# An Approach to Dyspnea: from atmosphere to hemoglobin

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Definition: a subjective sensation of abnormal or uncomfortable breathing

# Pathophysiologic Mechanisms of Dyspnea:

- Decreased arterial O<sub>2</sub> levels
- Increased CO<sub>2</sub> levels
- Low blood pH
- Neuromechanical dissociation
  - o a mismatch between respiratory effort and ventilation
    - elicits dyspnea in cases of abnormal chest wall compliance or airway resistance

## Etiologies: follow O<sub>2</sub> as we breathe

1) Drive to breathe

- The impulse to breathe is generated in the brainstem (medulla and pons) via mechanical and chemical stimuli
- Together with pulmonary and skeletal muscle stretch receptors, O<sub>2</sub> and CO<sub>2</sub> are the major contributors to the complex feedback control system of basic breathing
  - $\circ$  O<sub>2</sub> is sensed by peripheral chemoreceptors in the carotid and aortic bodies
  - $\circ$  CO<sub>2</sub> and pH are sensed primarily in the brain by medullary chemoreceptors
- Muted or absent responses can lead to a decreased "drive" to breathe, as in obesity hypoventilation

### 2) Generating negative pressure

- We generate negative pressure to move oxygen from the atmosphere into the lungs
- Abnormal neuromuscular function and/or poor respiratory system compliance can lead to neuromechanical dissociation and cause CO<sub>2</sub> retention
- Abnormal neuromuscular function:
  - Diaphragmatic paralysis
  - Myasthenia gravis
  - o Guillain-Barré
- Poor respiratory system compliance
  - Pulmonary fibrosis
  - Pleural effusions
  - Obesity
  - Ascites
  - o Pregnancy

#### 3) <u>Airways</u>

- Large airway disorders:
  - o COPD
  - o Asthma
  - o Bronchospasm
  - Obstructing tumor
  - Foreign body
- Small airway diseases (i.e. bronchiolitis, appearing as "tree-in-bud" opacities on CT imaging):
  - o Infectious: viral, bacterial, mycobacterial

- o Inflammatory: RA, SLE, vasculitis
- Fibro-proliferative: post lung transplantation
- o Inhalational: exposure to tobacco, toxic fumes, mineral dusts

# 4) Alveolar filling processes

- Cause dyspnea by impairing gas exchange due to shunt
  - Common etiologies:
    - Blood (alveolar hemorrhage)
    - Pus (pneumonia)
    - Water (pulmonary edema)
    - o Atelectasis can cause shunt due to complete collapse of alveoli

## 5) Alveolar-capillary membrane

- Gas exchange depends on a thin alveolar-capillary membrane and a large surface area
- Diseases that destroy (emphysema) or thicken (pulmonary fibrosis) the alveolar-capillary membrane decrease the total membrane surface area and slow the rate of diffusion
- Membrane abnormalities are rarely a cause of dyspnea. Rather, these disorders have other features as a source of dyspnea, as in COPD (hyperinflation, air trapping) or fibrosis (decreased compliance, increased work of breathing)
- One exception: alveolar-capillary membrane diseases can result in dyspnea during exercise due to shortened capillary transit time

## 6) O<sub>2</sub> transport in the blood

- The majority of O<sub>2</sub> is bound to hemoglobin, with some dissolved in blood as well
- Low O<sub>2</sub> content can result from decreased total hemoglobin or functional alterations of hemoglobin leading to impaired O<sub>2</sub> binding
- Two main etiologies:
  - o Anemia
  - Dyshemoglobinemias (CO poisoning or methemoglobinemia)

#### 7) Inefficient blood flow

- Cardiac disorders: systolic or diastolic heart failure, MI, arrhythmia, tamponade
- Pulmonary vascular diseases: pulmonary hypertension, PE

#### Pearls:

- 1. Dyspnea is not always from a pulmonary disorder
- 2. An easy way to build a differential diagnosis is to follow the oxygen molecule from atmosphere to hemoglobin
- 3. Blood, pus, or water are the main causes of alveolar shunt

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