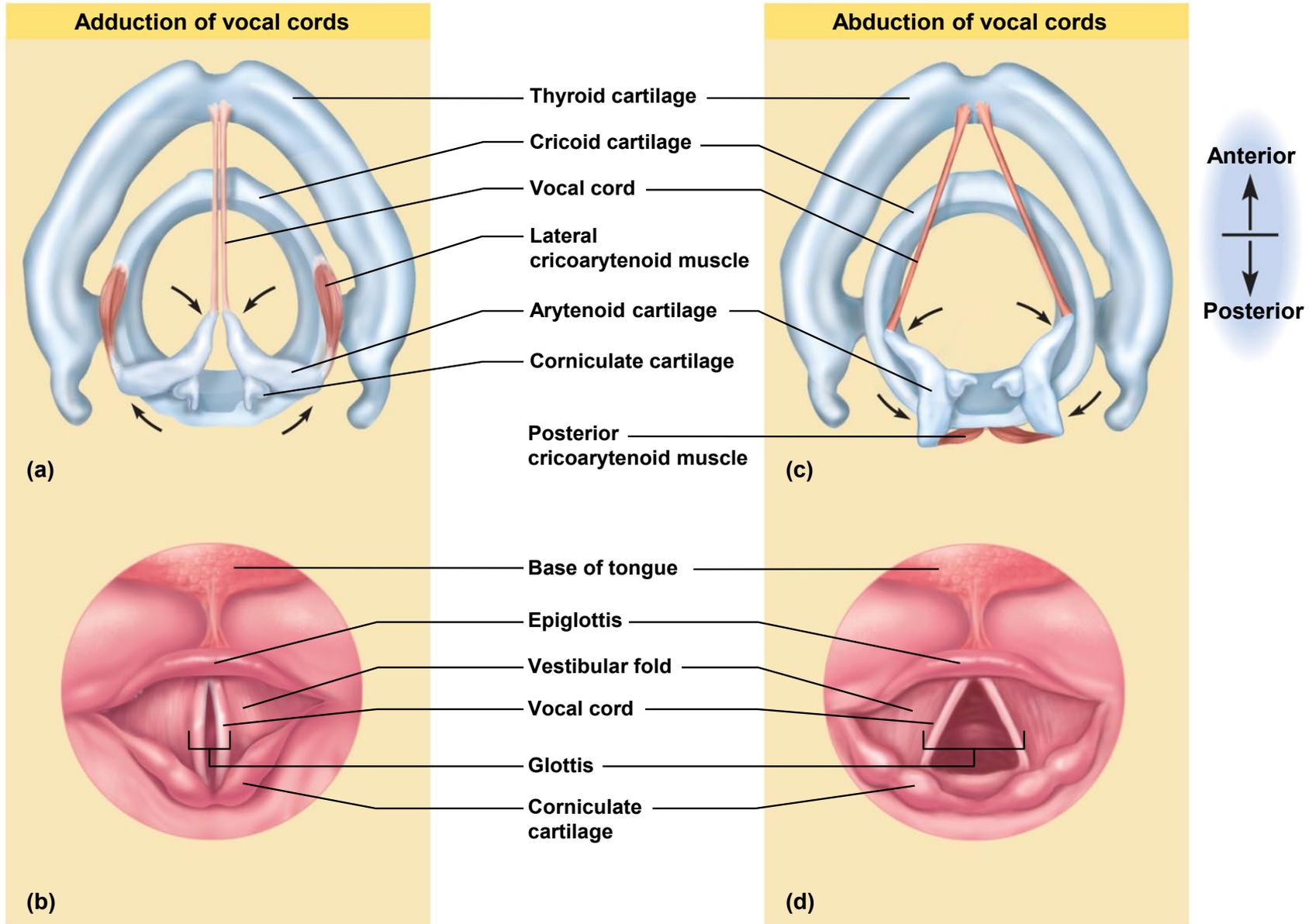
Vocal cord

Trachea

**Corniculate
cartilage**

Posterior

Action of Vocal Cords



Trachea (1 of 2)

Trachea (windpipe) – a rigid tube about 12 cm (4.5 in.) long and 2.5 cm (1 in.) in diameter

anterior to esophagus

supported by 16 to 20 **C-shaped** rings of **hyaline cartilage**

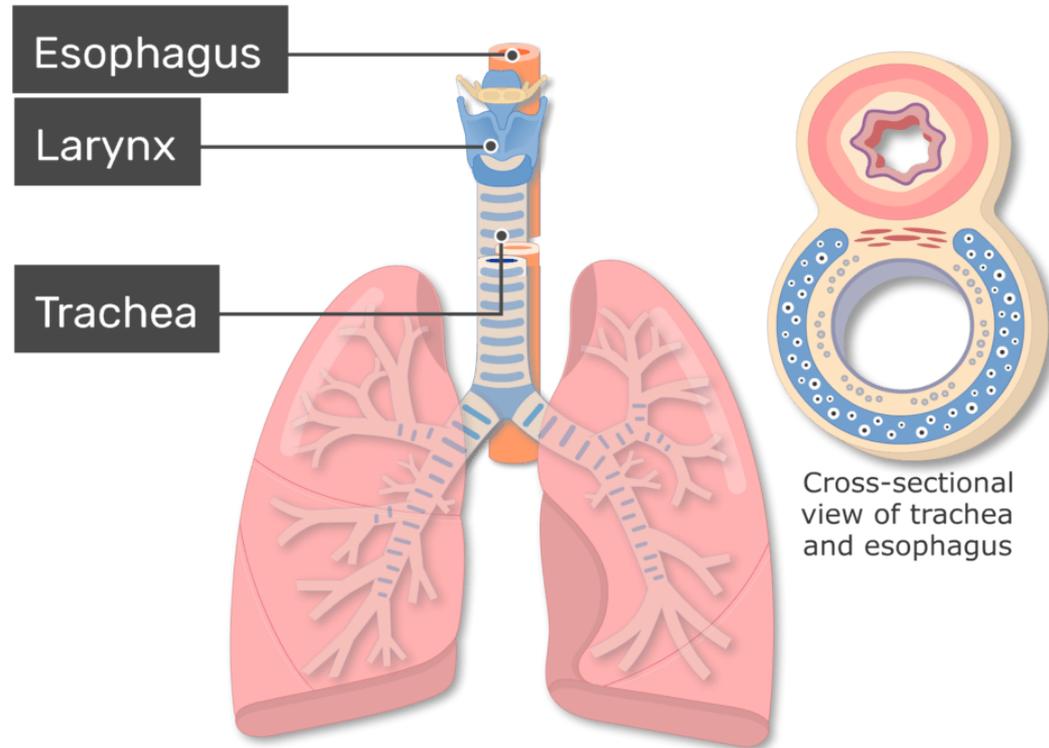
reinforces the trachea and keeps it from collapsing when you inhale

opening in rings faces posteriorly towards esophagus

trachealis muscle spans opening in rings

gap in C allows room for the esophagus to expand as swallowed food passes by

contracts or relaxes to adjust air flow



Trachea (2 of 2)

inner lining of trachea is a **ciliated pseudostratified columnar epithelium**

composed mainly of mucus-secreting cells, ciliated cells, and stem cells

mucociliary escalator – mechanism for debris removal

mucus traps inhaled particles

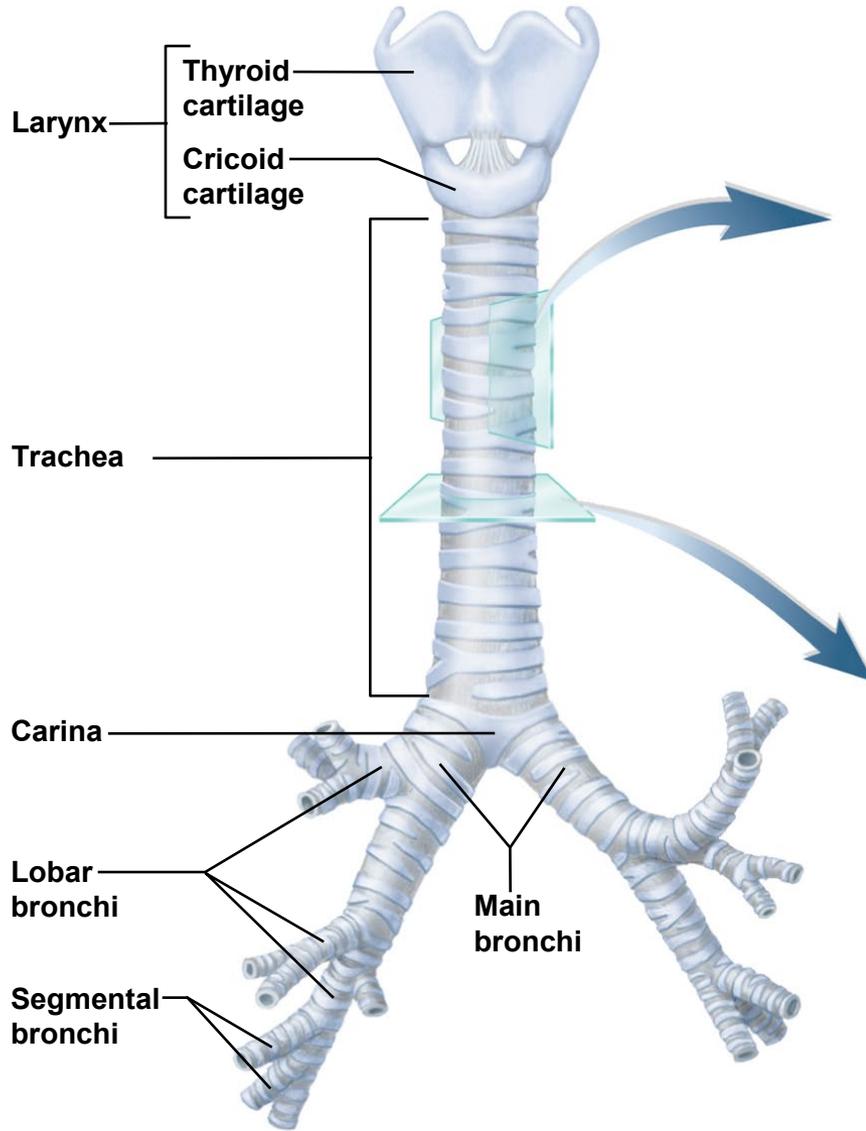
upward beating cilia drives mucus toward pharynx where it is swallowed

right and left main bronchi

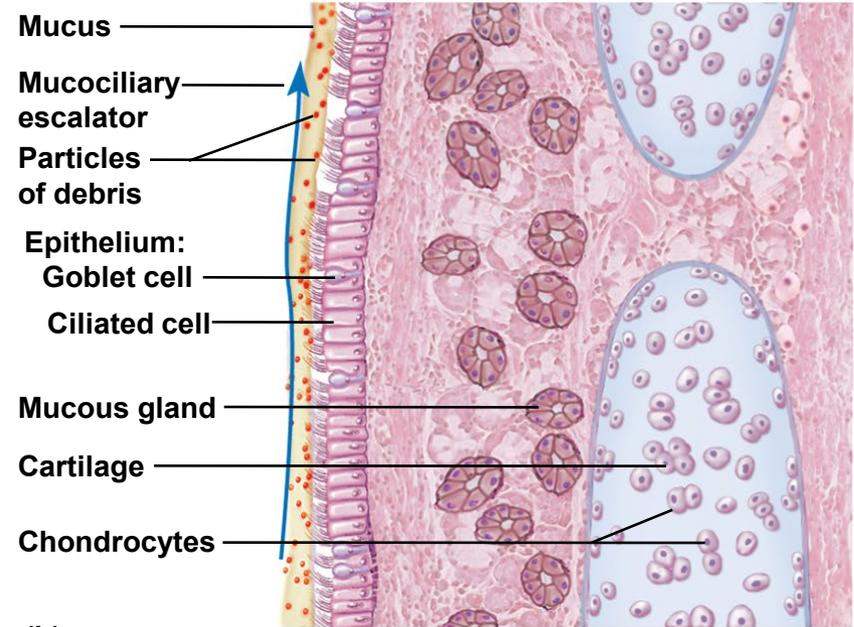
trachea forks at level of sternal angle /// **carina** – internal medial ridge in the lowermost tracheal cartilage

directs the airflow to the right and left

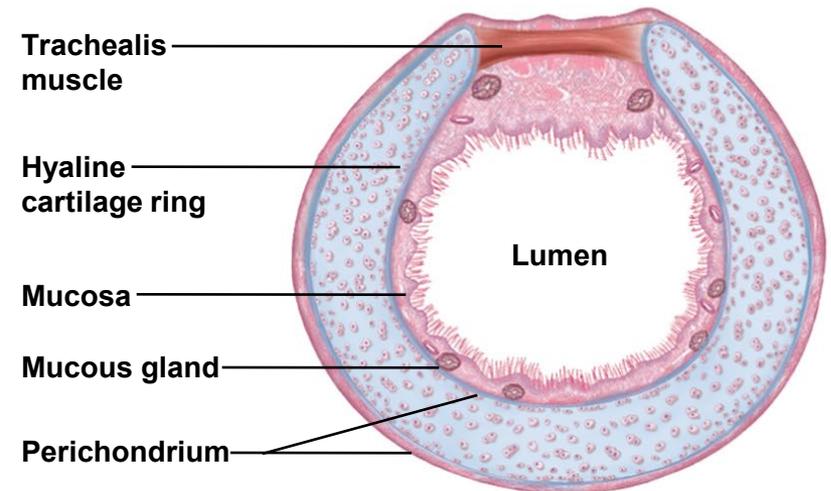
Lower Respiratory Tract



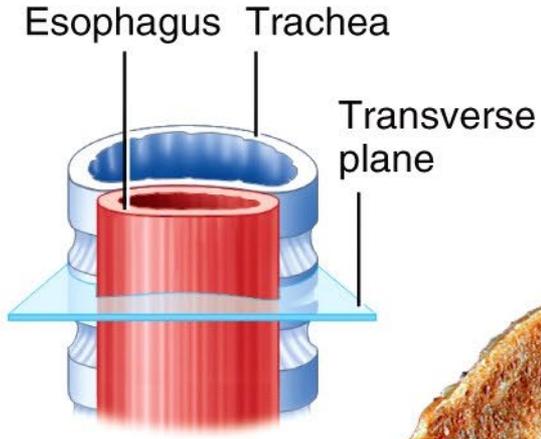
(a)



(b)



(c)



ANTERIOR

Left lateral lobe of thyroid gland

Tracheal cartilage

Right lateral lobe of thyroid gland

Trachealis muscle in fibromuscular membrane

Esophagus

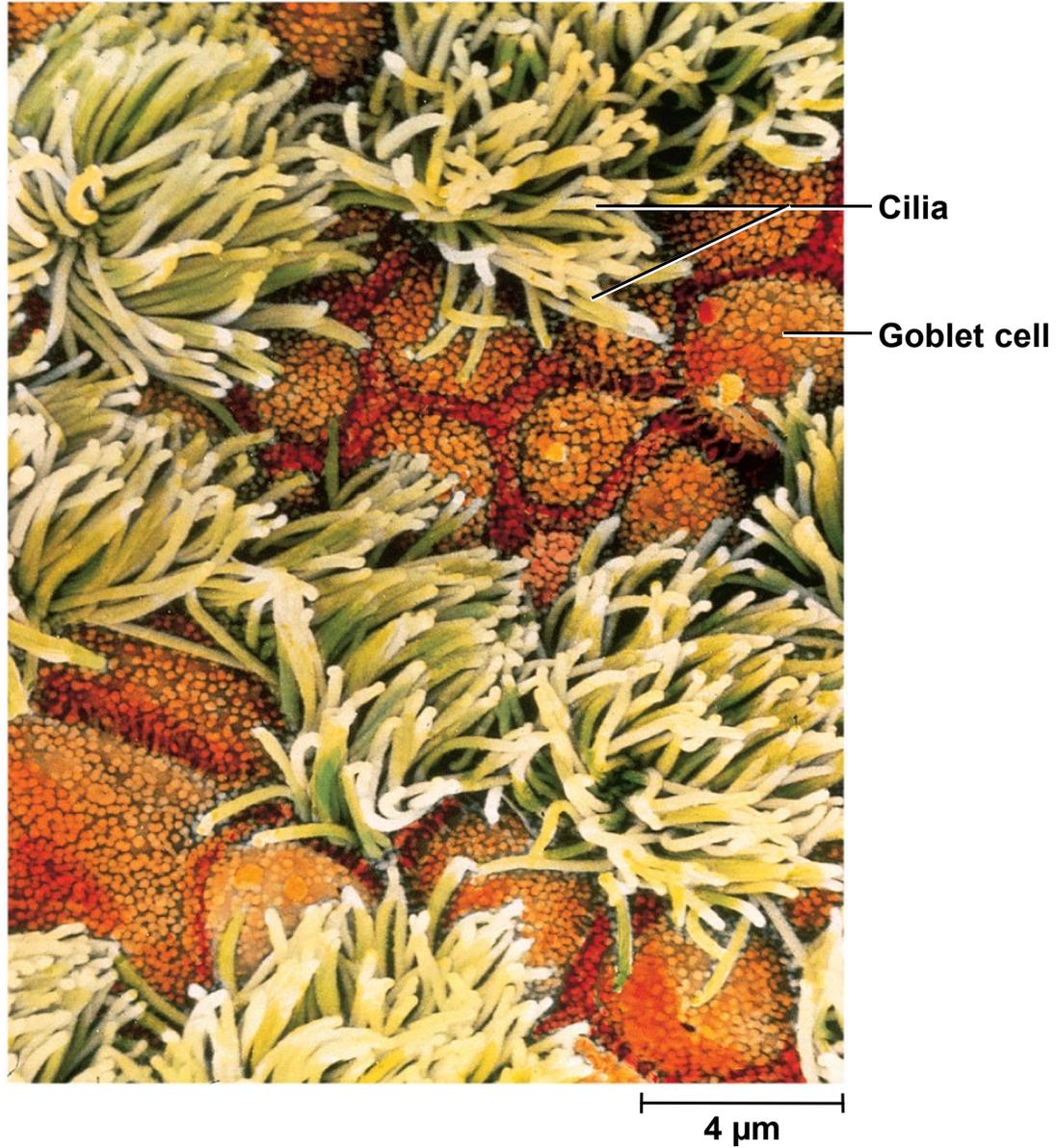
Dissection Shawn Miller,
Photograph Mark Nielsen

POSTERIOR

Superior view of transverse section of thyroid gland, trachea, and esophagus

Tracheal Epithelium

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Blood Vessels Associated with Lung Tissue

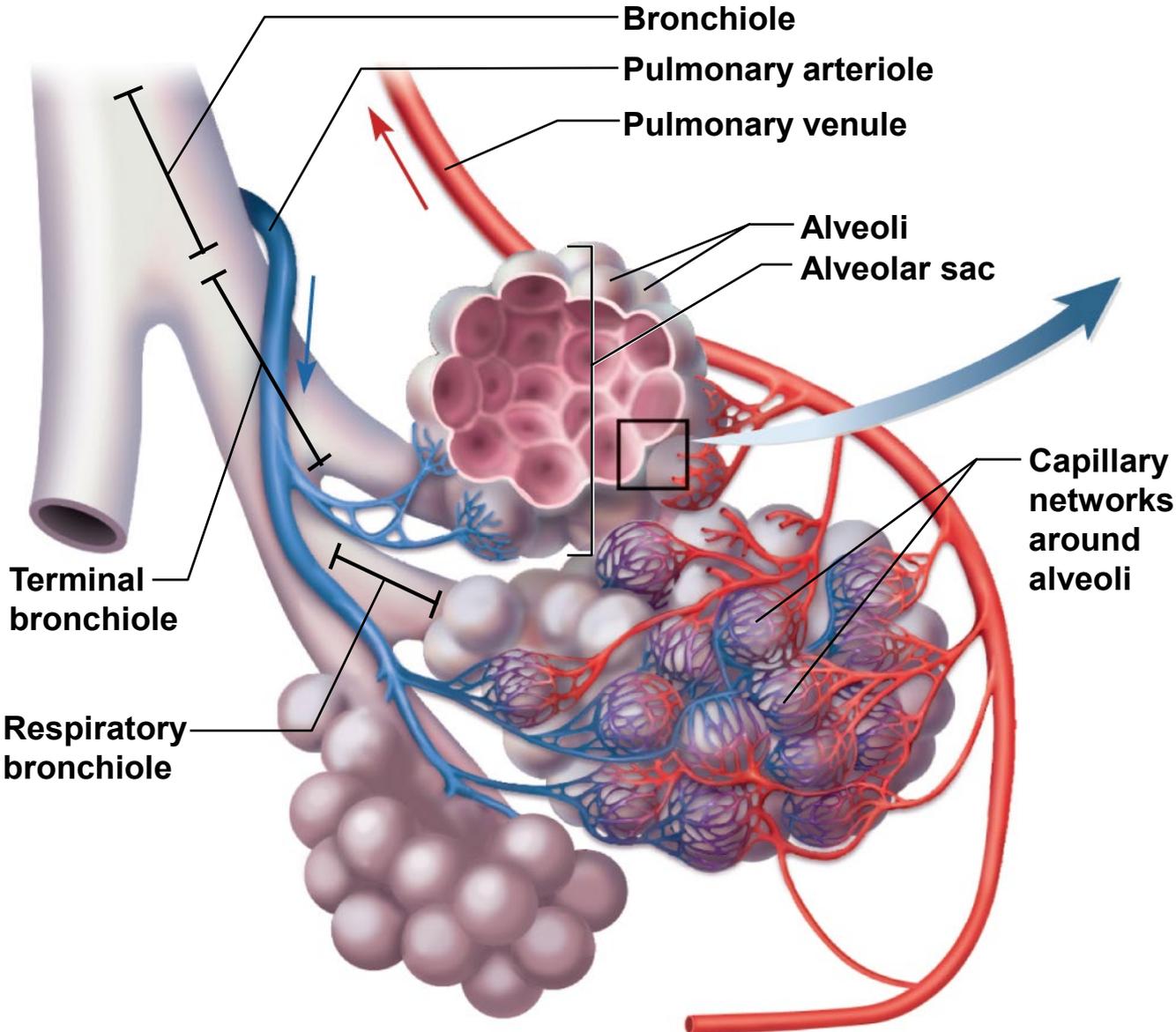
Pulmonary arteries & veins

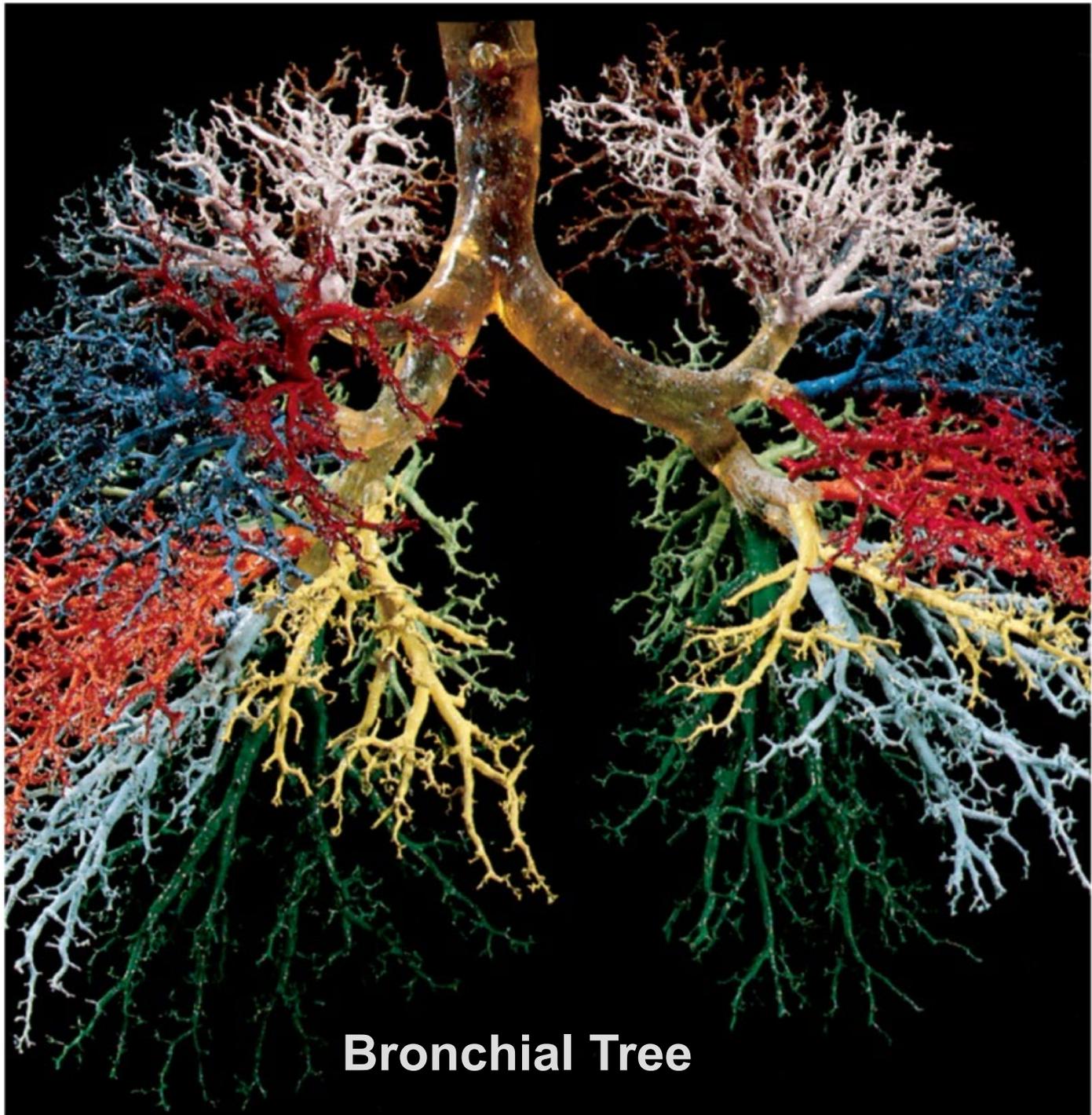
These blood vessels closely follow the bronchial tree (air passageways) as they make their way to the alveoli /// pulmonary circuit = blood flow from right ventricle to left atria // **pulmonary circuit**

Bronchial arteries & veins

These blood vessels branch of thoracic aorta /// delivers oxygen and removes waste products from lung tissues /// **systemic circuit**

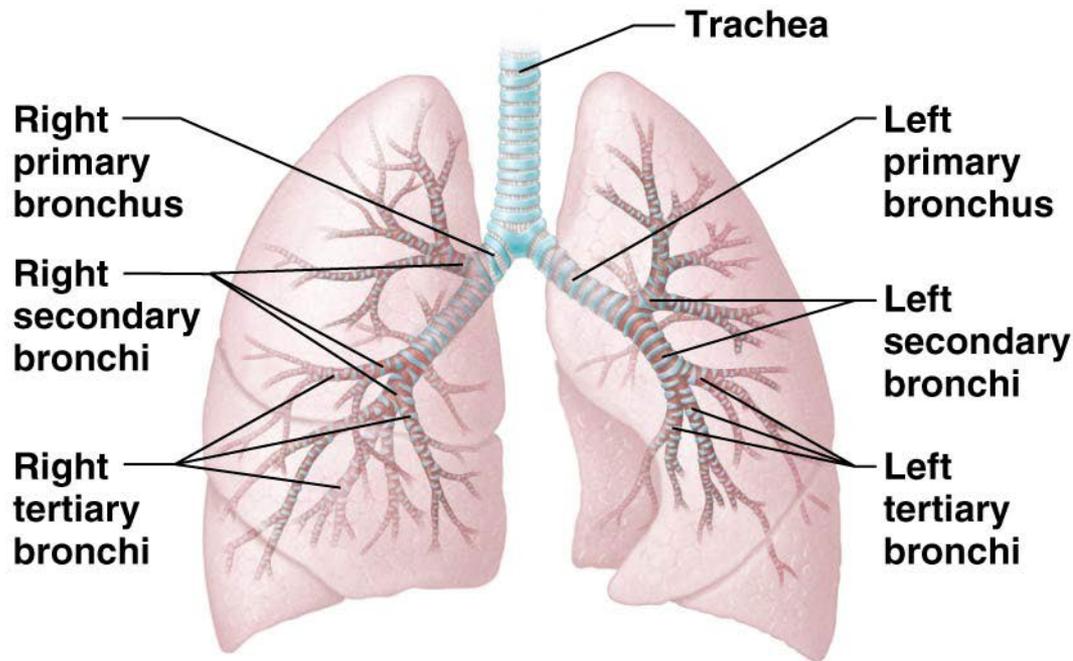
Blood Supply to the Alveolar Sacs



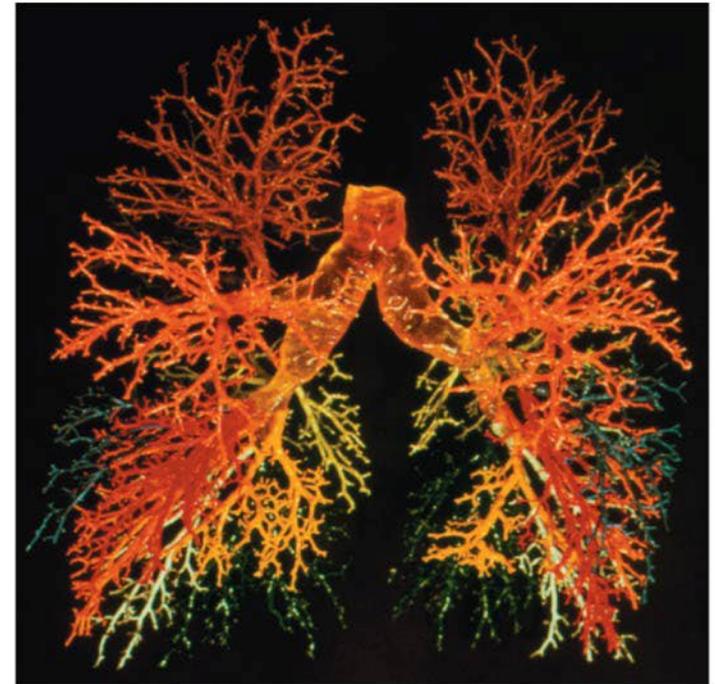


Bronchial Tree

Branching pattern of the bronchial tree.



(a) Conducting zone passages and bronchial tree



(b) Cast of bronchial tree

Main Brochi = Primary Bronchi // Secondary = Lobar // Tertiary = Segmental Within Lobes

Bronchial Tree

(Main Bronchi / Lobar Bronchi / Segmental Bronchi)

Bronchial tree – a branching system of air tubes located in each lung /// from one main bronchus extends 65,000 terminal bronchioles

Main bronchi (primary) – supported by c-shaped hyaline cartilage rings /// rt. main bronchus is a 2-3 cm branch arising from fork of trachea

–Right bronchus slightly wider and more vertical than left // this is why aspirated (inhaled) foreign objects lodge right bronchus more often the left

Left main bronchus is about 5 cm long /// slightly narrower and more horizontal than the right // less likely for food to end up in left bronchus

Bronchial Tree

(Main Bronchi / Lobar Bronchi / Segmental Bronchi)

Lobar bronchi (secondary) – supported by crescent shaped cartilage plates

three right lobar bronchi – superior, middle, and inferior /// one to each lobe of the right lung

two left lobar bronchi - superior and inferior /// one to each lobe of the left lung

Segmental (tertiary) bronchi - supported by crescent shaped cartilage plates

10 on right, and 8 on left

bronchopulmonary segment –functionally independent unit of the lung tissue

Bronchial Tree

(Main Bronchi / Lobar Bronchi / Segmental Bronchi)

All segments of the bronchial tree including the “ bronchioles” are lined with a mucosa lined with ciliated pseudostratified columnar epithelial cells

lamina propria = connective tissue /// has an abundance of mucous glands and lymphocyte nodules

bronchus-associated lymphoid tissue = BALT /// immediately deep to epithelium

all divisions of bronchial tree have a large amount of elastic connective tissue /// contributes to the recoil that expels air from lungs

the mucosa also has a well-developed layer of smooth muscle /// **muscularis mucosae** which contracts or relaxes to constrict or dilate the airway, regulating air flow

The Distal Bronchiole Tree

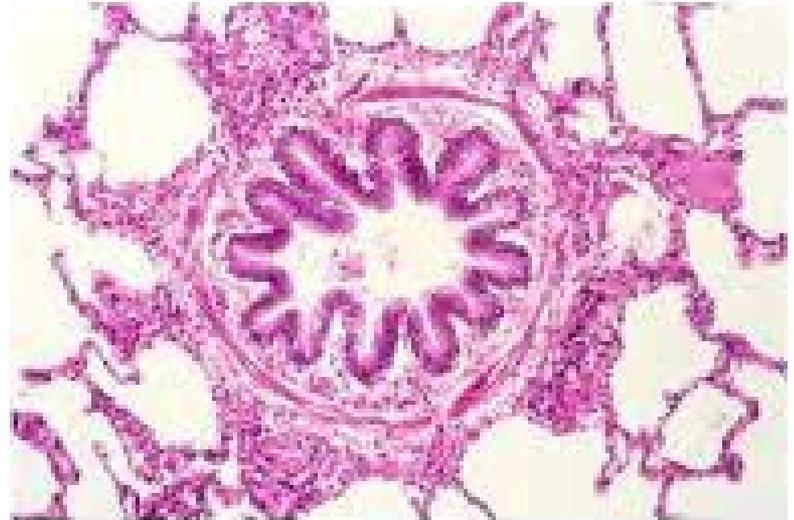
(Bronchioles / Terminal Bronchioles / Respiratory Bronchioles)

Bronchioles

Lack cartilage // thick smooth muscle walls prevent bronchioles from collapsing from negative pressure // only 1mm diameter

Pulmonary lobule = portion of lung ventilated by one bronchiole

Bronchioles have ciliated cuboidal epithelium and mucous glands



Terminal bronchioles have cilia but no mucus glands

Respiratory bronchioles do not have cilia and no mucous glands

Why does this make sense?

The Distal Bronchiole Tree

(Bronchioles / Terminal Bronchioles / Respiratory Bronchioles)

Terminal bronchioles

divides into 50 - 80 branches

final branches of conducting division

measure 0.5 mm or less in diameter

no mucous glands or goblet cells

but still have cilia // move mucus draining into them from bronchioles towards larynx by mucociliary escalator (last segment with cilia)

each terminal bronchiole gives off two or more smaller respiratory bronchioles

The Distal Bronchiole Tree

(Bronchioles / Terminal Bronchioles / Respiratory Bronchioles)

Respiratory bronchioles

no cilia and no mucous glands

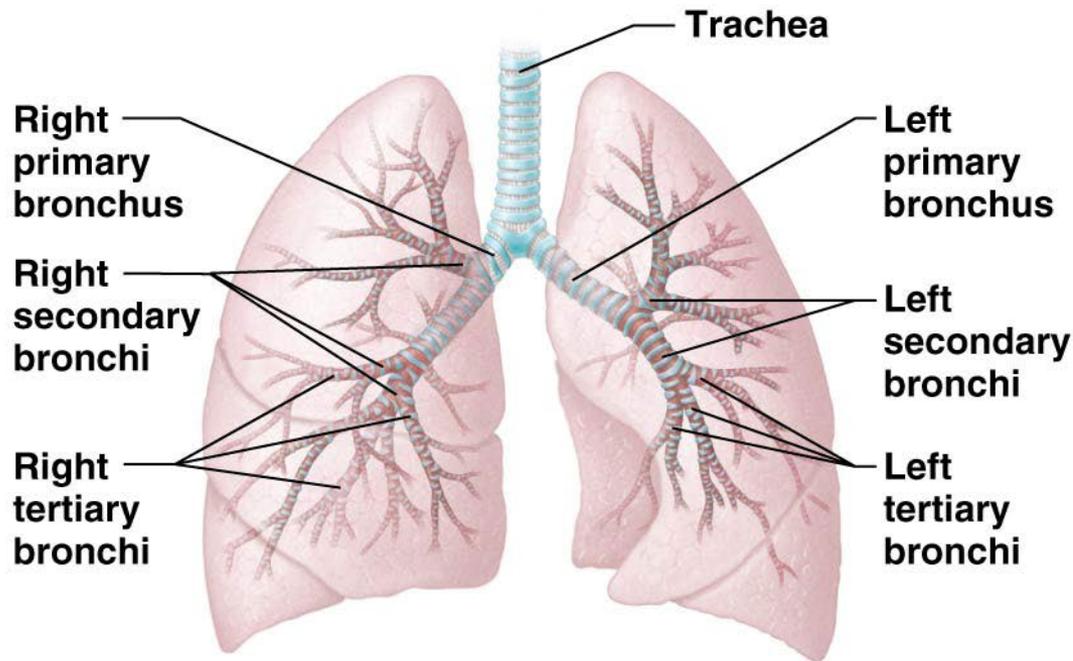
have alveoli budding from their walls

considered the **beginning of the respiratory division** since alveoli participate in gas exchange

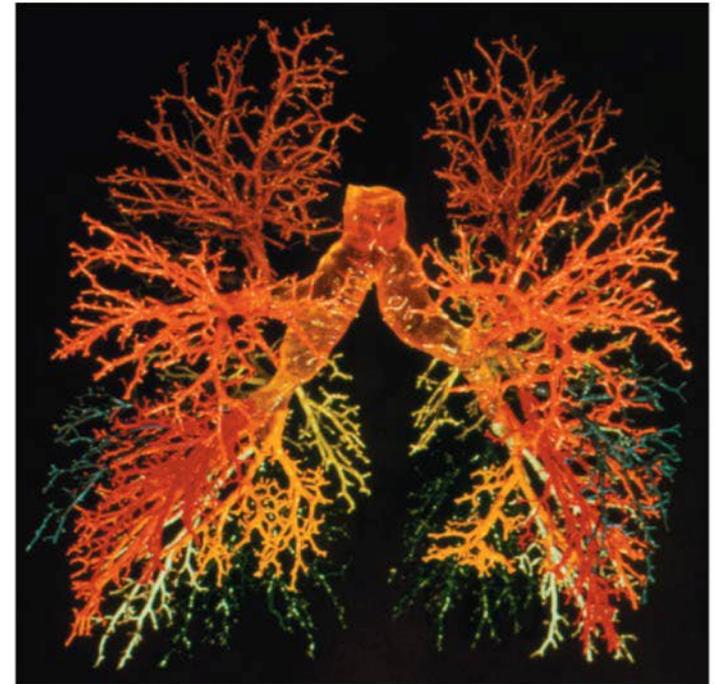
divide into 2-10 **alveolar ducts**

end in **alveolar sacs** – grape-like clusters of alveoli arrayed around a central space called the **atrium**

Branching pattern of the bronchial tree.

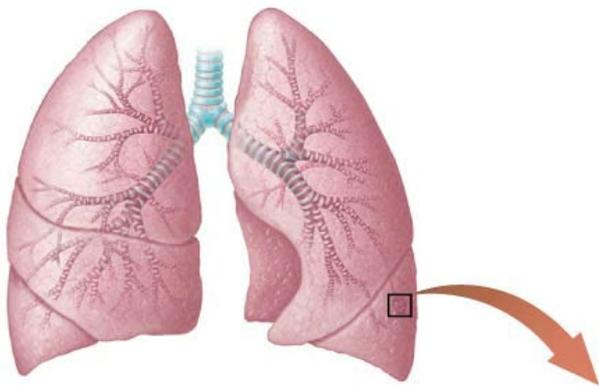


(a) Conducting zone passages and bronchial tree

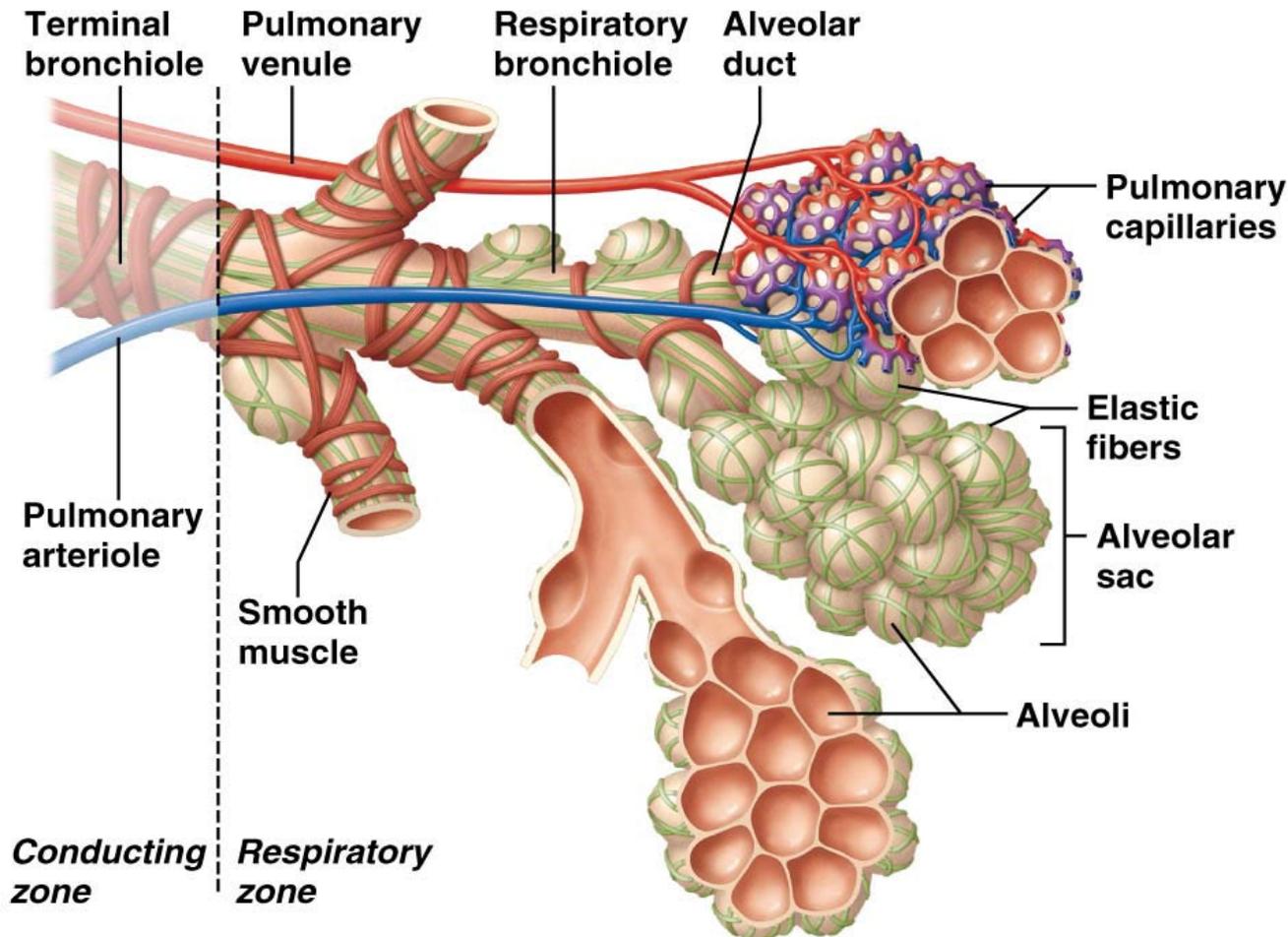


(b) Cast of bronchial tree

Main Brochi = Primary Bronchi // Secondary = Lobar // Tertiary = Segmental

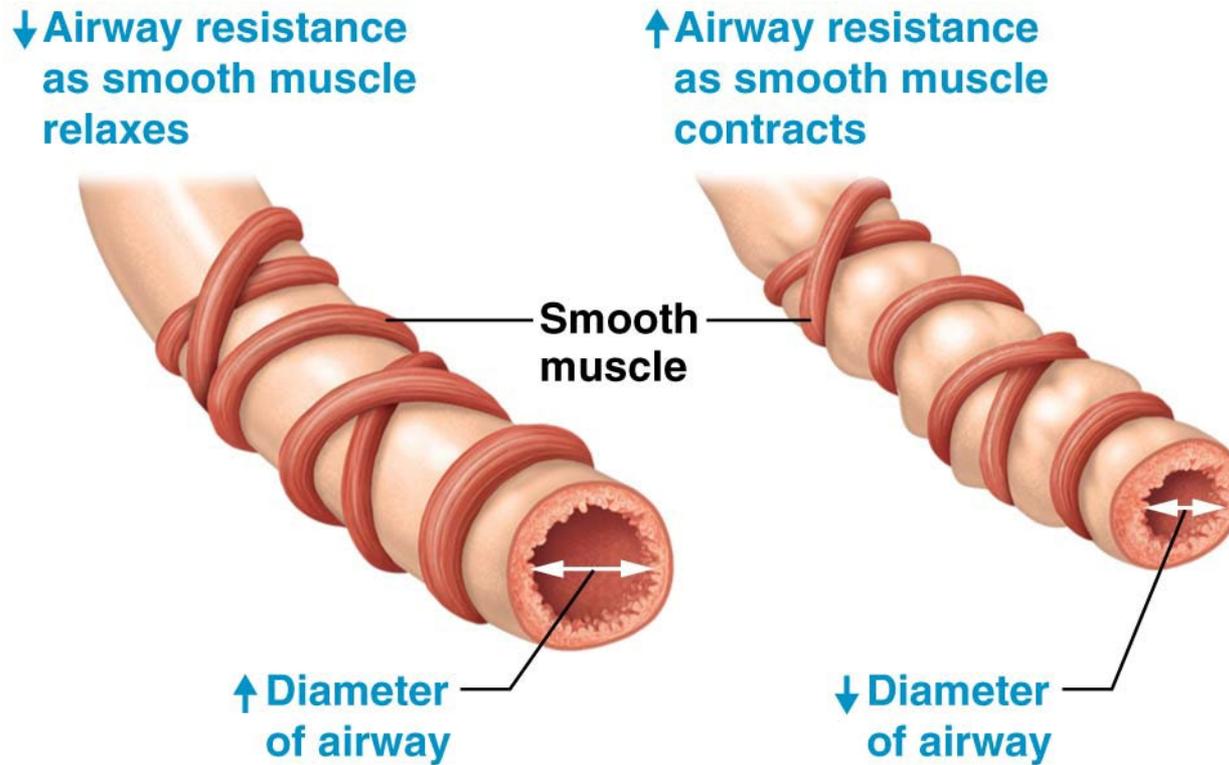


Neutrophils move into the area due to cigarette smoke and release an enzyme called **elastase**. The enzyme breaks down the elastic fibers. Causes reduced surface and gas exchange area.



(a) Structures of the respiratory zone

Relationship between airway resistance and airway diameter.



Histamine constricts bronchioles /// Epinephrine dilates bronchioles

What occurs when histamine is released by mast cells in the systemic tissues?

Three Cell Types of the Alveoli (1 of 3)

150 million **alveoli** in each lung

Providing about 70 m² of surface for gas exchange

Three cell type in the alveolus (type I / type II / dust cells)

Squamous alveolar cells (type I)

thin, broad cells that allow for rapid gas diffusion between alveolus and bloodstream

cover 95% of alveolus surface area

Three Cell Types of the Alveoli (2 of 3)

Great alveolar cells (type II)

round to cuboidal cells that cover the remaining 5% of alveolar surface

repair the alveolar epithelium when the squamous (type I) cells are damaged

secrete **pulmonary surfactant** /// a mixture of phospholipids and proteins that coats the alveoli and prevents them from collapsing when we exhale



Three Cells of the Alveoli (3 of 3)

Alveolar macrophages (dust cells)

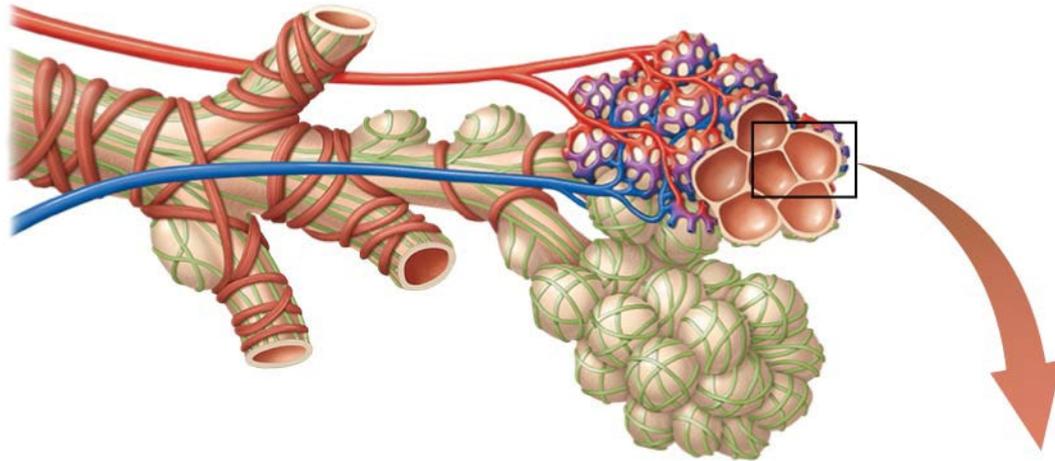
most numerous of all cells in the lung

wander the lumen and the connective tissue between alveoli

keep alveoli free from debris by phagocytizing dust particles

100 million dust cells perish each day as they ride up the mucociliary escalator to be swallowed and digested with their load of debris

Three Alveolar Cell Types

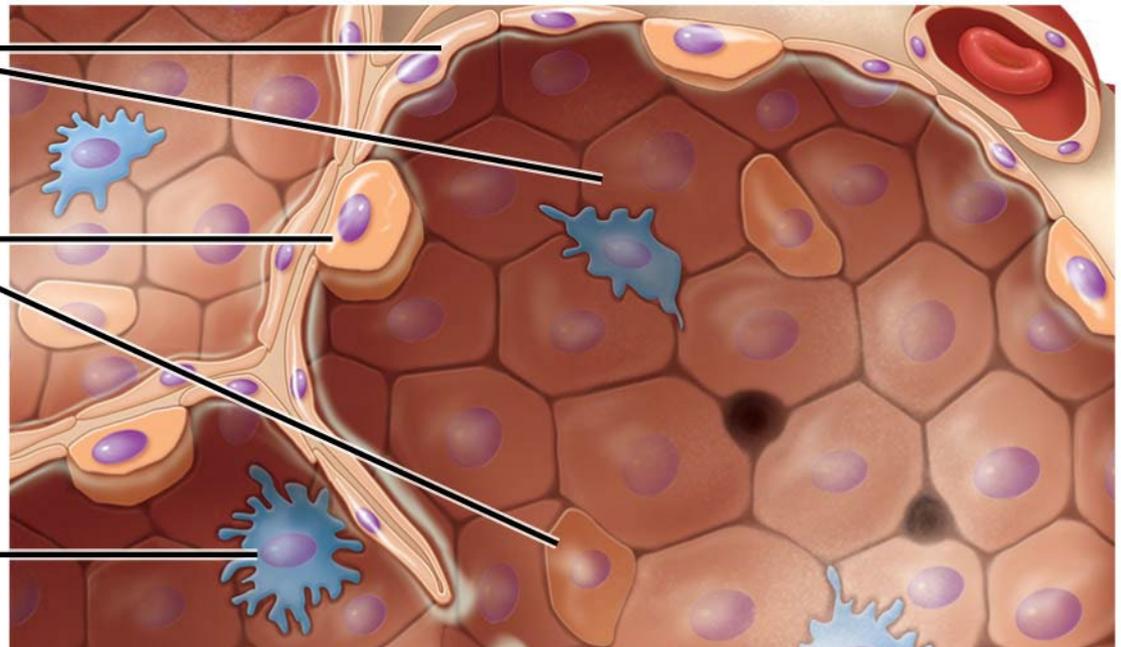


Know the function of each cell type.

Type I
alveolar
cells

Type II
alveolar
cells

Alveolar
macrophage



Respiratory Membrane

Each alveolus surrounded by a basket like capillary bed // network supplied with blood from pulmonary artery

Respiratory membrane – the barrier between the air inside the alveoli and the blood inside the pulmonary capillary

Respiratory membrane consists of

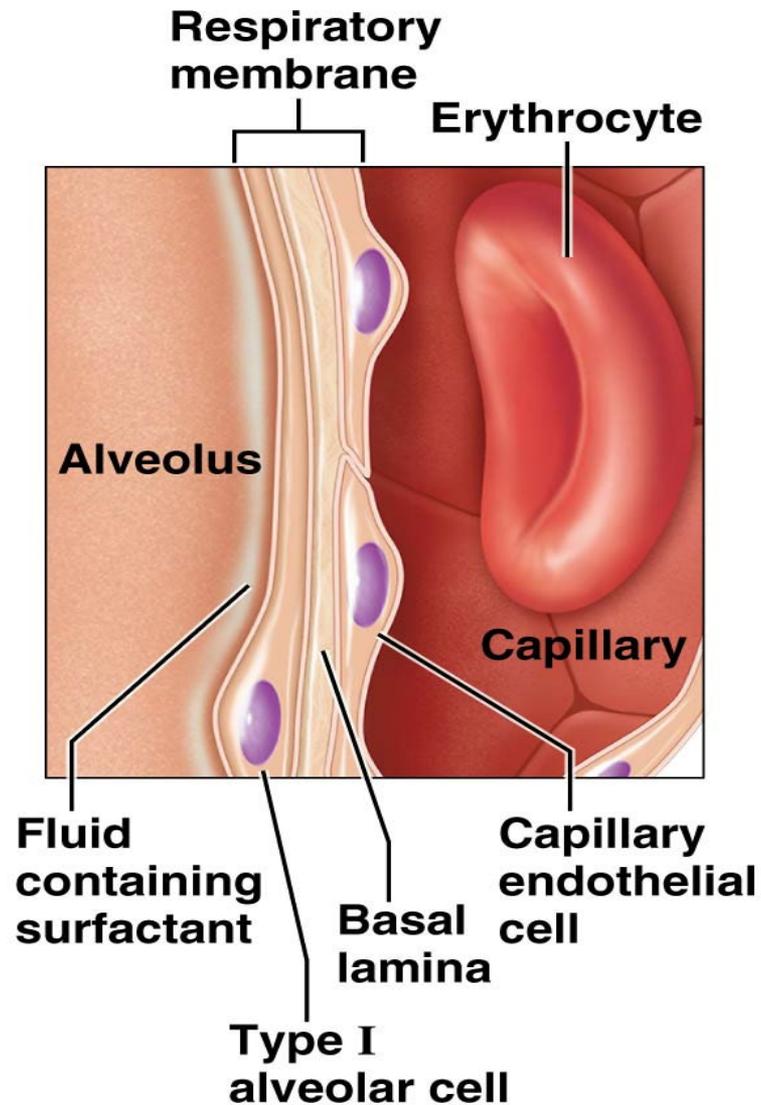
- squamous alveolar cells
- shared basement membrane
- endothelial cells of blood capillary

The shared basement membrane under certain conditions may swell due to edema.

Anything that increases distance across the respiratory membrane makes the diffusion of gasses slower

Anything that reduces the overall surface area of the respiratory membrane makes the diffusion of gasses slower

Structures of the respiratory membrane.



(b) Respiratory membrane

How Will a Layer of Water on the Inner Surface of Alveoli Influences Inspiration and Expiration?

Pulmonary surfactant produced by the great alveolar cells

Decreases surface tension by disrupting the hydrogen bonding in water

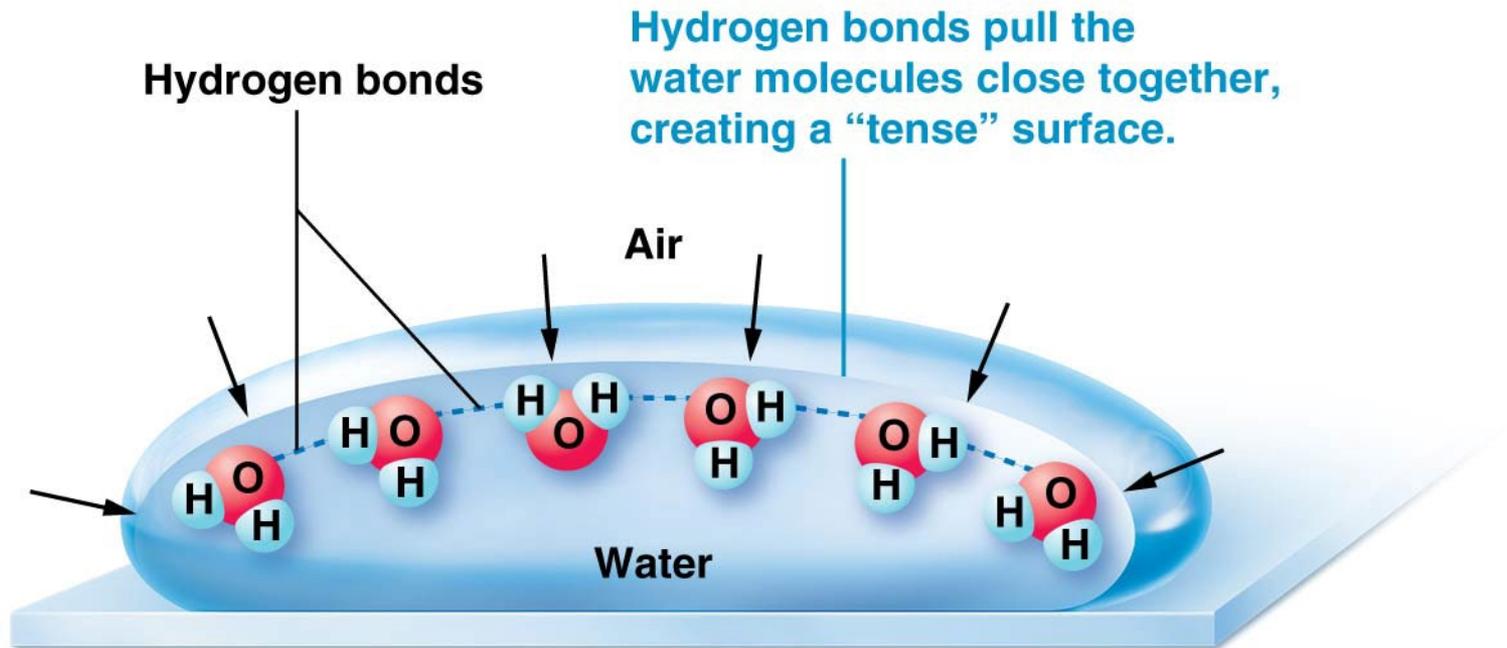
As lungs collapse, the water molecules on inner face of alveoli stack on top of each other.

These water molecules are connected by Hydrogen Bonds. /// surfactant disrupts Hydrogen Bonds, so it is easier to inflate lungs!!!

Because premature infants that lack surfactant they suffer from **infant respiratory distress syndrome (IRDS)** // great difficulty in breathing and may die

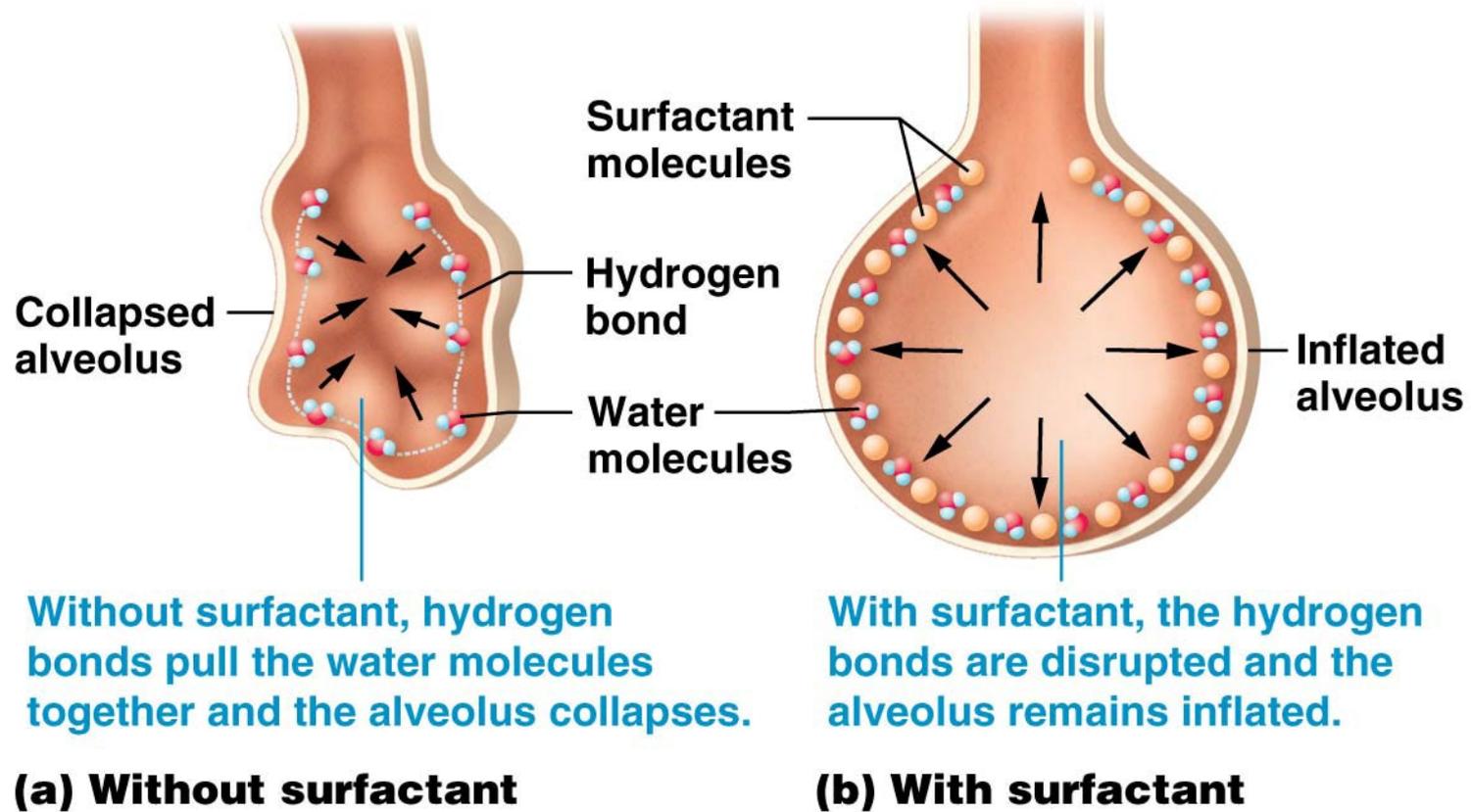
Knowledge about hydrogen bonding and the function of surfactant allows for treatment with artificial surfactant until lungs can produce its own surfactant

Water is a polar covalent molecule.



- It takes energy to separate water molecules
- This bonding helps to “deflate” the lungs.
- To reinflate the lungs these bonds must be broken.
- Pulmonary surfactant disrupts hydrogen bonds making it easier to inflate lungs.

Effect of surfactant on alveolar surface tension.



Without surfactant it will take more energy to inflate alveoli.

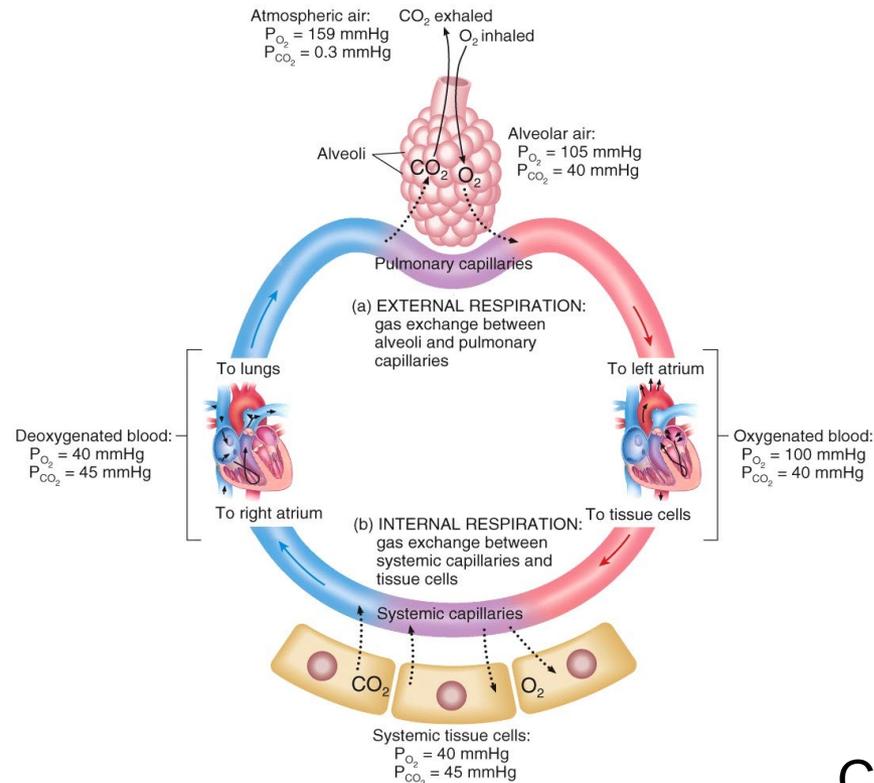
TABLE 22.4

**Composition of Inspired
(Atmospheric) and Alveolar Air**

Gas	Inspired Air*		Alveolar Air	
N ₂	78.6%	597 mm Hg	74.9%	569 mm Hg
O ₂	20.9%	159 mm Hg	13.7%	104 mm Hg
H ₂ O	0.5%	3.7 mm Hg	6.2%	47 mm Hg
CO ₂	0.04%	0.3 mm Hg	5.3%	40 mm Hg
Total	100%	760 mm Hg	100%	760 mm Hg

*Typical values for a cool clear day; values vary with temperature and humidity. Other gases present in small amounts are disregarded.

How is the concentration of O₂ and CO₂ different between the atmosphere and systemic tissues?

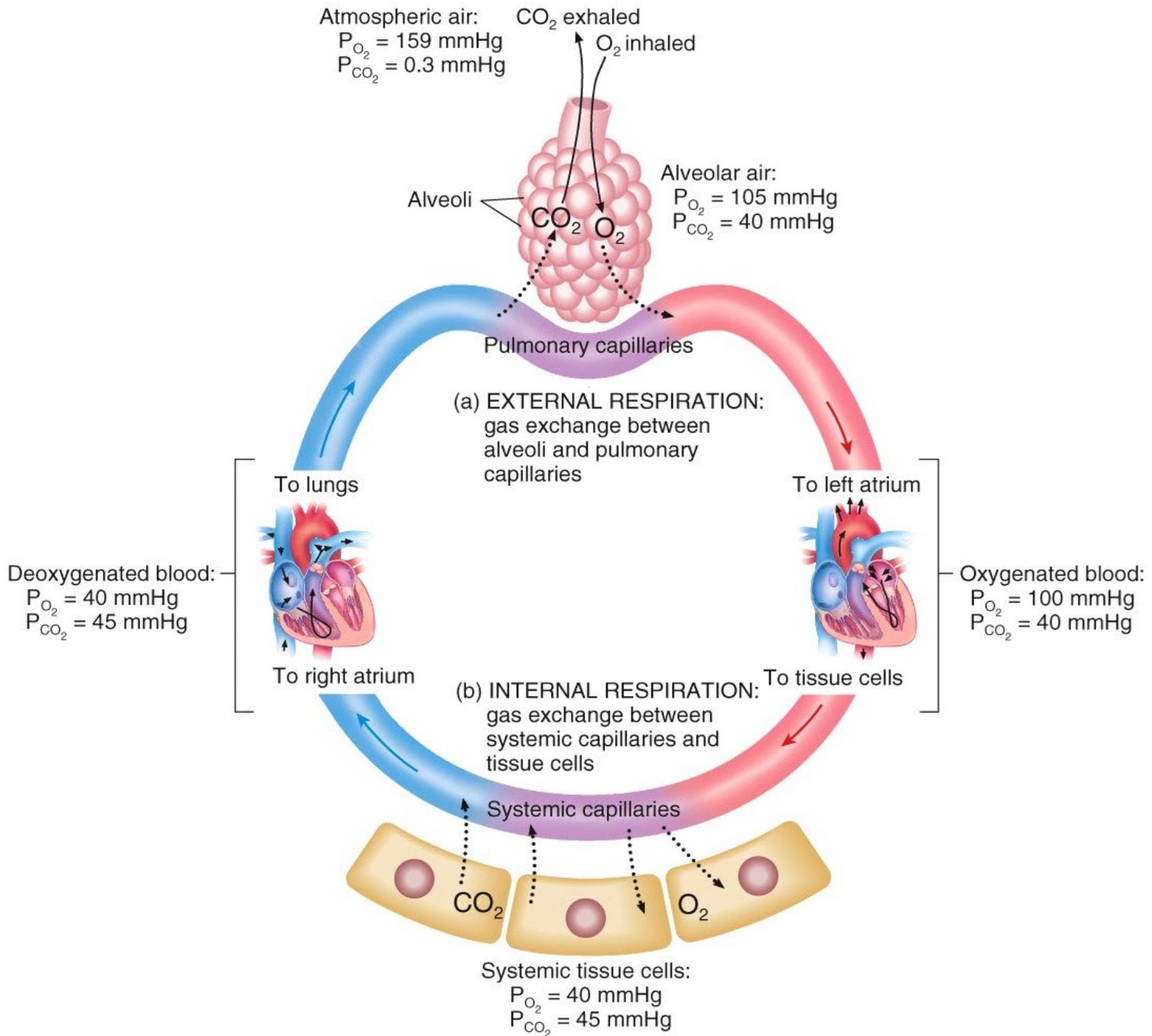


Oxygen

159 mmHg - atmosphere
 105 mmHg - alveolar
 100 mmHg - blood arterial
 40 mmHg - systemic tissue
 40 mmHg - blood venous

Carbon Dioxide

.3 mmHg - atmosphere
 40 mmHg - alveolar
 40 mmHg - blood arterial
 45 mmHg - systemic tissue
 45 mmHg - blood venous



Pressure Gradients Determine Airflow

Respiratory airflow is governed by the same principles that regulates the flow of blood: **pressure and resistance**

Fluid movement is directly proportional to the pressure difference between two points

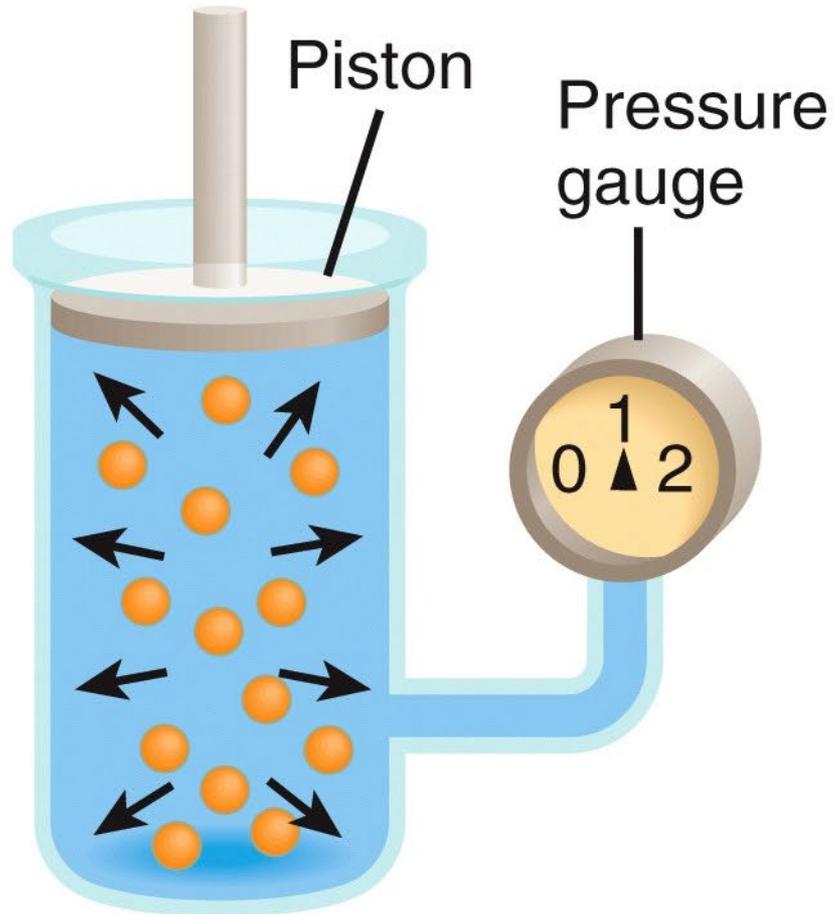
And fluid flow is inversely proportional to the resistance

So, if we want to ventilate the lungs (move air in and out) then we need to create a **transient pressure gradient** between the atmosphere and the lung tissue

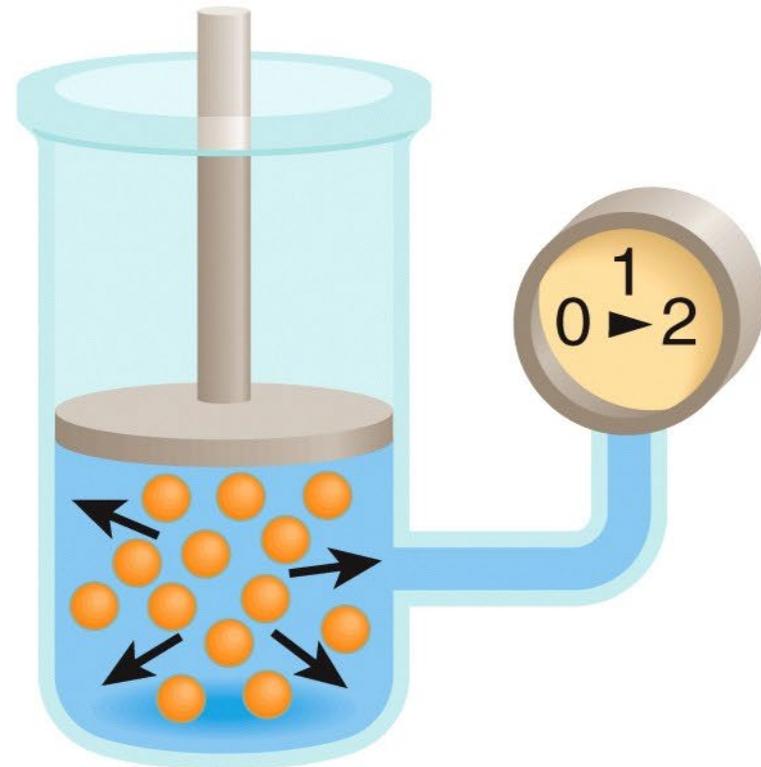
*Lower pressure in pleural cavity than the atmospheric pressure **creates inspiration**) /// greater pressure in pleural cavity than atmospheric pressure **creates expiration***

Laws of physic explain how the lungs move air!

Boyle's Law

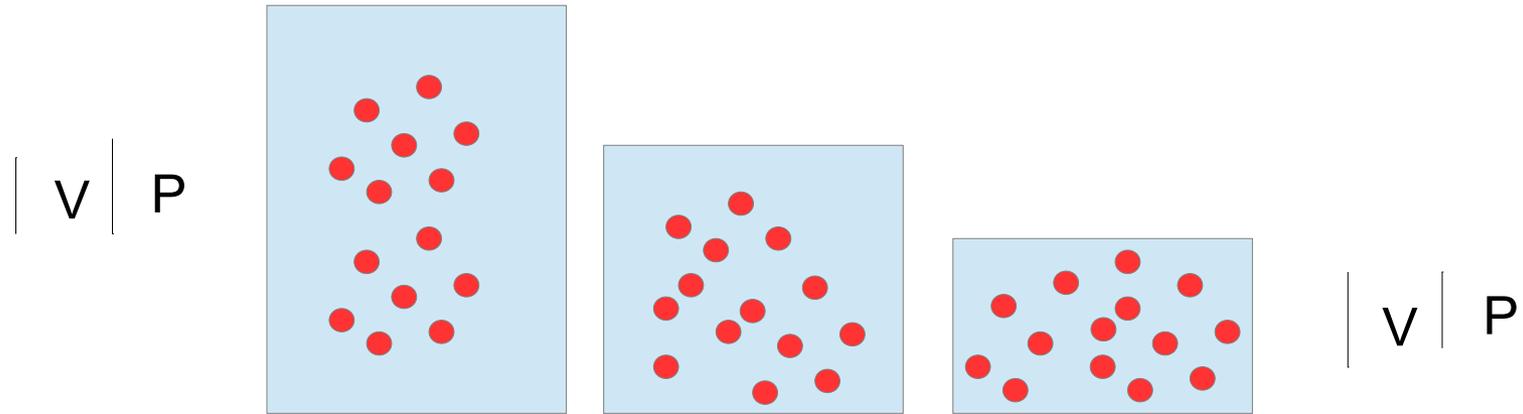


Volume = 1 liter
Pressure = 1 atm



Volume = 1/2 liter
Pressure = 2 atm

Boyle's Law



Assume constant temperature

Same number of gas atoms in each container

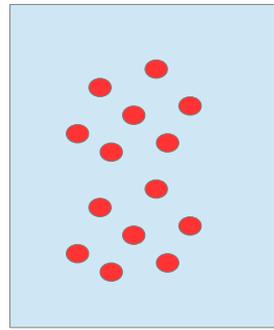
Fewer collisions in larger container (lower pressure)

More collisions in smaller container (higher pressure)

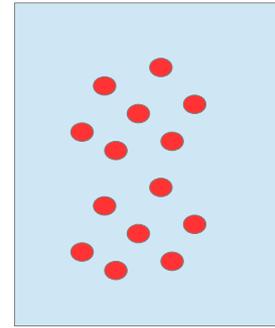
Pressure is inversely proportional to volume.

How does this help respiration? Significance?

Charles' Law



25 degree
Celcius



37 degree
Celcius

Higher Pressure

Here the **volume is constant but the temperature increases.**

Since gas molecules move faster at a higher temperature, there are more gas molecule collisions against the sides of the container which results in an increase in pressure. We did not increase the number of molecules, just the number of collisions.

When we breath in air, it is warmed by the mucosa.

How does this help respiration? Significance?

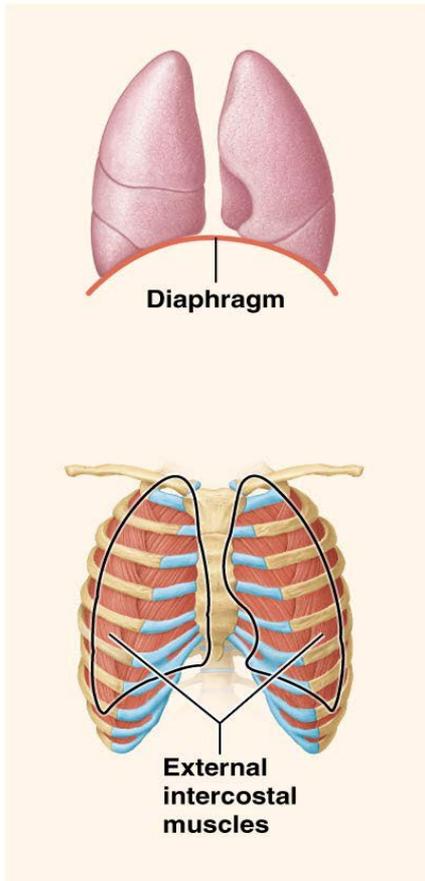
Note: Dalton's Law of Gases and Henry's

Law also help to explain respiratory physiology

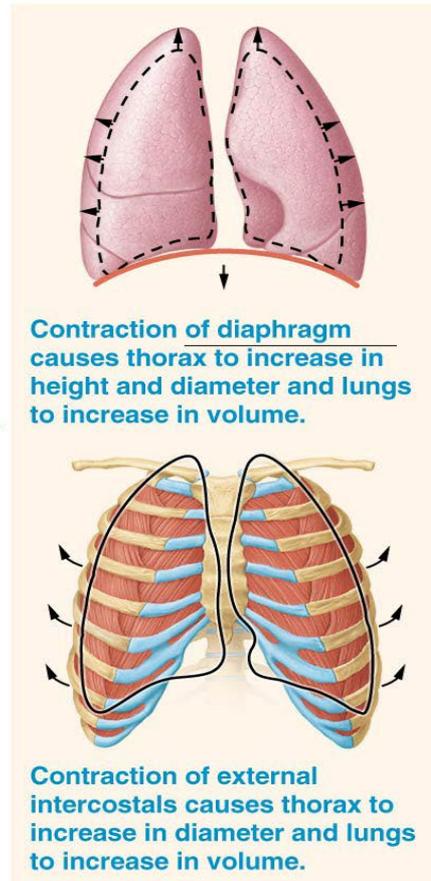
(see reference video for more details)

Volume changes in pulmonary ventilation: structure and function of the inspiratory muscles in quiet breathing.

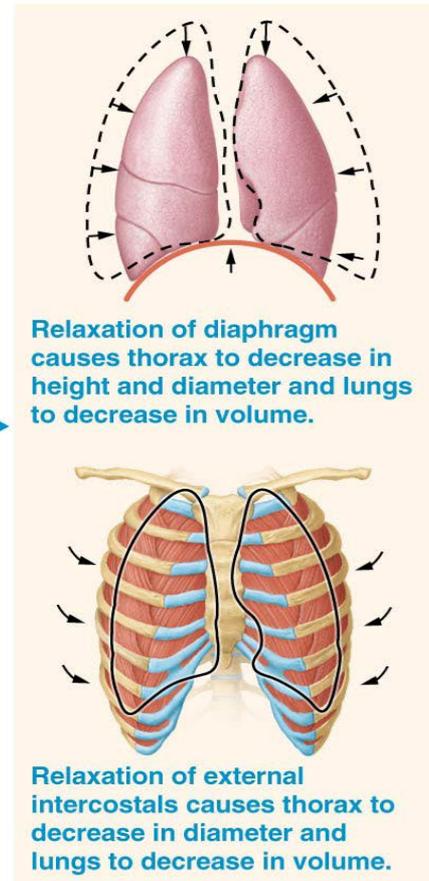
① Between breaths



② Inspiration: Diaphragm and external intercostals contract.



③ Expiration: Diaphragm and external intercostals relax.



- In “quiet breathing” **inspiration is active and expiration is passive.**
- Inspiration requires contraction of diaphragm and expiration occurs because of the recoil of elastic tissue expanded during inspiration.

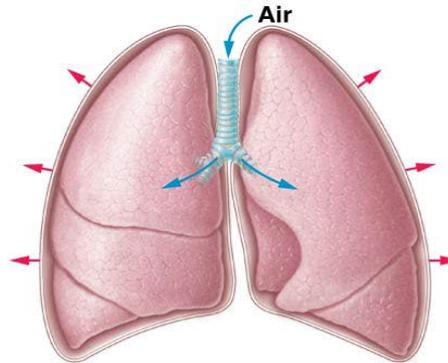
Pressure changes in pulmonary ventilation during respiratory cycle.

① **Between breaths:** Intrapulmonary pressure equals atmospheric pressure.



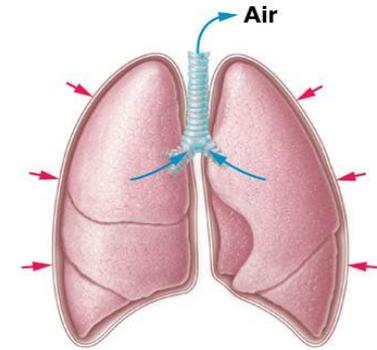
LOCATION	PRESSURE (mm Hg)
Atmospheric (air)	760
Intrapulmonary (alveoli)	760
Intrapleural (pleural cavity)	756

② **Inspiration:** Lung volume increases, intrapulmonary pressure decreases, and air rushes into the lungs.



LOCATION	PRESSURE (mm Hg)
Atmospheric (air)	760
Intrapulmonary (alveoli)	↓ 758
Intrapleural (pleural cavity)	↓ 754

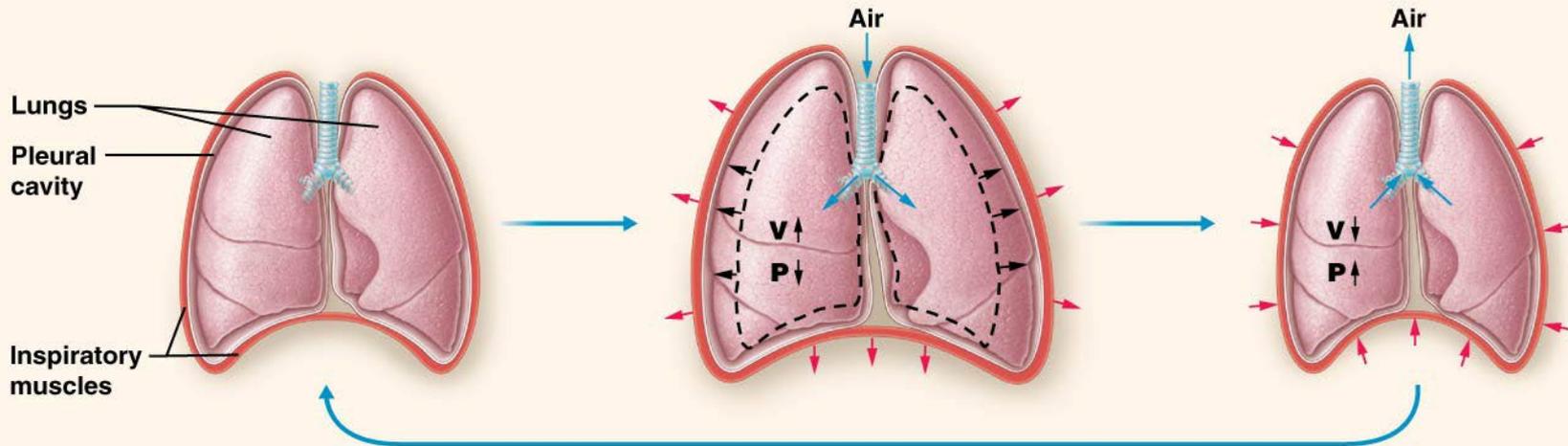
③ **Expiration:** Intrapulmonary pressure increases, lung volume decreases, and air moves out of the lungs.



LOCATION	PRESSURE (mm Hg)
Atmospheric (air)	760
Intrapulmonary (alveoli)	↑ 762
Intrapleural (pleural cavity)	↑ 758

- Respiratory cycle is **five seconds** long.
- Inspiration occurs for **two seconds** as diaphragm contracts (active / volume).
- Expiration occurs for **three seconds** as diaphragm relaxes by elastic recoil.

The Big Picture of Pulmonary Ventilation



Between breaths

Intrapulmonary pressure equals atmospheric pressure.

No air movement occurs.

Inspiration

Inspiratory muscles contract.

Thoracic volume increases.

Lung volume increases.

Intrapulmonary pressure decreases to below atmospheric pressure.

Air flows into lungs.

Expiration

Inspiratory muscles relax.

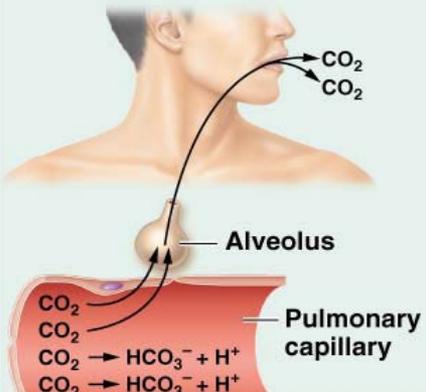
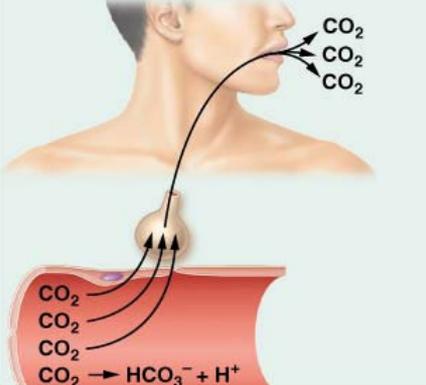
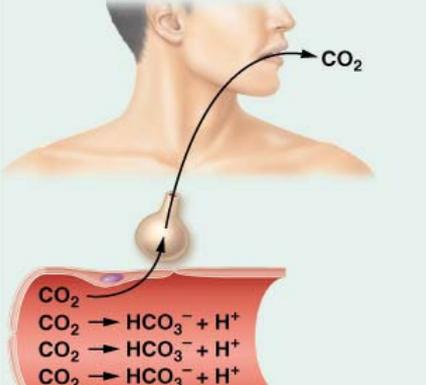
Thoracic volume decreases due to elastic recoil.

Lung volume decreases.

Intrapulmonary pressure increases to above atmospheric pressure.

Air flows out of lungs.

Passive

PATTERN OF VENTILATION	BLOOD pH
<p>Normal ventilation</p>  <p>CO₂ CO₂</p> <p>Alveolus</p> <p>CO₂ CO₂ CO₂ → HCO₃⁻ + H⁺ CO₂ → HCO₃⁻ + H⁺</p> <p>Pulmonary capillary</p>	<p>Remains stable pH = 7.35–7.45</p>
<p>Hyperventilation</p>  <p>CO₂ CO₂ CO₂</p> <p>CO₂ CO₂ CO₂ CO₂ → HCO₃⁻ + H⁺</p>	<p>Increases, pH > 7.45</p> <p>More CO₂ is expired.</p> <p>↓</p> <p>Blood P_{CO₂} decreases.</p> <p>↓</p> <p>Carbonic acid concentration decreases.</p> <p>↓</p> <p>Hydrogen ion concentration decreases.</p> <p>↓</p> <p>Blood pH increases.</p>
<p>Hypoventilation</p>  <p>CO₂</p> <p>CO₂ CO₂ CO₂ → HCO₃⁻ + H⁺ CO₂ → HCO₃⁻ + H⁺ CO₂ → HCO₃⁻ + H⁺</p>	<p>Decreases, pH < 7.35</p> <p>Less CO₂ is expired.</p> <p>↓</p> <p>Blood P_{CO₂} increases.</p> <p>↓</p> <p>Carbonic acid concentration increases.</p> <p>↓</p> <p>Hydrogen ion concentration increases.</p> <p>↓</p> <p>Blood pH decreases.</p>

Effect of blood pH on ventilation patterns



Carbon dioxide plus water makes carbonic acid that then changes into proton plus bicarbonate

Carbonic acid is an acid because it can donate a proton

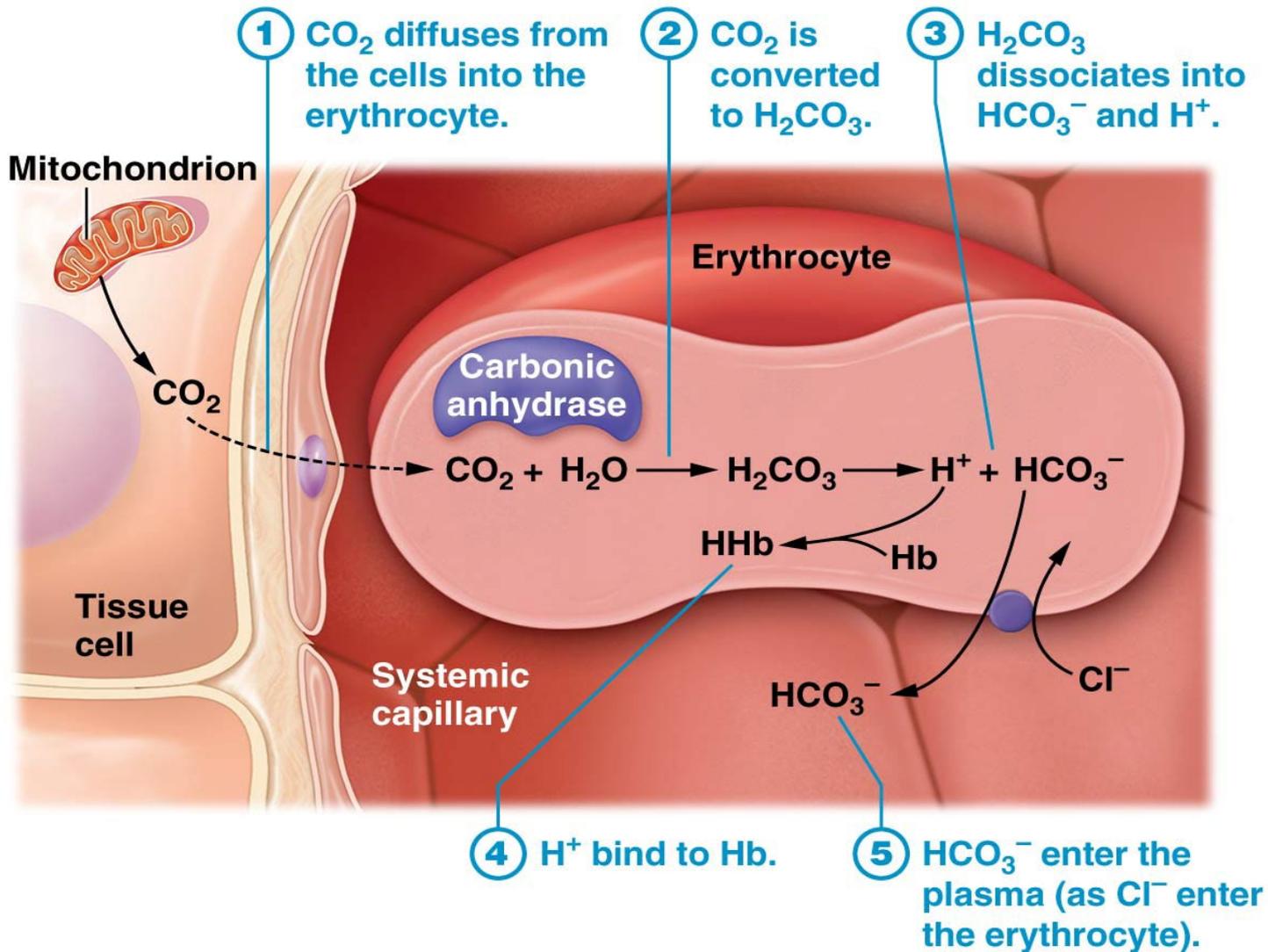
Bicarbonate is a weak base, and the proton is an acid

This chemical reaction occurs inside RBC and is catalyzed by carbonic anhydrase (CA)

Hyperventilate reduces ventilation

Hypoventilation stimulates ventilation

Transport of carbon dioxide: the conversion of carbon dioxide and water into carbonic acid in erythrocytes.



Bicarbonate forms ionic bond with sodium and is transported to lungs as sodium-bicarbonate

Hydrogen ion binds to Hb-O₂ and causes oxygen to be released from Hb // oxygen now free to diffuse to mitochondria

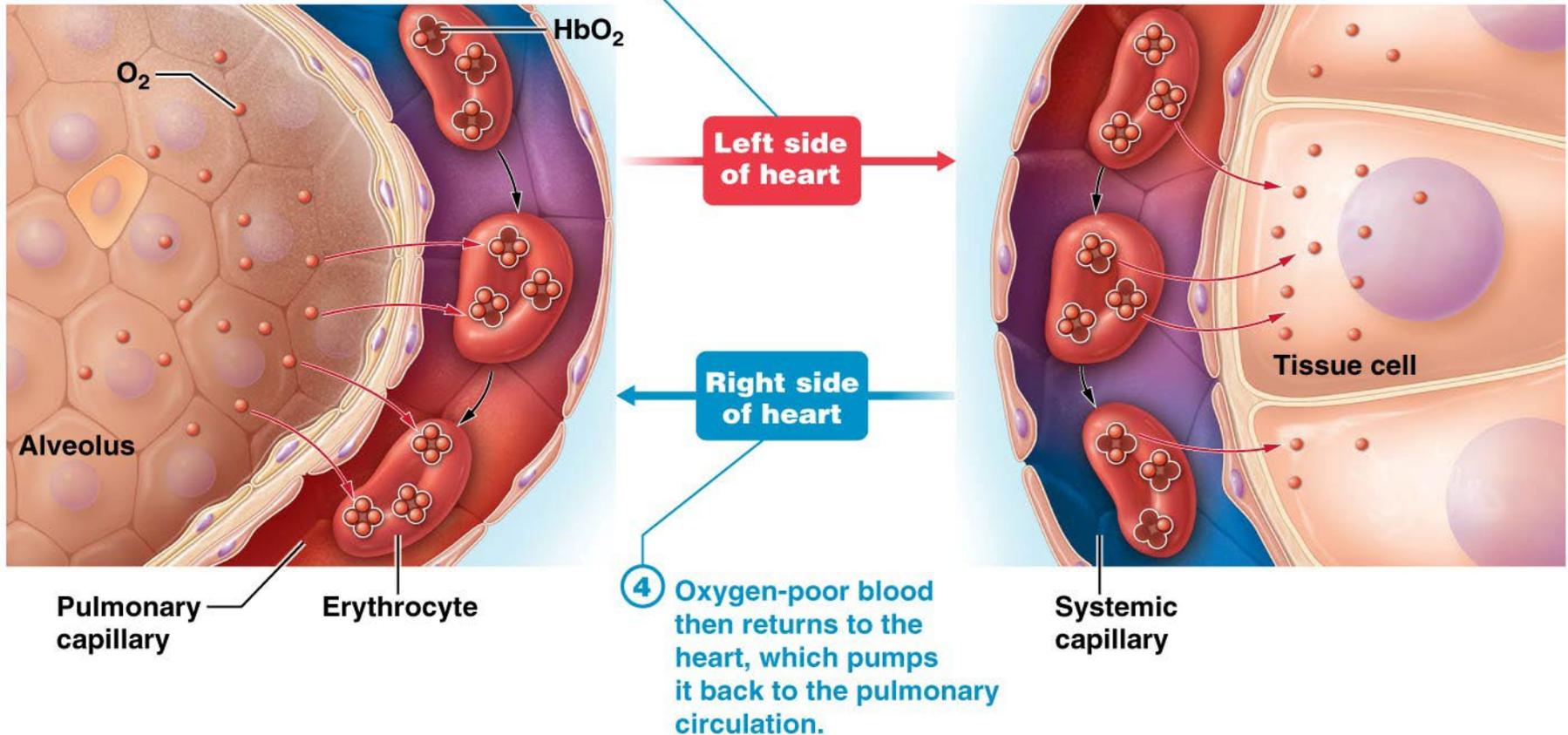
(a) Bicarbonate formation in an erythrocyte in a systemic capillary

Transport of oxygen: loading and unloading of oxygen on hemoglobin in erythrocytes.

① During loading, oxygen from alveoli binds to hemoglobin (Hb) in the pulmonary capillaries, converting it to oxyhemoglobin (HbO₂).

② Oxygen-rich blood travels to the heart, which pumps it to the systemic circulation.

③ During unloading, Hb in the systemic capillaries releases oxygen to the tissue cells.



pH

More basic
(alkaline)



Neutral



More acidic

pH

14

13

12

11

10

9

8

7

6

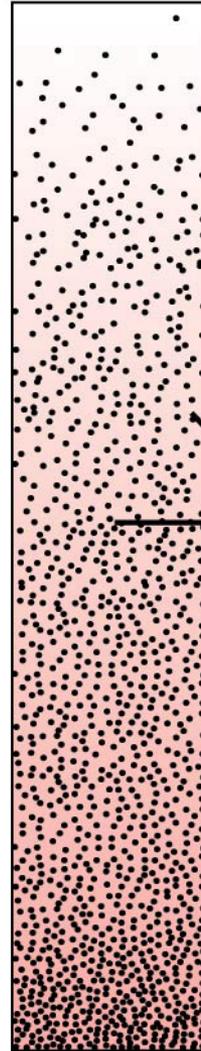
5

4

3

2

1



Hydrogen
ions

Primary Control of Ventilation

The primary control centers for breathing (ventilation of lungs) are the **chemoreceptors** that respond to carbon dioxide levels in the blood. // Two locations

Central chemoreceptors // Located in the medulla

Peripheral chemoreceptors // Located in the carotid bodies and aortic arch

Both chemoreceptors detect changes in carbon dioxide level & hydrogen ions /// increase CO₂ stimulates breathing

Hypercapnia (high PCO₂) Main Stimulus for Ventilation

As carbon dioxide levels in the blood increases // pH lower – blood becomes more acid like.

Carbon dioxide easily diffuses into cerebral spinal fluid within medulla oblongata (respiratory control centers).

This lowers pH and stimulates respiratory center

Increased rate and depth of respiration

Hypercapnia (high PCO₂) Main Stimulus for Ventilation

Hyperventilation is a condition in which you start to breathe very fast. Healthy breathing occurs with a healthy balance between breathing in oxygen and breathing out carbon dioxide.

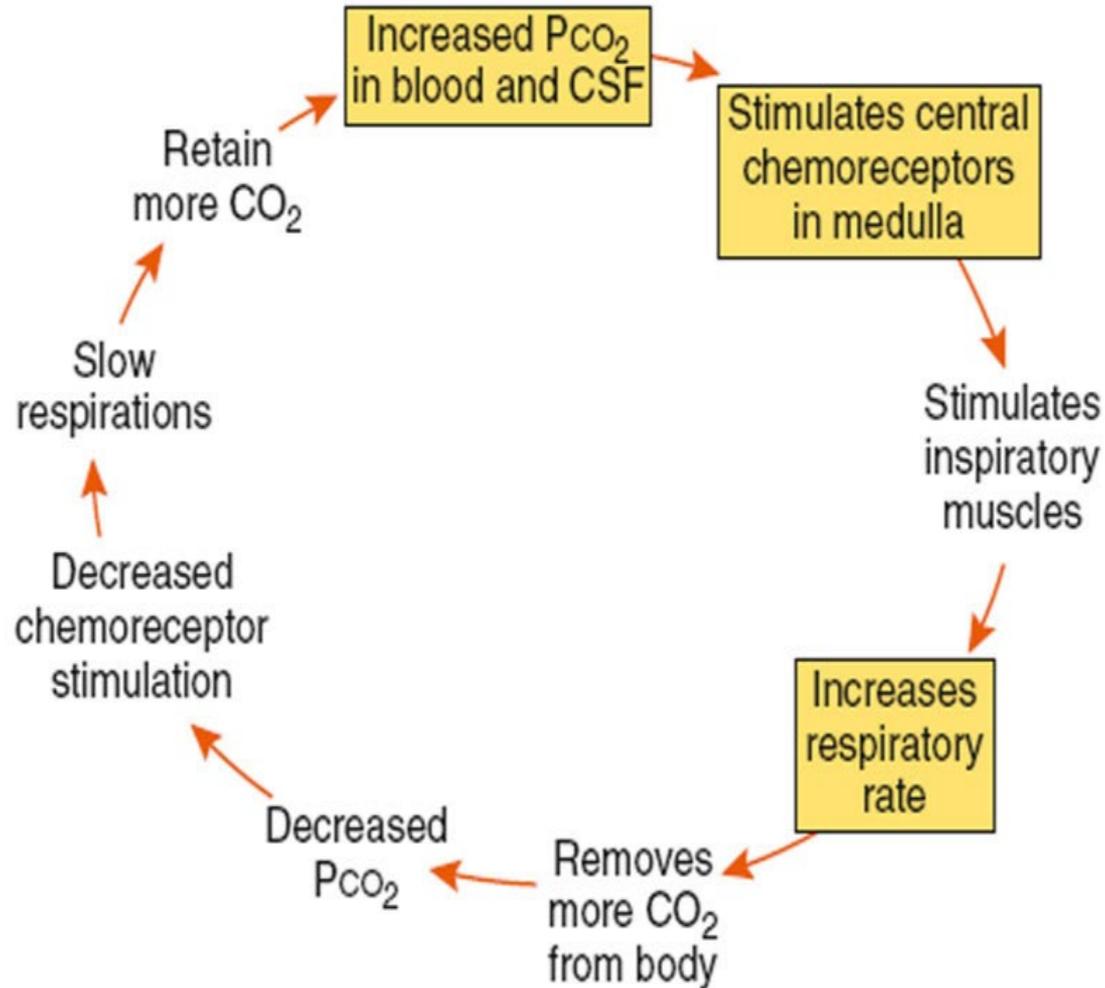
You upset this balance when you hyperventilate by exhaling more CO₂. This upsets acid-base balance in blood.

Hyperventilation causes **respiratory alkalosis**, and this will depress central nervous system.

Swimmers often hyperventilate thinking this will add more oxygen to their blood so they can hold their breath longer, however. This is not true. The hyperventilation reduces carbon dioxide blood concentration so the swimmer may faint when they dive into the water. Very dangerous practice!

Normal Respiratory Control

A. NORMAL CYCLE



Hydrogen Ions & Pulmonary Ventilation

Pulmonary ventilation is adjusted to maintain the pH within brain

Central chemoreceptors in the medulla oblongata produce about **75% of the change in respiration induced by pH shift**

Yet H^+ does not cross the blood-brain barrier very easily

CO_2 crosses blood brain barrier rapidly and in CSF reacts with water and produces carbonic acid

Dissociates into bicarbonate and hydrogen ions /// most H^+ remains free and greatly stimulates the central chemoreceptors

Hydrogen ions also stimulate **peripheral chemoreceptors** which produce about **25% of the respiratory response to pH change**

HOWEVER - peripheral chemoreceptors also sensitive to PO_2 // play important role in respiration after chronic high PCO_2 or very low oxygen concentrations

When Peripheral Chemoreceptors Control of Ventilation

The peripheral chemoreceptors are stimulated by low oxygen levels (but not the central chemoreceptors)

Occurs if PO₂ falls below normal 100 mmHg and above 50 mmHg // within this range PO₂ stimulates breathing rate

Below 50 mmHg peripheral chemoreceptors do not stimulate inspiration

Severe PO₂ deficiency will depress central chemoreceptors // DRG do not respond to any inputs and sends fewer impulses to inhalation muscles // breathing slows – as O₂ continues to drop this starts a fatal positive feedback loop!

O2 Secondary Control of Ventilation

Hypoxemia // Marked decrease in oxygen

Primary stimulus is under normal physiologic conditions with elevated CO₂ however.....

Extended period of high CO₂ will cause chemoreceptors to fatigue, and they will not respond elevated CO₂

Peripheral chemoreceptors now may react to low O₂ levels

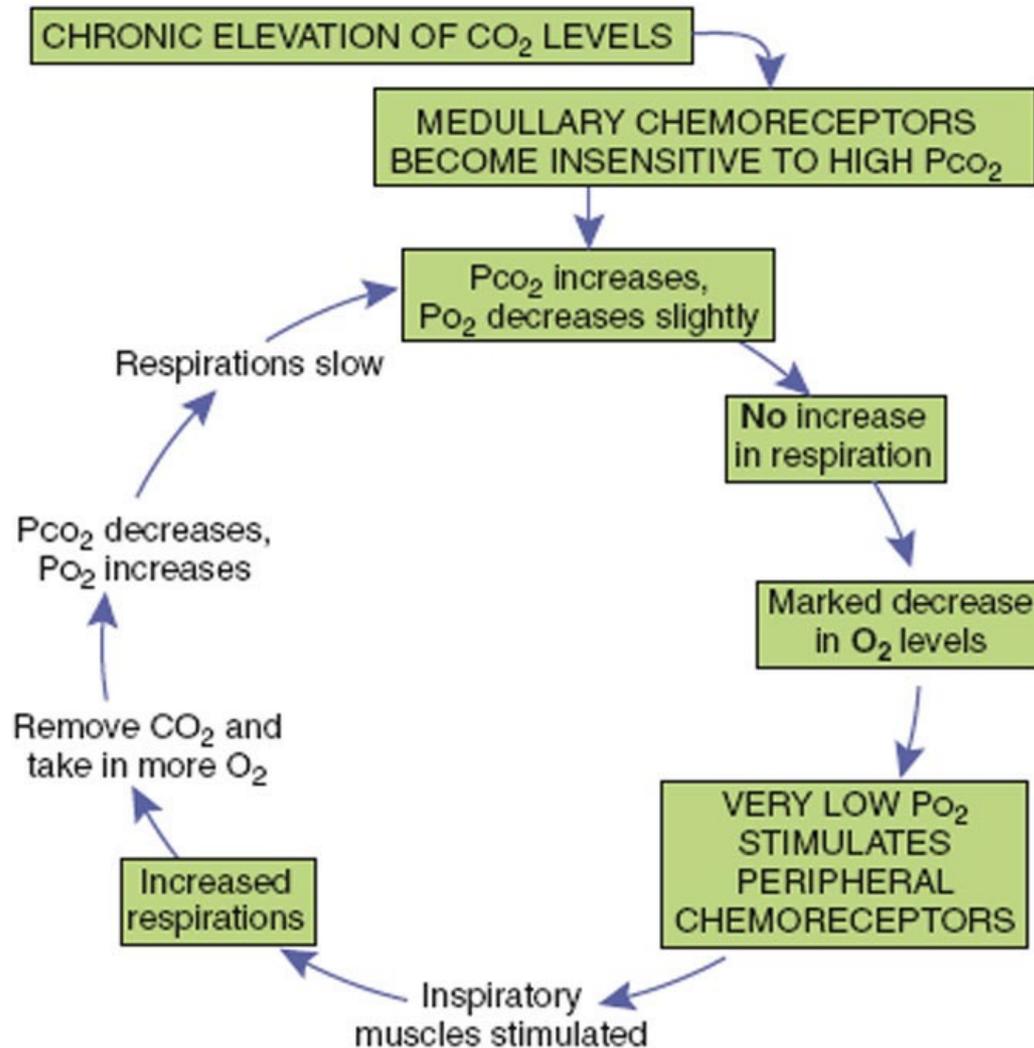
This is why O₂ becomes most important control mechanism in individuals with chronic lung disease (e.g. emphysema)

This changes respiratory system into **hypoxic drive**

Central control centers no longer respond to CO₂ stimulus, and now peripheral chemoreceptors dominates with PO₂ as “controller”

Hypoxic Drive

B. HYPOXIC DRIVE WITH CHRONIC ELEVATED P_{CO_2} LEVELS (e.g., emphysema)



Air Flow Into Lung Tissue and Gas Exchange

Only air that reaches the alveoli is available for gas exchange

Not all inhaled air gets to alveoli

500 mL moved into respiratory system // tidal volume

About 150 mL of the tidal volume fills the conducting division of the airway

Only 350 mL of air reaches the alveoli

Usable Air Within Lung Tissue

Anatomic dead space /// conducting division of airway where there is no gas exchange

This can be altered slightly by sympathetic and parasympathetic stimulation

Pulmonary diseases

Respiratory membrane thickened by edema or fibrosis

Some alveoli may be unable to exchange gases because they lack blood flow (perfusion and ventilation are not matched)

Some diseases reduce amount of respiratory membrane

These events all add to anatomic dead space to create total dead space

Usable Air Within Lung Tissue

Physiologic dead space (also called total dead space)

Sum of **anatomic dead space** and any **pathological alveolar dead space** caused by disease

If a person **inhales 500 mL of air** /// If 150 mL stays in anatomical dead space /// **only 350 mL reaches alveoli**

Alveolar ventilation rate (AVR)

Air that ventilates alveoli (350 mL) X respiratory rate (12 bpm) = 4200 mL/min

Of all the measurements, this one is most directly relevant to the body's ability to get oxygen to the tissues and dispose of carbon dioxide /// **lung diseases reduces AVR**

Measuring Ventilation Volumes

Spirometer – a device that recaptures expired breath and records such variables such as rate and depth of breathing, speed of expiration, and rate of oxygen consumption ([view video](#))

Respiratory volumes ([view video](#))

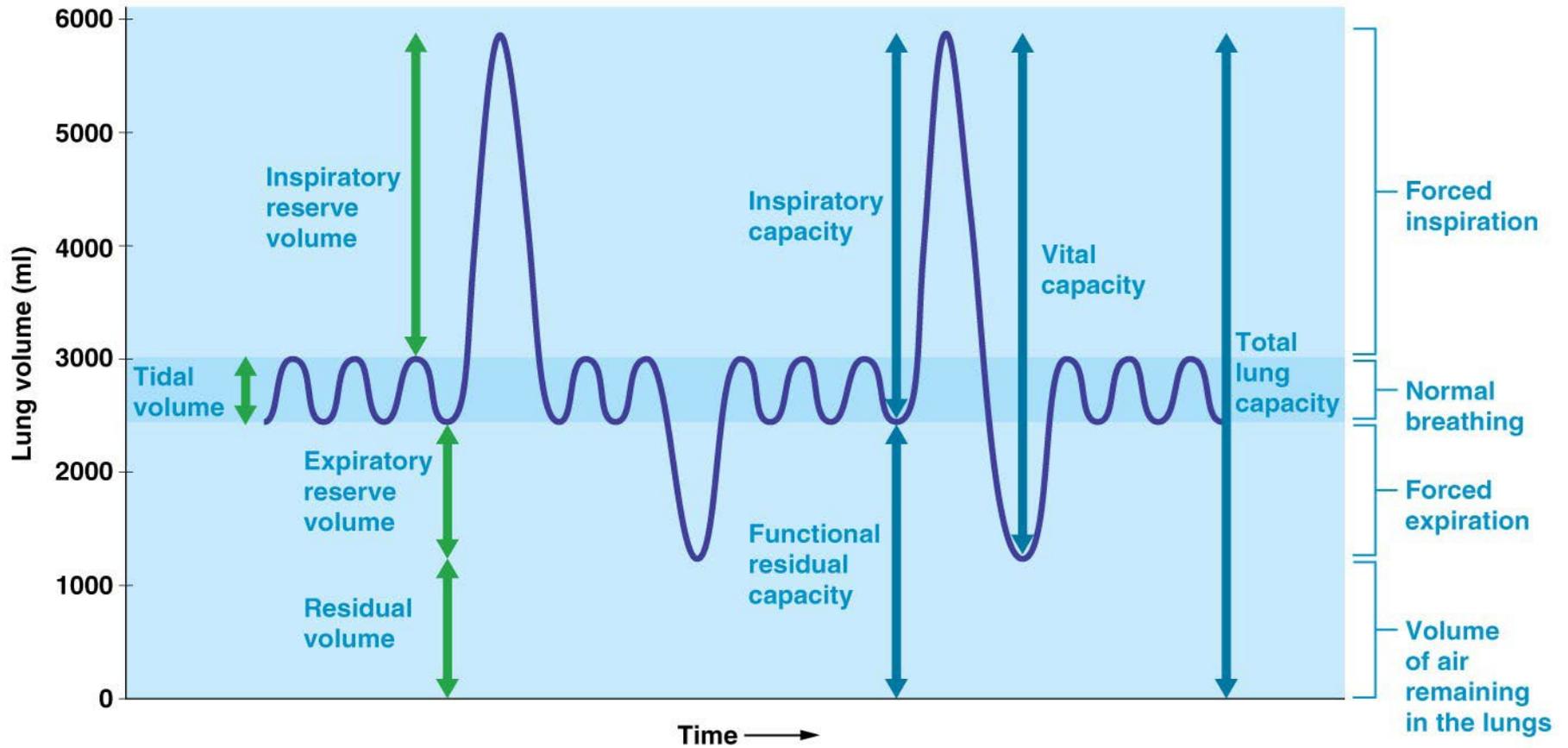
Tidal volume - volume of air inhaled and exhaled in one cycle during quiet breathing (**500 mL**)

Inspiratory reserve volume - air in excess of tidal volume that can be inhaled with maximum effort (3000 mL)

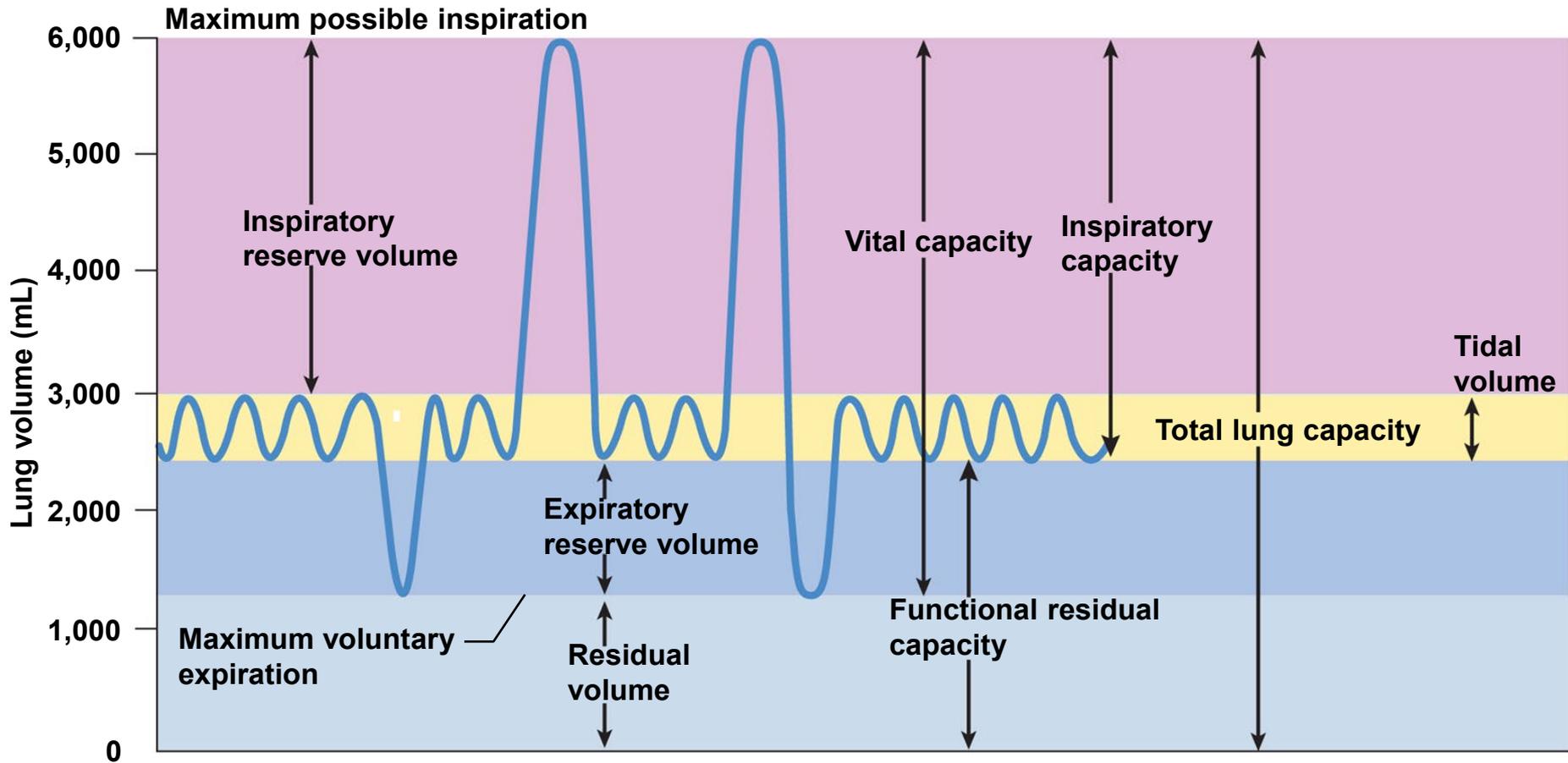
Expiratory reserve volume - air in excess of tidal volume that can be exhaled with maximum effort (1200 mL)

Residual volume - air remaining in lungs after maximum expiration (1300 mL)

Graph of pulmonary volumes and capacities.



Lung Volumes and Capacities



Factors Affecting Gas Exchange

membrane thickness / membrane surface area / ventilation-perfusion coupling

Respiratory Membrane thickness

Only 0.5 μm thick

Presents little obstacle to diffusion

Pulmonary edema caused by left ventricular failure results in “thickening” of the respiratory membrane

Pneumonia also causes thickening of respiratory membrane

Results = greater distance for gases to travel between blood and air

Cannot equilibrate fast enough to keep up with blood flow

Factors Affecting Gas Exchange

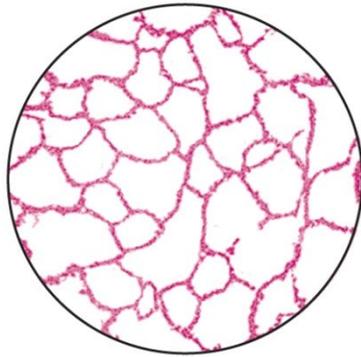
Membrane surface area

100 ml blood in alveolar capillaries, spread thinly over 70 m²

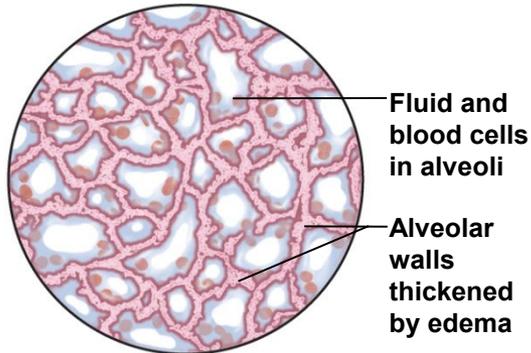
Emphysema, lung cancer, and tuberculosis all decrease surface area for gas exchange

These conditions results in hypoxia

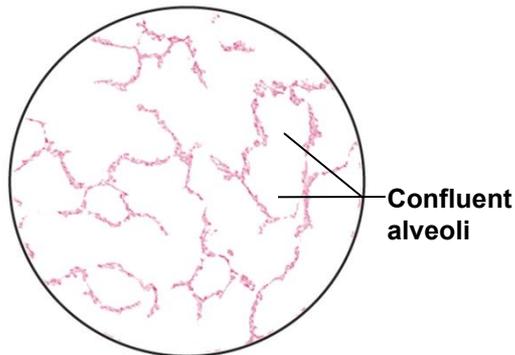
Lung Disease Affects Gas Exchange



(a) Normal



(b) Pneumonia



(c) Emphysema

Abnormally few but large alveoli with total reduction of respiratory membrane.

Autoregulation to Adjust Blood and Air Flows Within Lungs

Ventilation-perfusion coupling // the ability to match ventilation to perfusion (see next two slides)

Gas exchange requires both good ventilation of alveolus and good perfusion of the capillaries

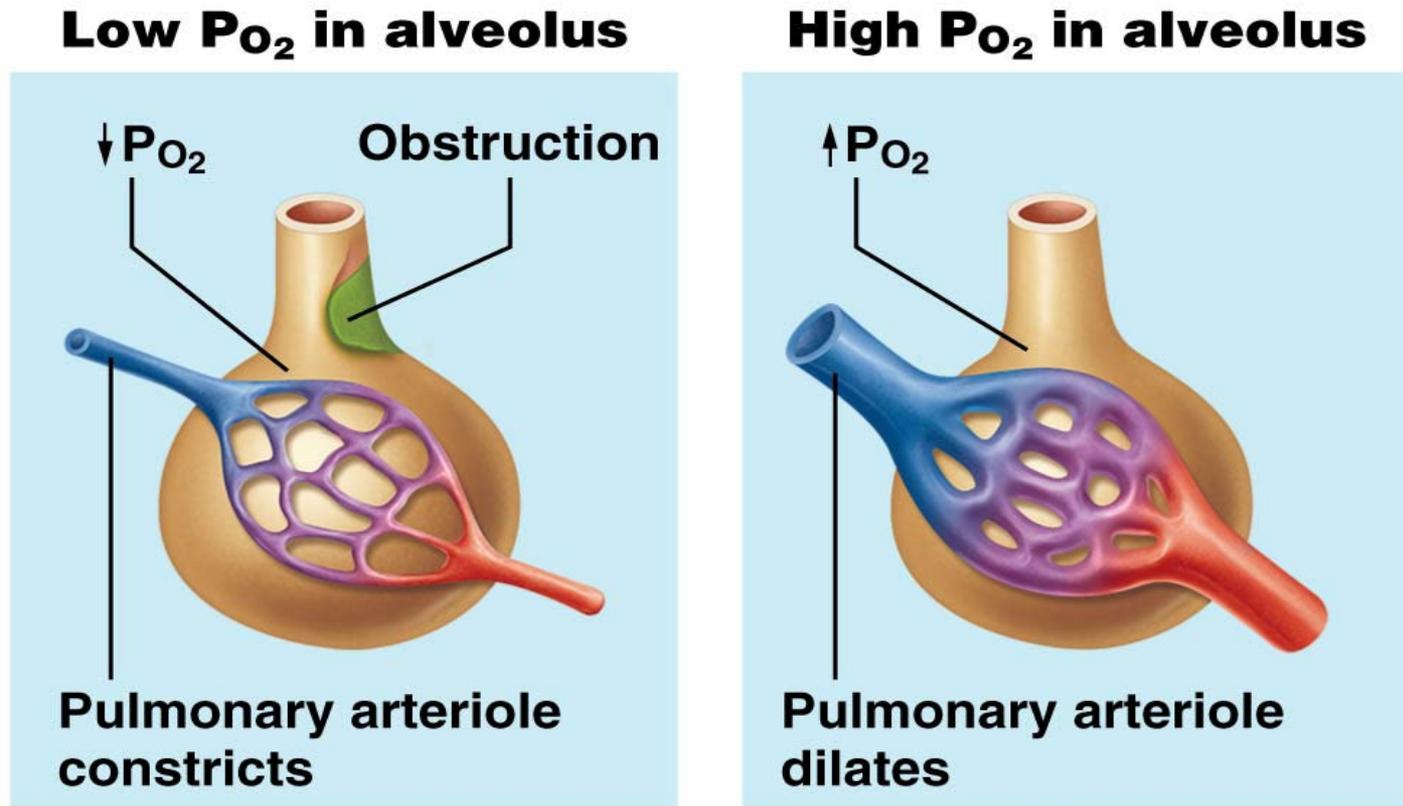
Ventilation-perfusion ratio of 0.8 (a flow of 4.2 L of air and 5.5 L of blood per minute at rest)

- Ventilation is air movement through alveoli
- Perfusion is blood flow through pulmonary capillary

Ventilation and perfusion is matched within the lung tissue by “autoregulation” // smooth muscles in bronchioles and pulmonary arterioles will selectively open and close to match ventilation to perfusion or perfusion to ventilation.

Ventilation – Perfusion Coupling

(Oxygen concentration in alveoli regulates smooth muscle in pulmonary arterioles.)



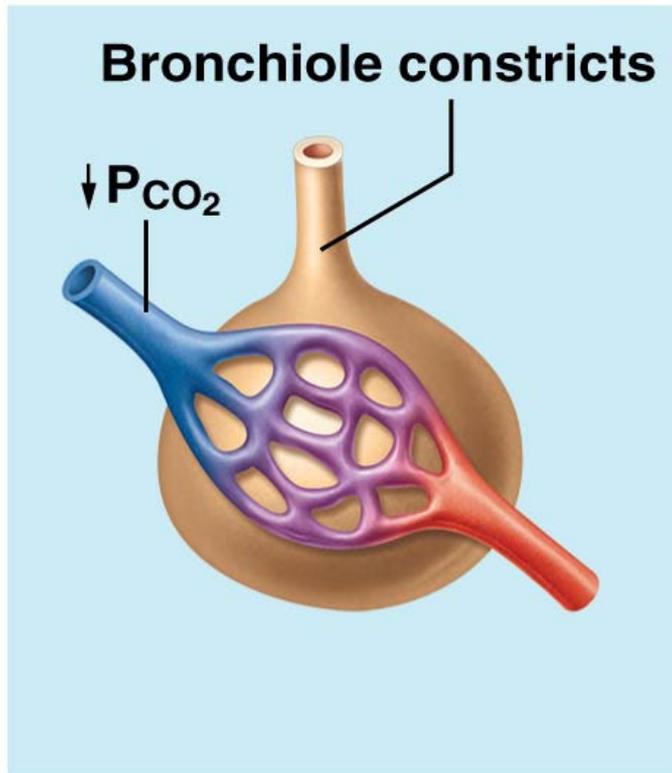
(a) Changes in ventilation (alveolar P_{O_2}) lead to changes in perfusion.

What happens when you change ventilation (bronchioles)?
You change smooth muscle tension in pulmonary arteriole.

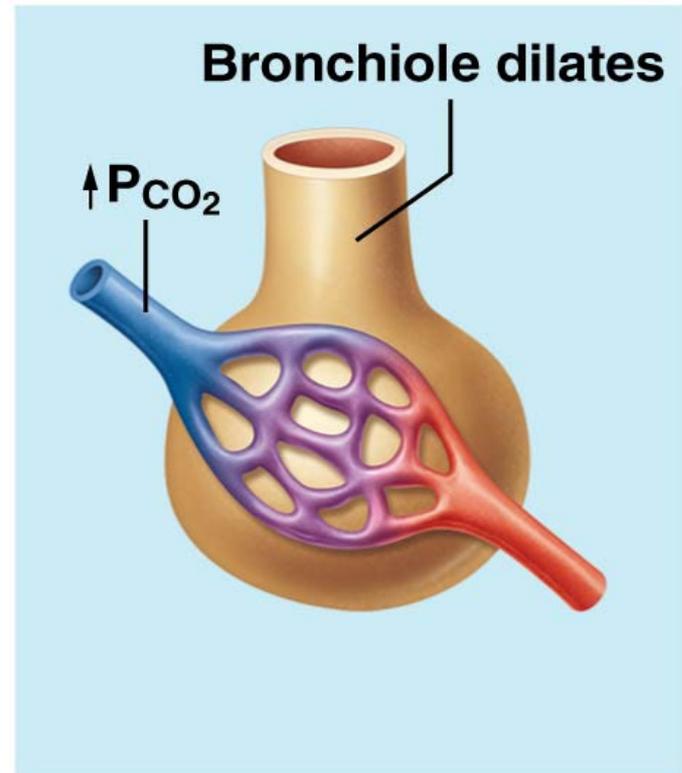
Perfusion - Ventilation Coupling

(Carbon dioxide concentration in pulmonary arteriole regulate smooth muscle in bronchioles)

Low P_{CO_2} in arteriole



High P_{CO_2} in arteriole



(b) Changes in perfusion (arteriolar P_{CO_2}) lead to changes in ventilation.

What happens when you change perfusion (pulmonary arterioles)?
You change smooth muscle tension in bronchioles.

Gas Transport

The process of carrying gases

- from the alveoli to the systemic tissues
- from the system tissues to the alveoli

Oxygen transport

- 98.5% bound to hemoglobin
- 1.5% dissolved in plasma

Carbon dioxide transport

- 70% as bicarbonate ion
- 23% bound to hemoglobin (carbamino)
- 7% dissolved in plasma

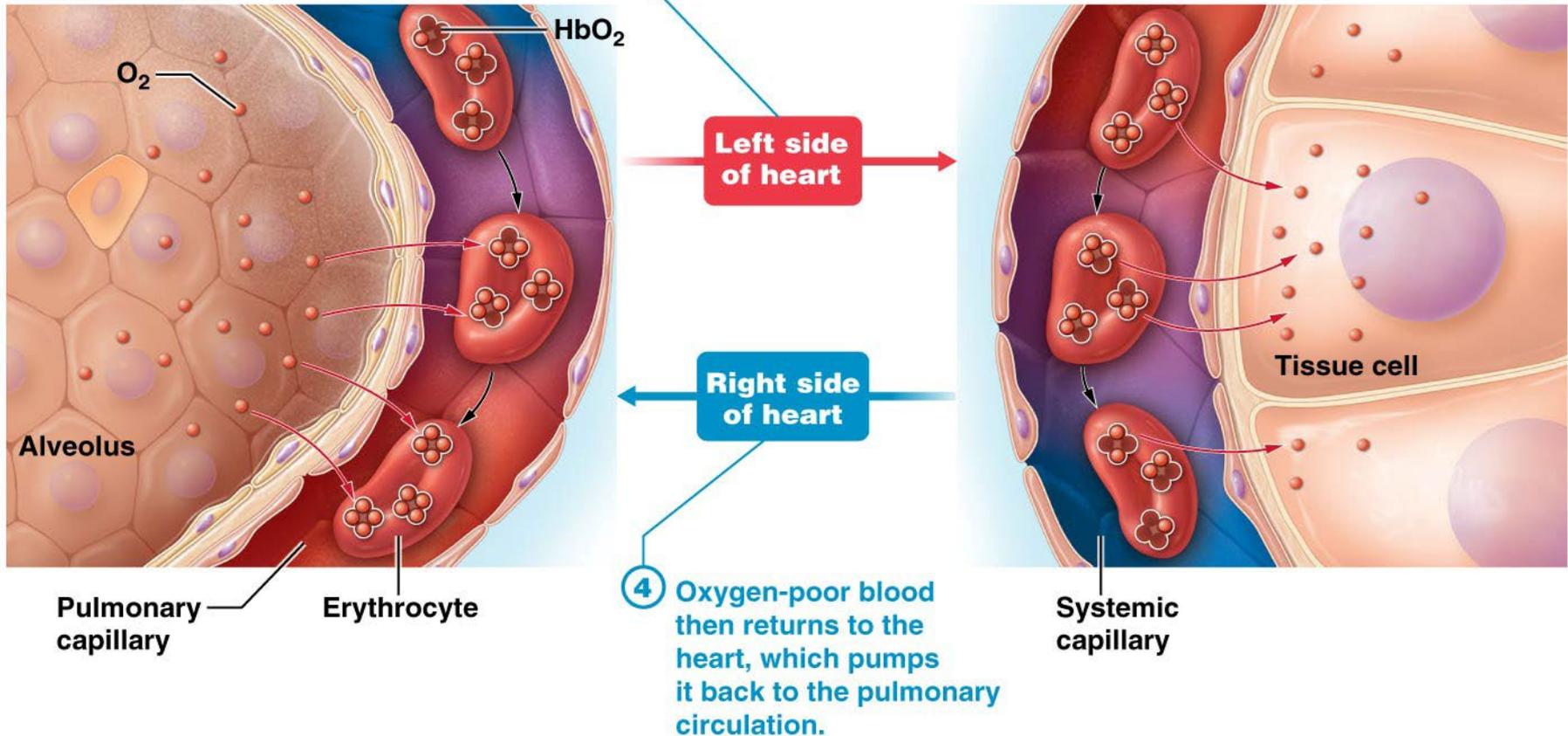
Transport of oxygen: loading and unloading of oxygen on hemoglobin in erythrocytes.



① During loading, oxygen from alveoli binds to hemoglobin (Hb) in the pulmonary capillaries, converting it to oxyhemoglobin (HbO₂).

② Oxygen-rich blood travels to the heart, which pumps it to the systemic circulation.

③ During unloading, Hb in the systemic capillaries releases oxygen to the tissue cells.



Oxygen Transport

Hemoglobin (Hb) – molecule specialized in oxygen transport

Hb consist of four separate protein molecules (globulin)

Each globulin molecule has a heme group which binds one O_2 to its ferrous ion (Fe^{2+})

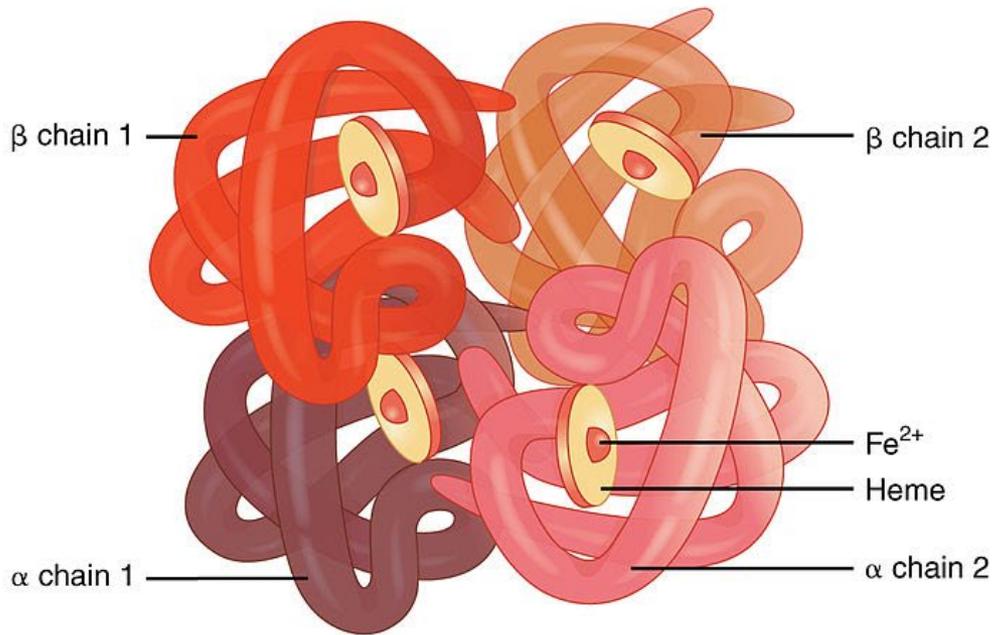
Therefore, one hemoglobin molecule can carry up to 4 O_2

100% oxyhemoglobin (HbO_2) – 4 O_2 bound to hemoglobin

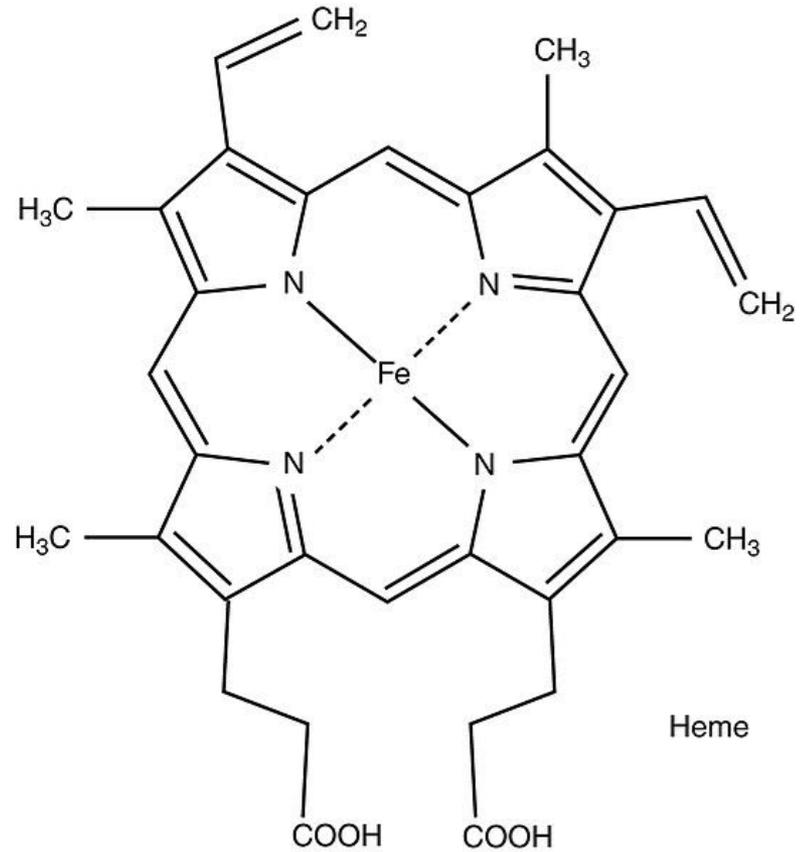
Deoxyhemoglobin (HHb) – hemoglobin with no O_2

50% saturation Hb with 2 oxygen molecules

Oxygen Transport



(a)



(b)

Carbon Dioxide Transport

Carbon dioxide transported in blood using these three forms

- carbonic acid
- carbamino compounds
- dissolved in plasma

How is oxygen and carbon dioxide load and unload during Systemic Gas Exchange?

Unloading of O₂ /// from blood into tissue

Loading CO₂ /// from tissue into blood

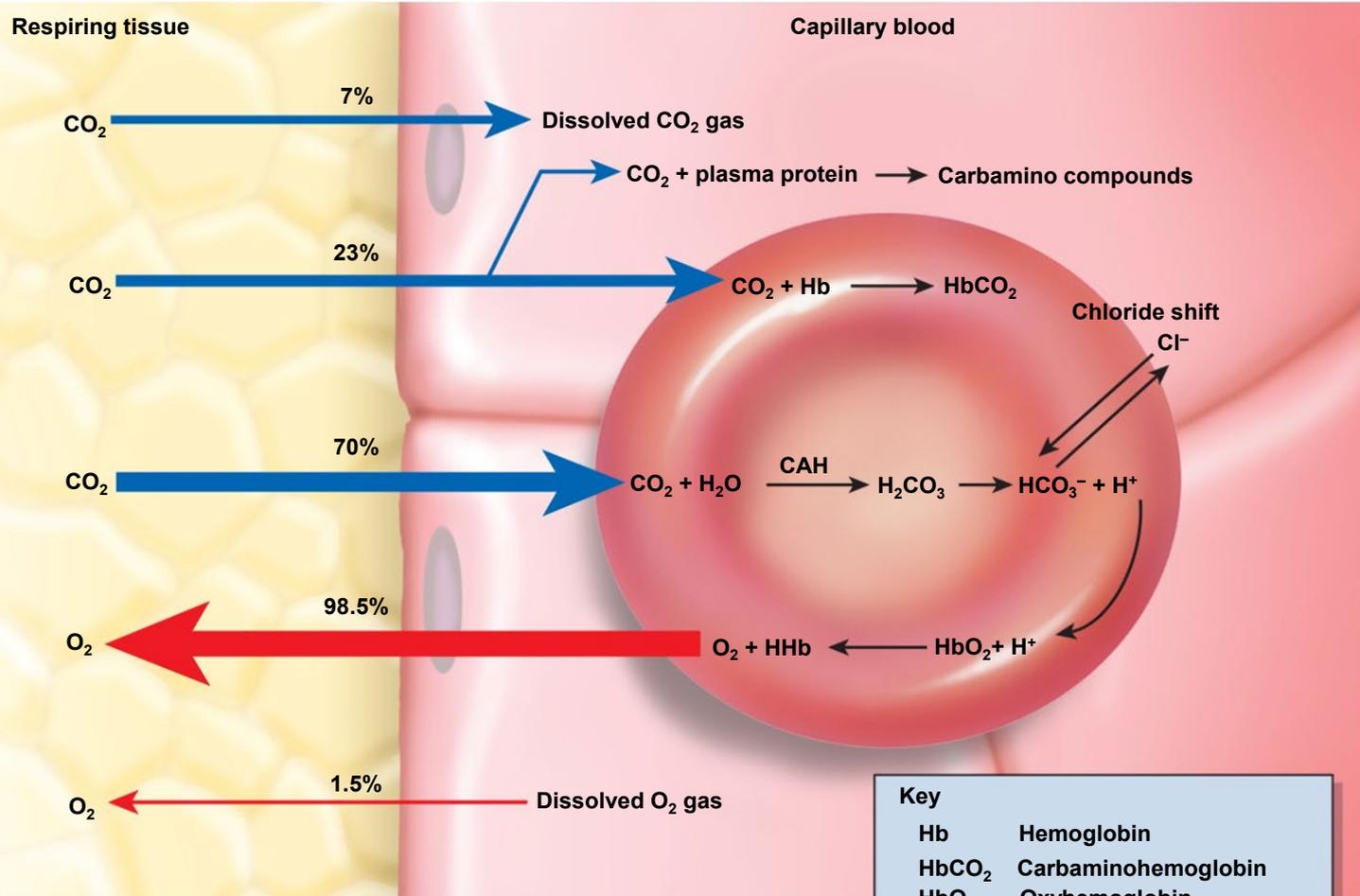
Key event occurs inside RBC // requires carbonic anhydrase



The “Chloride Shift”

- keeps reaction proceeding // exchanges HCO₃⁻ for Cl⁻
- H⁺ binds to hemoglobin

Systemic Gas Exchange



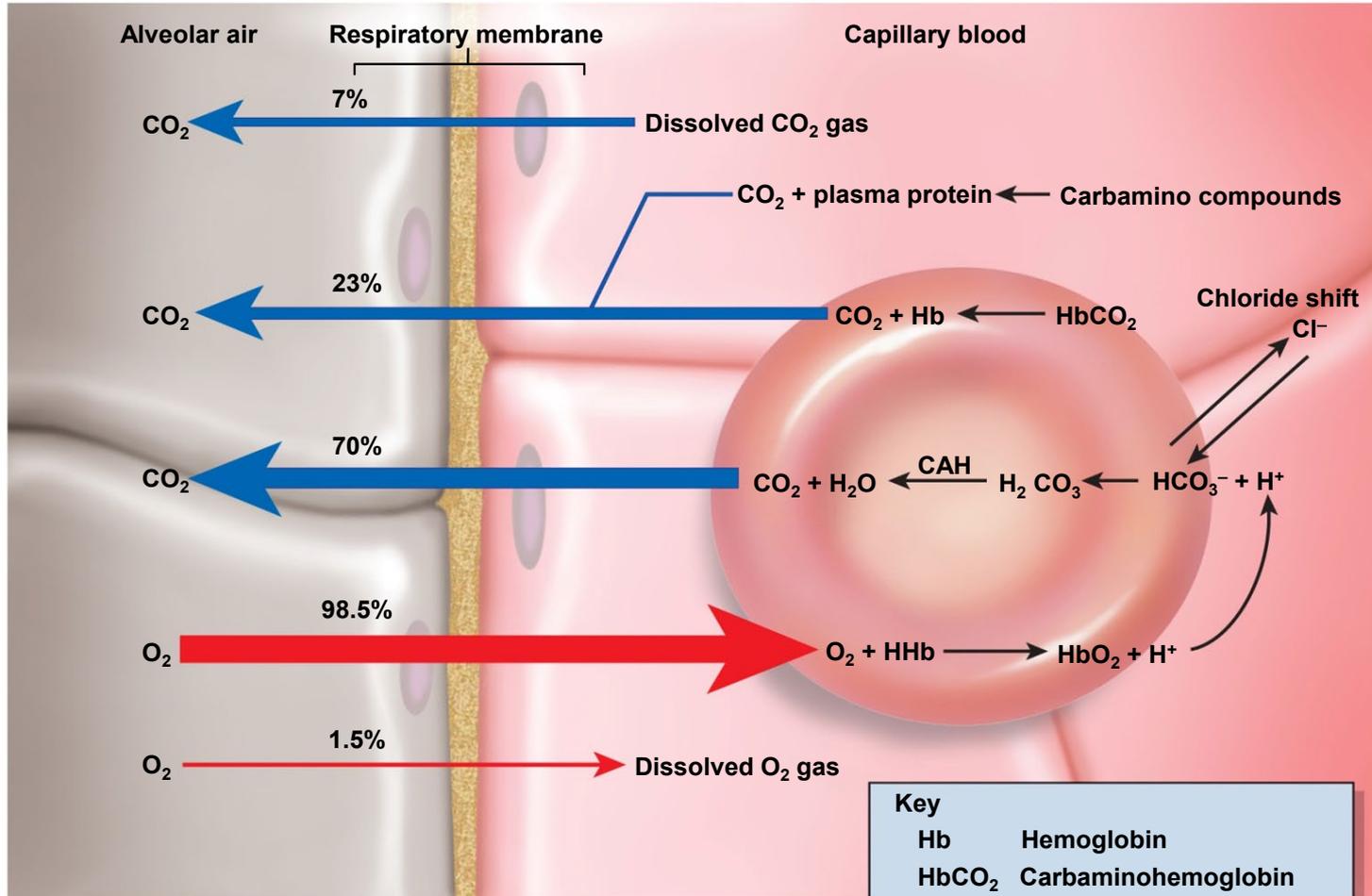
Carbonic anhydrase (CAH) makes H₂CO₂ // inside RBC

Hydrogen ion (H⁺) causes the release of O₂ from Hb

What moves into cell as bicarbonate moves out?

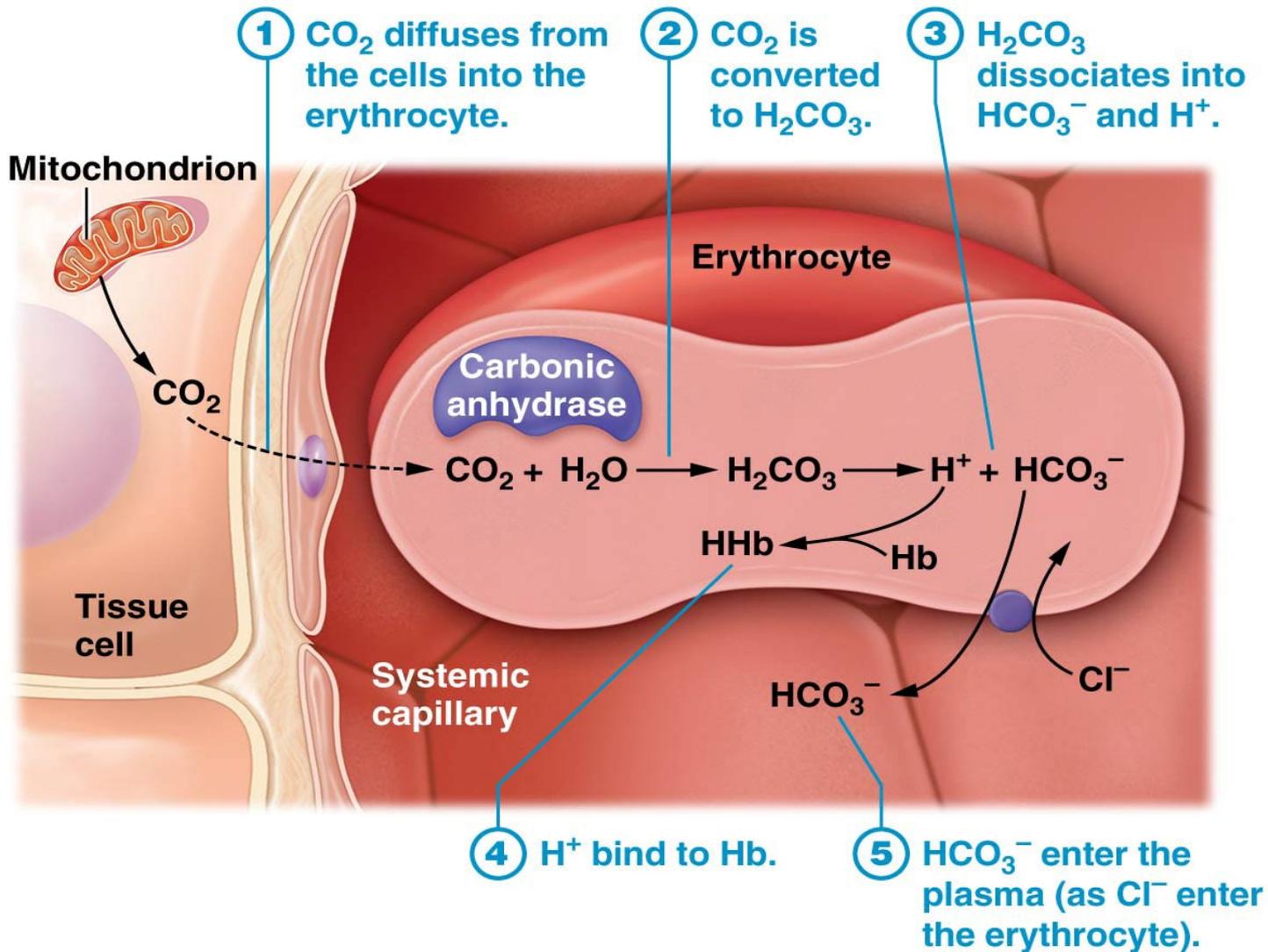
Alveolar Gas Exchange

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Everything moves in the opposite direction from the systemic gas exchange! Follow the diffusion of O₂ into RBC and see outcome.

Transport of carbon dioxide: the conversion of carbon dioxide and water into carbonic acid in erythrocytes.

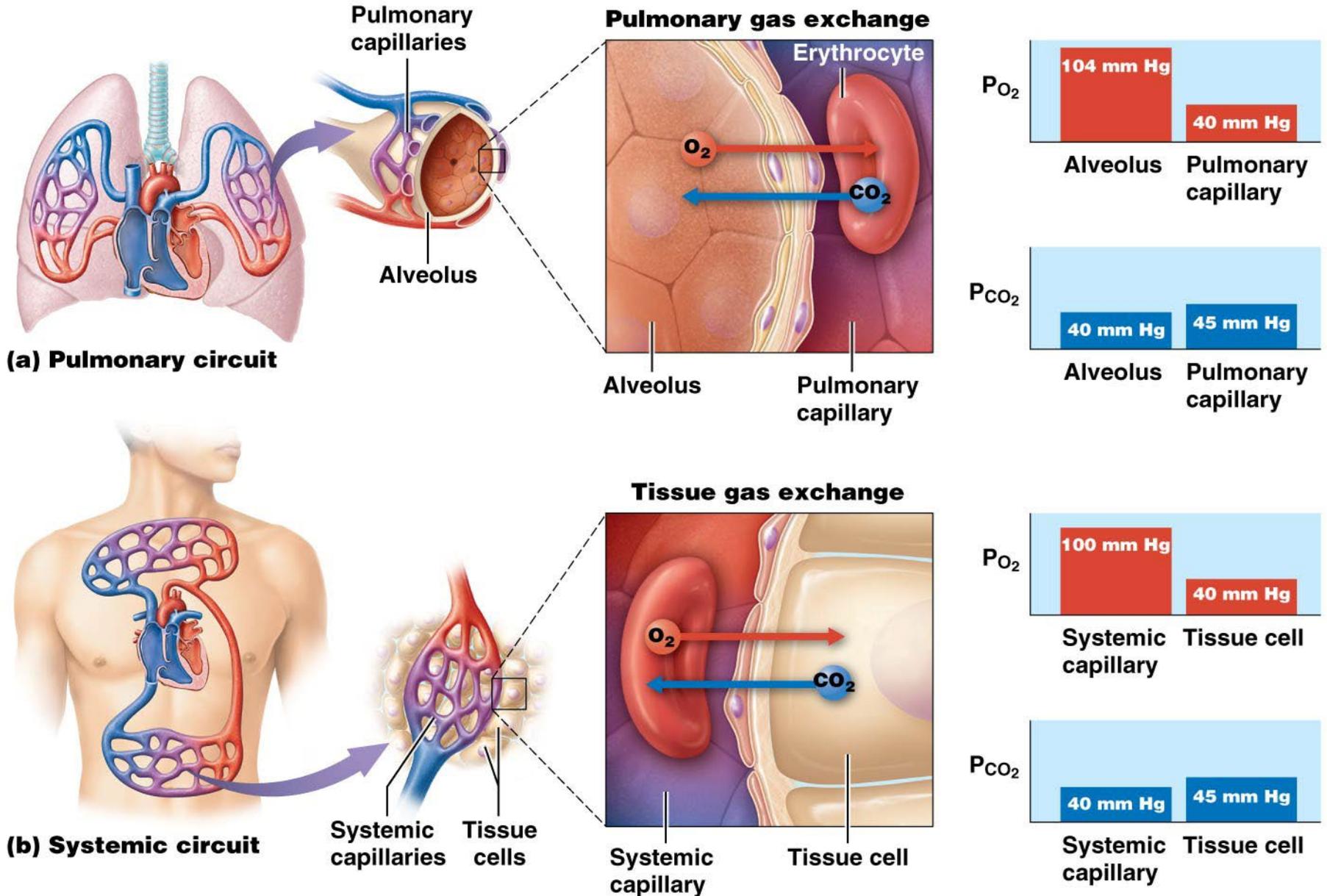


Bicarbonate forms ionic bond with sodium and is transported to lungs as sodium-bicarbonate

Hydrogen ion binds to Hb-O₂ and causes oxygen to be released from Hb // oxygen now free to diffuse to mitochondria

(a) Bicarbonate formation in an erythrocyte in a systemic capillary

Gas Transport in Systemic and Pulmonary Circuits



How is oxygen unloaded from RBC in the Systemic Gas Exchange?

O₂ unloading

H⁺ binding to HbO₂ reduces its affinity for O₂ /// H⁺ responsible for release of O₂ from iron

Iron in hemoglobin release oxygen

HbO₂ arrives at systemic capillaries 97% saturated (close to 100%!) /// leaves 75% saturated - unloads 25%

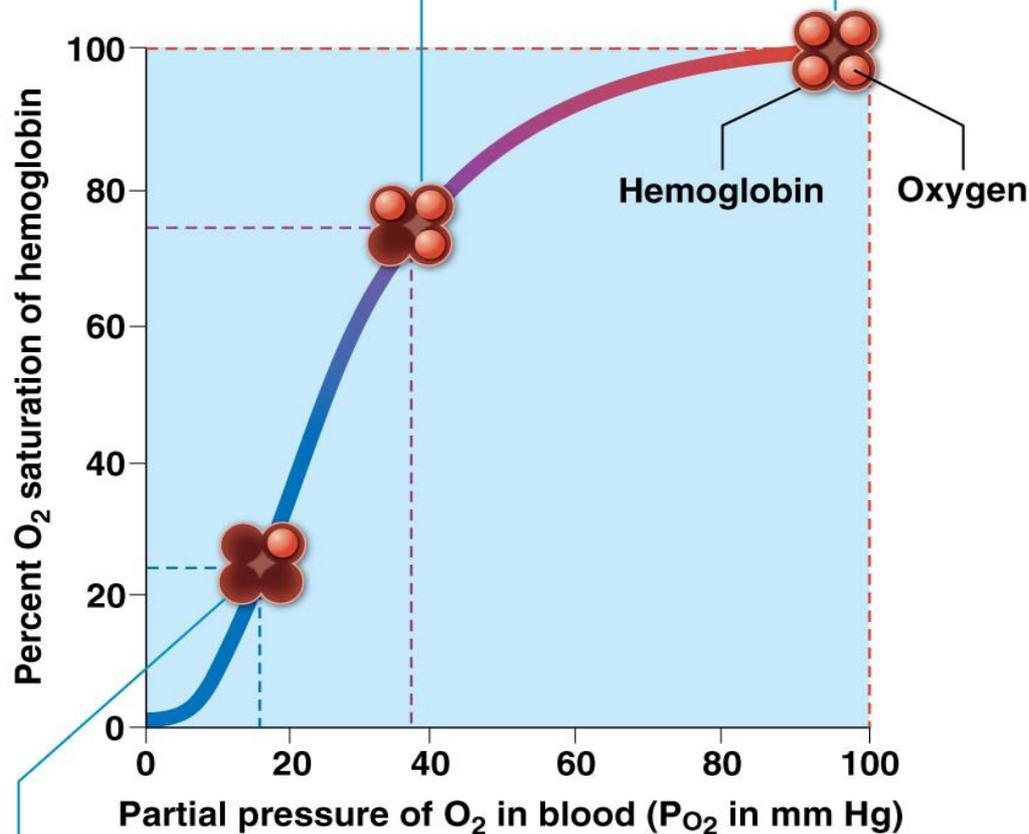
Venous reserve – oxygen remaining in the blood after it passes through the capillary beds

Utilization coefficient – gives up 22% of its oxygen load

How much oxygen is unloaded during periods of rest or exercise?

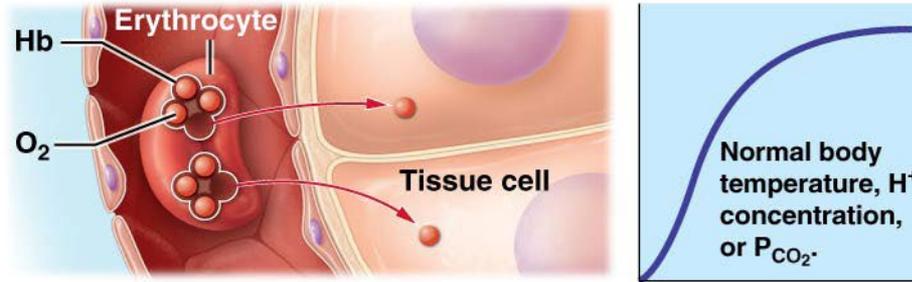
Point A: Hemoglobin (Hb) is almost 100% saturated in systemic arterial blood, as no O₂ has been unloaded to the tissues.

Point B: When a person is resting, Hb unloads only about 25% of its O₂ to the tissues. Therefore, Hb in systemic venous blood is about 75% saturated.

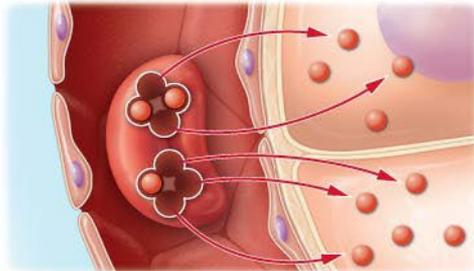


Point C: When a person exercises vigorously, Hb unloads most of its oxygen in the tissues. In such cases, the Hb entering venous blood is only about 25% saturated.

Effect of temperature, hydrogen ion concentration, and P_{CO_2} on oxygen unloading from hemoglobin.

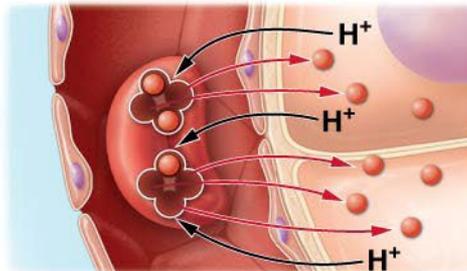


Temperature increases.



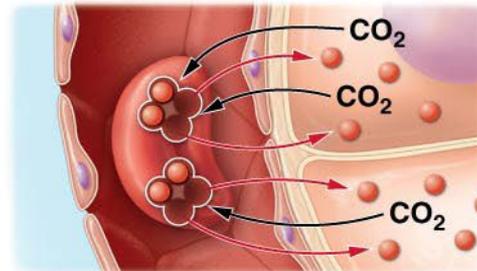
(a) Increased temperature causes more O_2 unloading.

H^+ concentration increases.

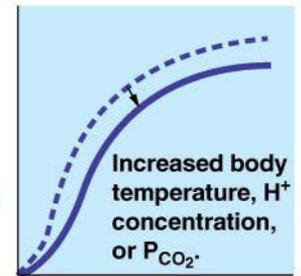


(b) Increased H^+ causes more O_2 unloading.

P_{CO_2} increases.



(c) Increased P_{CO_2} causes more O_2 unloading.



These variables all increase when you exercise. This is when we need Hb to release more oxygen so we can deliver more O_2 to our mitochondria (organelle that uses reduced coenzymes and oxygen) to make ATP

Another Molecule Used to Unload Oxygen

Bisphosphoglycerate (BPG) – intermediate in glycolysis // as concentration of BPG increases it indicates high level of anaerobic metabolism

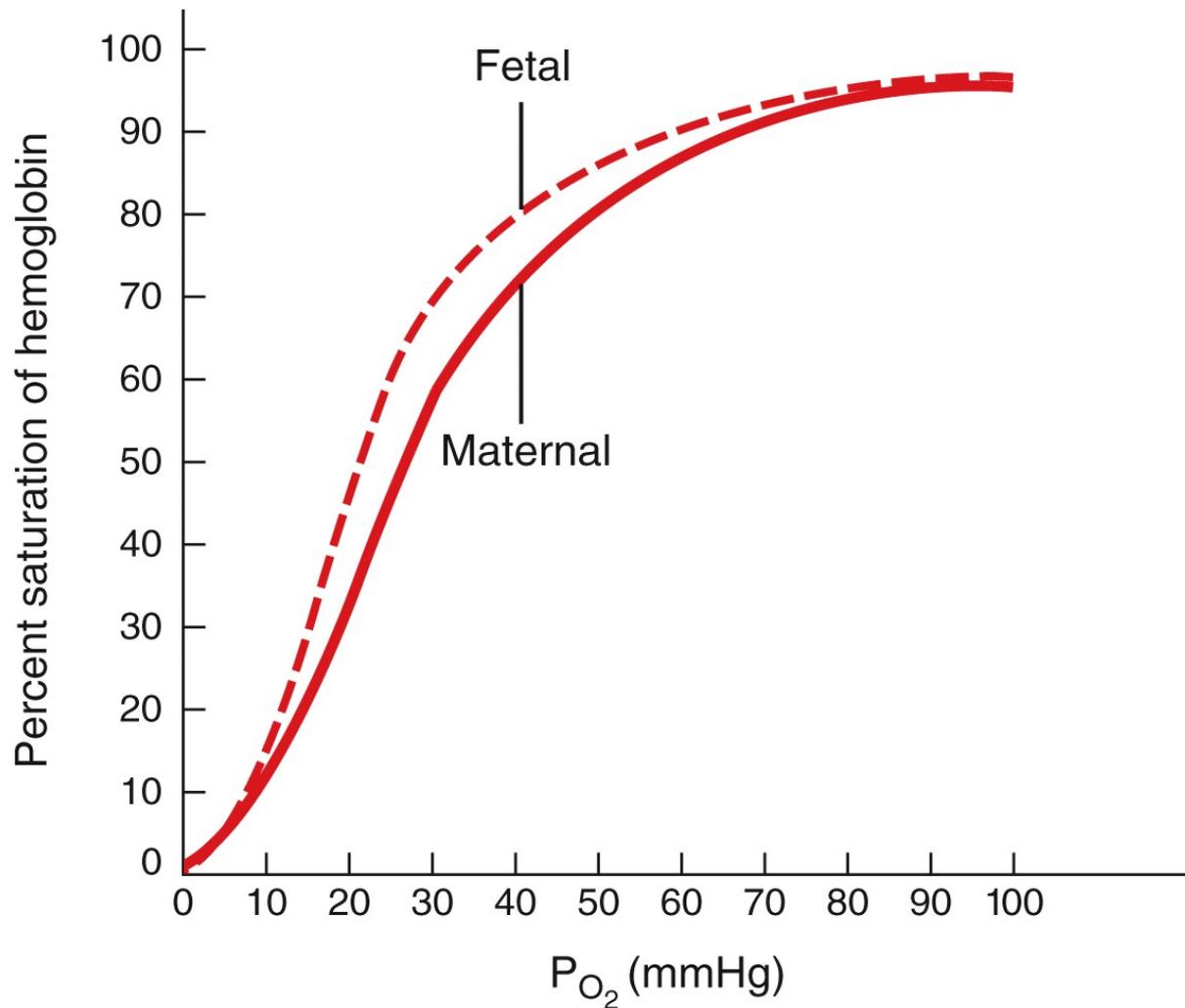
As RBCs produce more BPG, it binds to Hb and more O₂ is unloaded

↑ body temp (fever), thyroxine, growth hormone, testosterone, and epinephrine all raise BPG and cause more O₂ unloading

↑ metabolic rate requires ↑ oxygen

Haldane effect – rate of CO₂ loading is also adjusted to varying needs of the tissues /// low level of oxyhemoglobin enables the blood to transport more CO₂

This Chart Shows Fetal Hb Has a Greater
Affinity for O₂ Than Maternal Hb



Terminology: Variations in Respiratory Rhythm

eupnea – relaxed quiet breathing /// characterized by tidal volume 500 mL and the respiratory rate of 12 – 15 bpm

apnea – temporary cessation of breathing

dyspnea – labored, gasping breathing; shortness of breath

hyperpnea – increased rate and depth of breathing in response to exercise, pain, or other conditions

hyperventilation – increased pulmonary ventilation in excess of metabolic demand

hypoventilation – reduced pulmonary ventilation

Kussmaul respiration – deep, rapid breathing often induced by acidosis

respiratory arrest – permanent cessation of breathing

tachypnea – accelerated respiration rate

bradypnea – slow respiration rate