



มหาวิทยาลัยราชภัฏนครปฐม



Mechanism and Illness of Human Body 2

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Disorders of the Respiratory System

Pulmonary function test

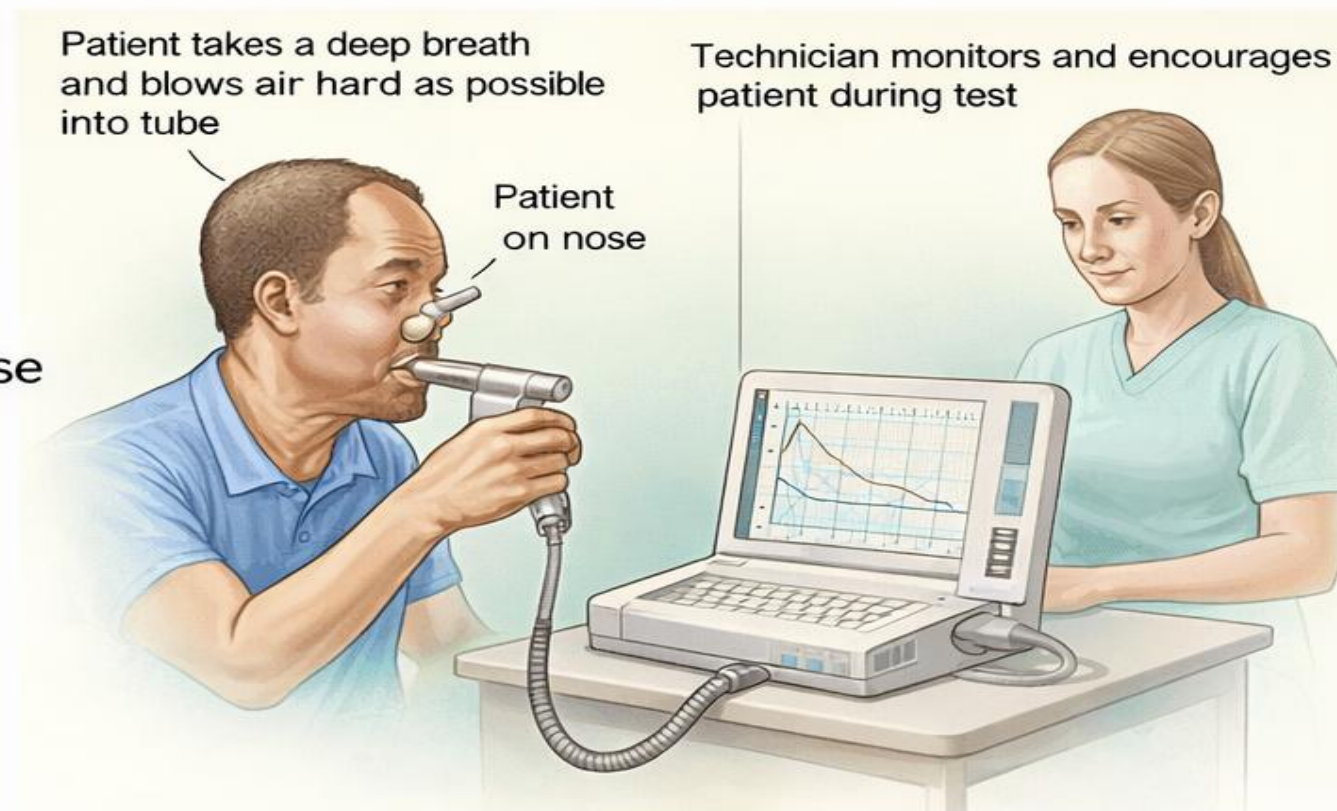
- **FEV1 (forced expiratory volume in 1 second)** measures flow that is exhaled forcefully in a one second period.
If low, indicates limited or obstructed airflow
- **FVC (forced vital capacity)** measures the amount of air that is forcibly exhaled and is the volume plateau or maximum achieved after 3-6 seconds exhalation

if $FEV1/FVC < 90\%$, indicates limited airflow
(obstructive lung disease)

if $FVC < 80\%$, suggests restrictive lung disease

if $FEF25-75 < 70\%$, may indicate small airway disease

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disease



Spirometry

Disorders of the Respiratory System

Pulmonary function test

- **PEFR (peak expiratory flow rate)** measures “peak flow” which is the highest flow achieved during forced exhalation. Often tested in Asthma

Bronchodilator (BD) response test

- Perform spirometry (pre-BD) 1st
- Then administer bronchodilator in nebulizer / DPI / MDI
- Wait 15-30 minutes, then perform post-BD
- Check FEV1 or PEFR, often tested in COPD



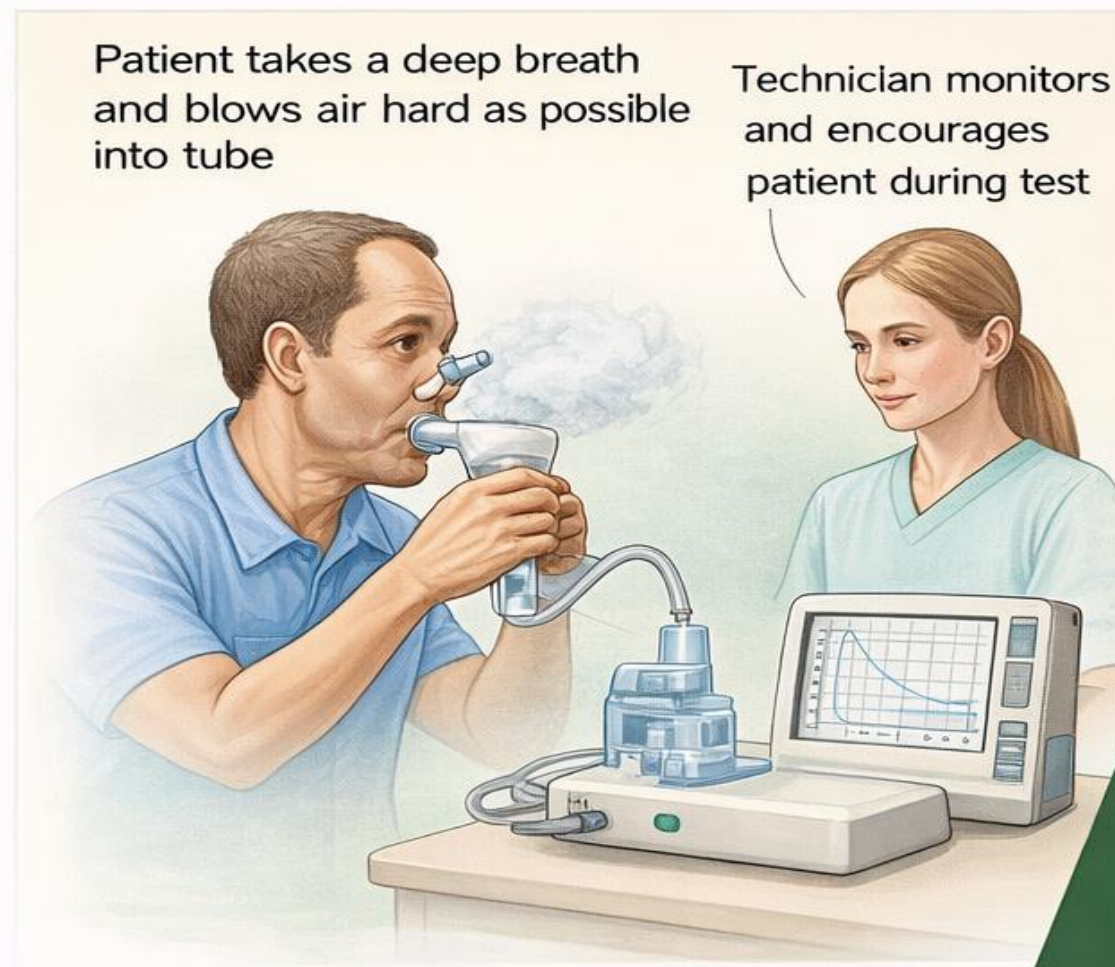
Nebulizer



DPI



MDI



Disorders of the Respiratory System

V/Q mismatch

- **Ventilation perfusion mismatch (V/Q mismatch)** is a condition involving an imbalance between the air (ventilation; V) and blood flow (perfusion; Q) around the alveoli.

Normal V/Q ratio should be $> 0.8-1.0$

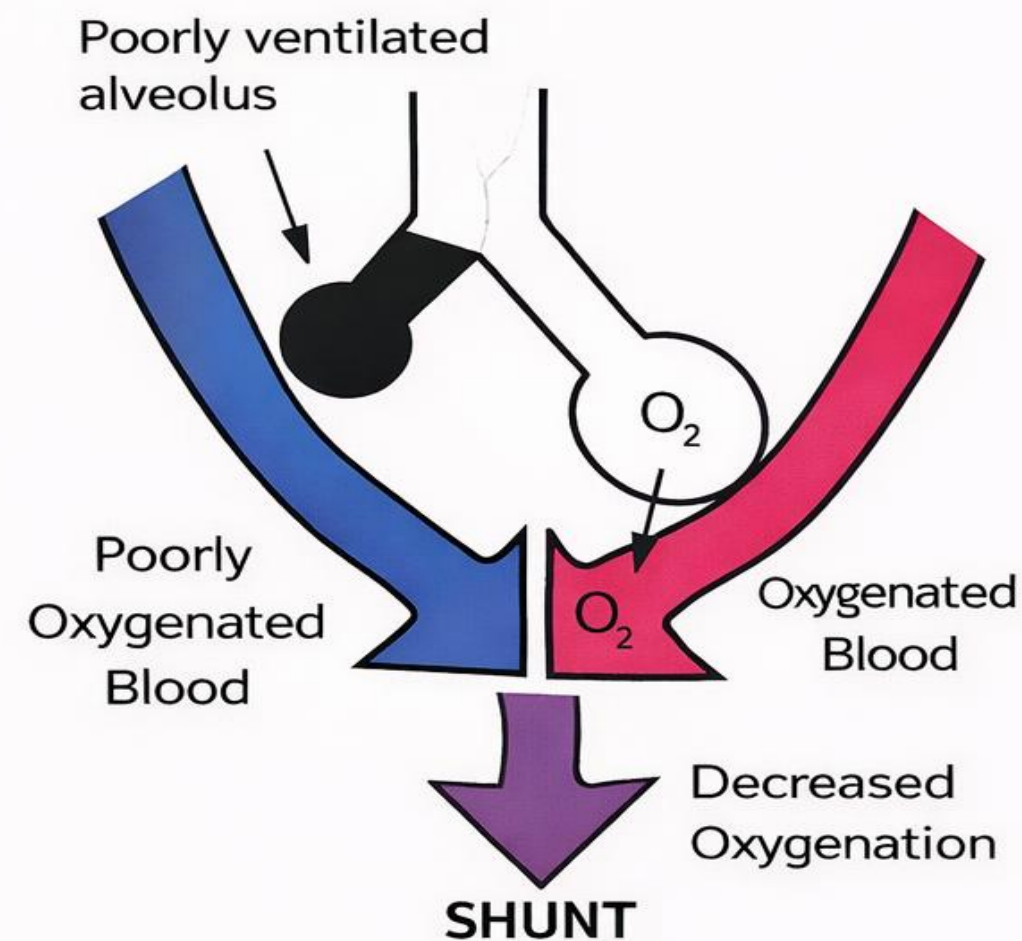
- Low V/Q ratio: Inadequate oxygenation of the body's cells

Causes of low V/Q ratio: Ventilation: V

1. Lung compliance (c)
2. Surface tension (s)
3. Constriction of respiratory muscles

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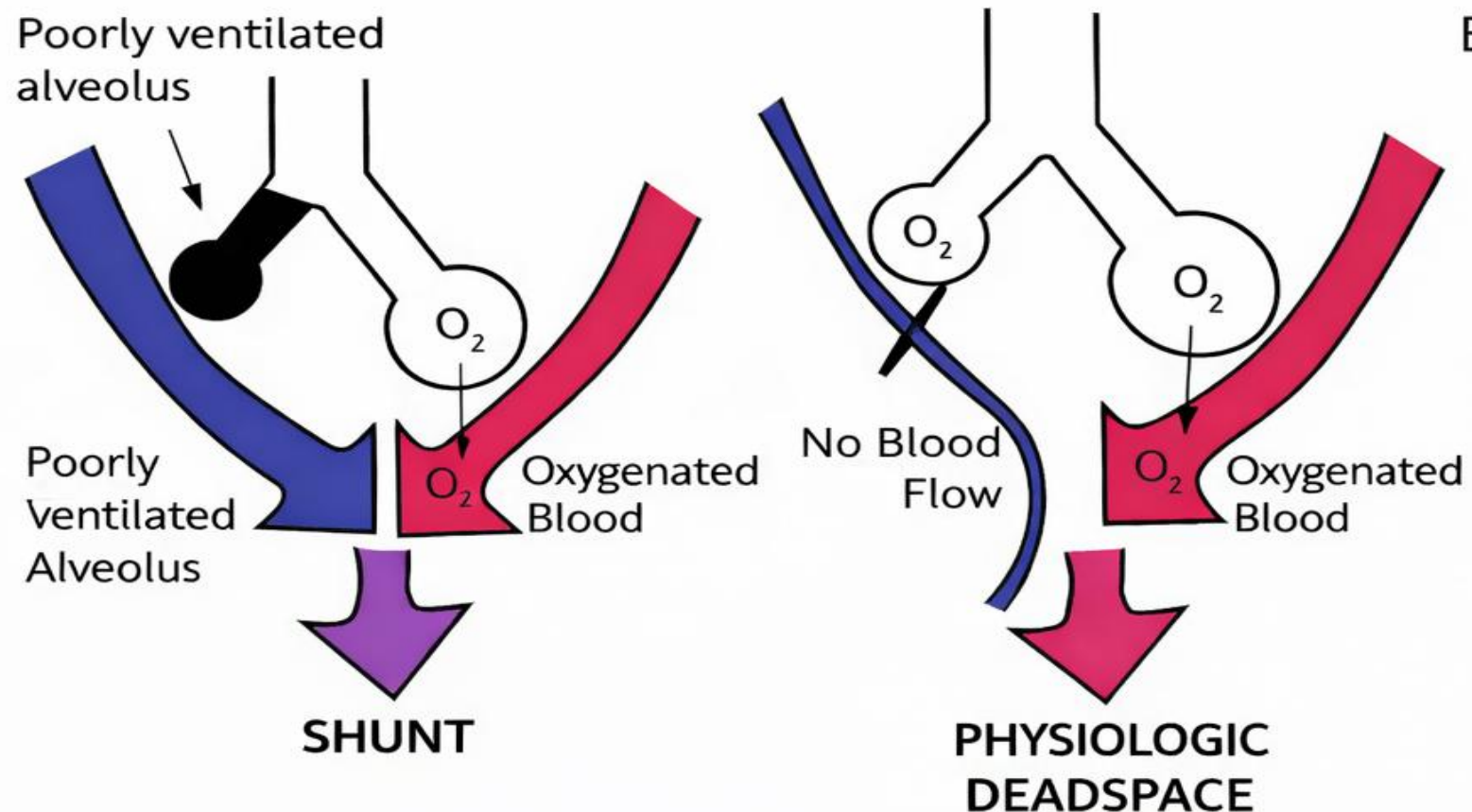


Disorders of the Respiratory System

V/Q mismatch

Normal resting V/Q should be approximately ~ 5,000 mL/min.

- $V/Q < 0.8$ is V/Q mismatch characteristic of a **shunt effect**
- $V/Q > 1$ is V/Q mismatch characteristic of a **dead space effect**



Examples of $V/Q > 1$ include:

- Decreased **perfusion** due to dehydration, blood loss or low cardiac output
- Increased **ventilation** from tachypnea, rapid breathing or high tidal volumes



Disorders of the Respiratory System

Diffusion impairment

Diffusion impairment occurs when the surface area for gas exchange is reduced or thickened, increasing diffusion time.

1. Differences between O_2 concentration in capillary blood and air in alveoli lead to **faster diffusion**
2. Alveolar Capillary Membrane **thickens, slowing diffusion rate**
3. Thickening of gas exchange tissue, reduces oxygen diffusion
4. Characteristics of diffusion impairment:
 - O_2 diffuses into capillary blood **longer than 0.25 sec**, averaging about **20 sec**
 - CO_2 diffuses from capillary blood into alveoli **faster than O_2** due to **higher solubility**
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Disorders of the Respiratory System

Dead space

A region where Ventilation (V) occurs but Perfusion (Q) does not, resulting in no gas exchange (no function).

1. Anatomical dead space:

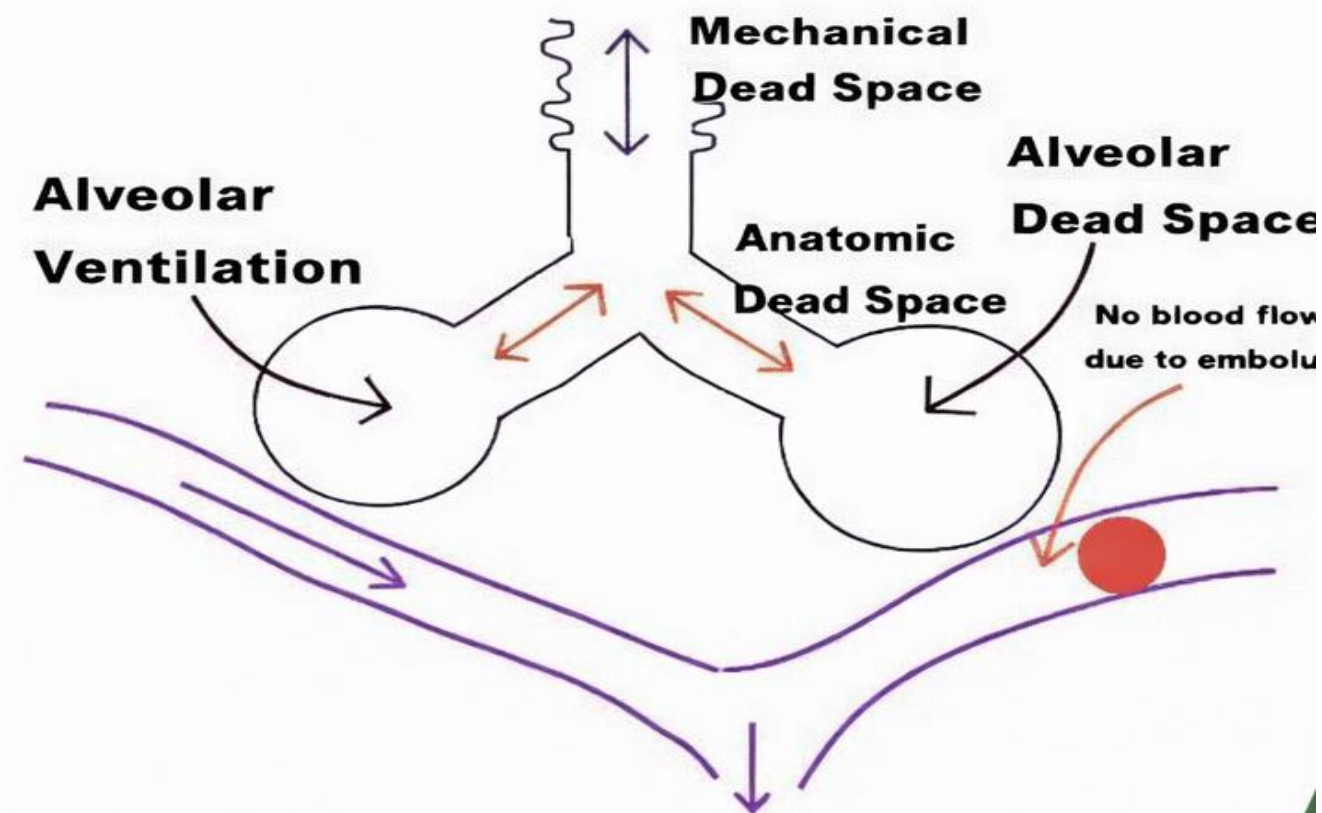
Air within the airway that does not participate in gas exchange

2. Alveolar dead space:

Air within the alveolus that does not participate in gas exchange

3. Mechanical dead space:

Air that is within ventilated mechanical conduits.



Disorders of the Respiratory System

Shunting

- **Right to Left shunt**

- **Right to Left shunt**

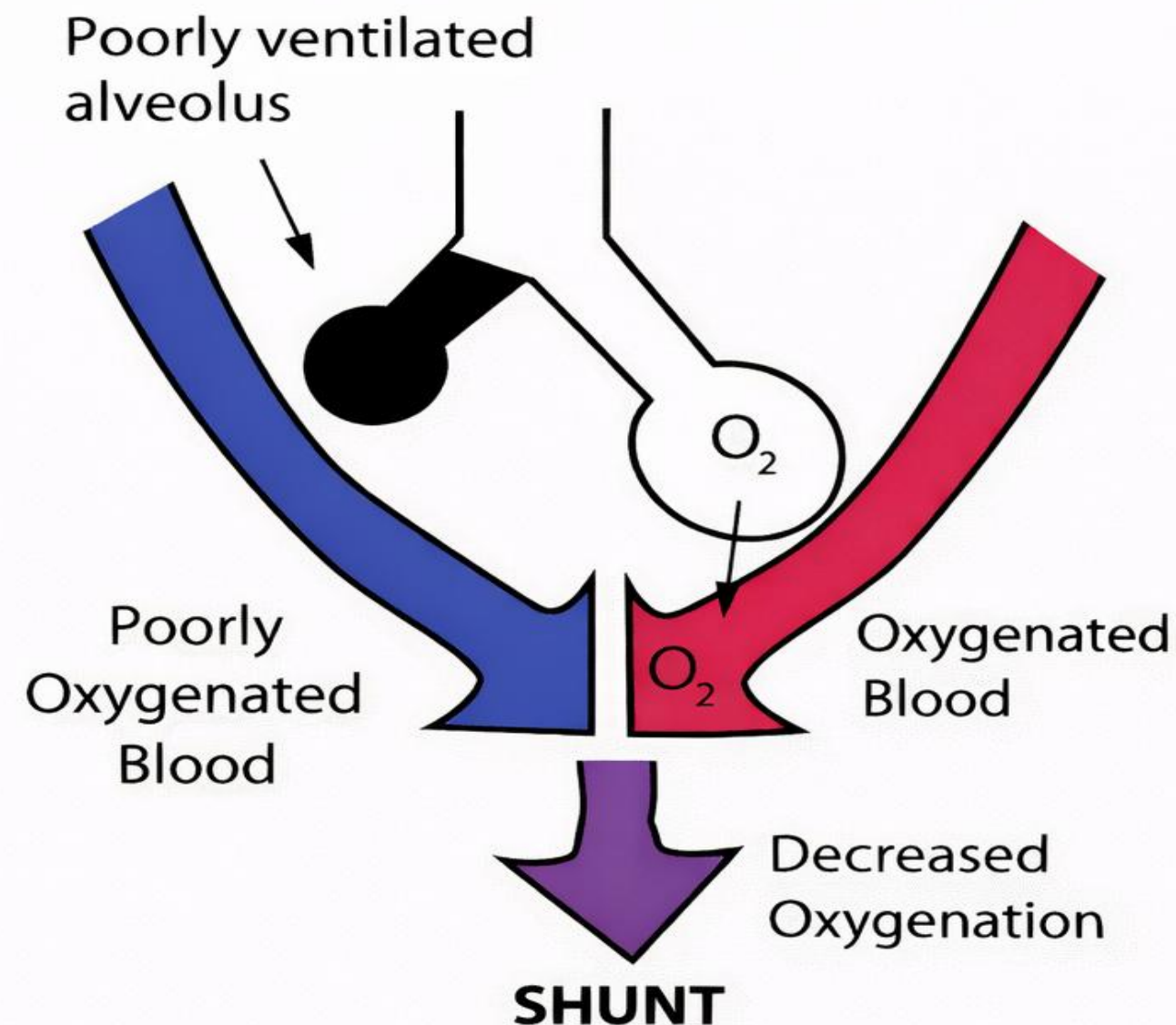
in ISD, VSD, PDA

- **Left to Right shunt**

Pulmonary stenosis

- **Capillary shunt**

Atelectasis, ARDS (acute respiratory distress syndrome)





Disorders of the Respiratory System

Type of Hypoxia

Hypoxia is a condition where tissues have **inadequate oxygen** (O_2).

Hypoxemia is a condition where the blood has **lower than normal O_2 levels**.

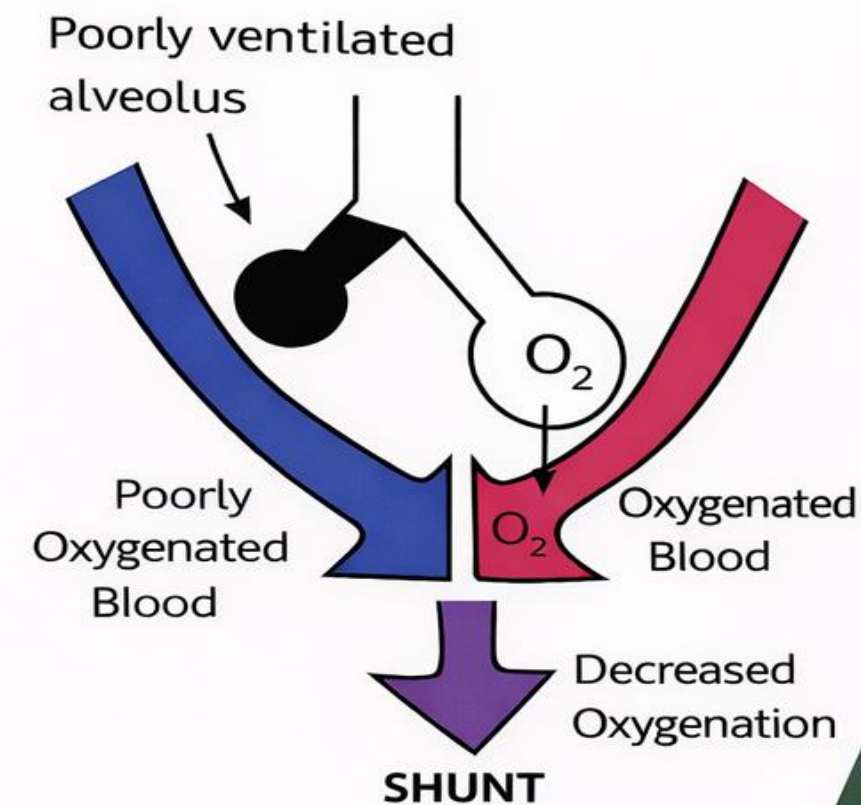
- **Histotoxic hypoxia or Cellular hypoxia:**

Anemia decreases Hb, releasing O_2 in hemoglobin (Hb).

Toxins or poisons like CO (which binds to hemoglobin more than 200-250 times greater than O_2), sulfanilamide, Cyanide, Carbon Monoxide can prevent tissue from using oxygen

- **Hypoxemic hypoxia**

1. Low inspired partial pressure of O_2
2. Alveolar hypoventilation
3. Intrapulmonary shunt
4. Dead space ventilation
5. Ventilation – Perfusion mismatch
6. Diffusion defect



Disorders of the Respiratory System

Type of Hypoxia

- **Circulatory hypoxia or Stagnant hypoxia:**

This is hypoxia resulting from **decreased flow** of blood toward tissues due to:

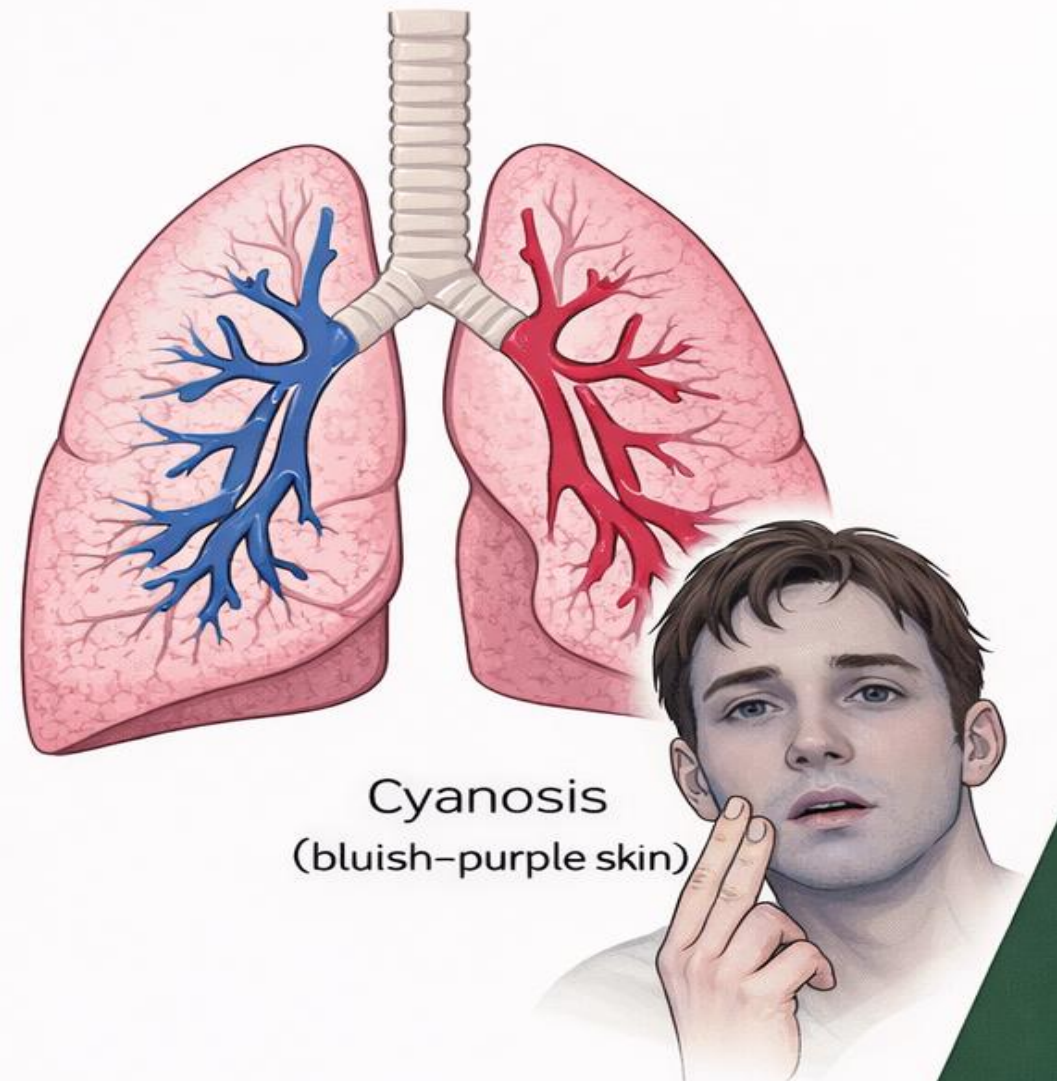
- Hypovolemic shock
- Distributive shock
- Cardiogenic shock or Obstructive shock

- **Histotoxic hypoxia or Cellular hypoxia:**

This is hypoxia caused by cells that receive O_2 but cannot use it due to disruption of cellular oxidative processes by:

- Cyanide poisoning
- Sulfanilamide
- Carbon monoxide

Cyanosis can be easily observed in hypoxic hypoxemic hypoxia and Circulatory hypoxia.



Disorders of the Respiratory System

Signs of Hypoxia

Nervous System

- Accounts for basal oxygen consumption (20%). Uses O_2 45 mL/min to produce ATP needed for control of Na-K pump, neurotransmitter synthesis, synapse signaling
- if $PaO_2 < 40-45$ mmHg → acidosis increases cerebral cortex permeability → cerebral edema occurs.



Mild hypoxemia: PaO_2 60 - 80 mmHg
Moderate hypoxemia: PaO_2 40-60 mmHg
Severe hypoxemia: $PaO_2 < 40$ mmHg

Cardiovascular System

- Sympathetic stimulation increases cardiac output (SV, HR).
- Vasodilation increases blood flow to essential organs: function of heart increases blood flow, oxygen demand and heart rate.
- Metabolic acidosis leads to clubbing finger, 2,3-diphosphoglycerate (2,3-DPG) rises O_2 delivery increases.

Respiratory System

- Low O_2 + lactic acidosis stimulates carotid body to send signals to speed up rapid, forceful breathing. May use accessory muscles e.g. belly muscles.
- $PaO_2 < 30$ mmHg → medullary center depressed → Cheyne-Stokes breathing → apnea in severe cases.



Kidneys & Muscle System

- Low O_2 + lactic acidosis ↓ blood flow to kidneys.
- Renin erythropoietin factor (REF) stimulates erythropoietin (EPO) to stimulate excessive RBC formation leading to high viscosity blood, ↑ pulmonary hypertension risk.
- Low O_2 leads to centrilobular fibrosis.
- Muscle tissue breaks down stored ATP (creatinine phosphate) leading to glycolysis





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